

# COVID-19

## Selection of peer-reviewed articles

Special focus on Pharmaceutical and non-Pharmaceutical  
interventions from 18-06-20

UPDATE OF  
19 AUGUST 2020

REACTing shares a selection of the most relevant articles published on COVID-19 on a weekly basis. This literature review not only presents a selection of references, but also highlights the key points and messages from each article. It does not include pre-print articles.

Our objective is to help the scientific community, health-workers and public health decision makers, being up to date with the latest scientific research.

### Coordinated by:

REACTing:  
Renaud Vatrinet  
& Eric D'Ortenzio

### Documented by:

Inserm- Collective  
Expertise Unit:  
Bénédicte Varignon  
& Laurent Fleury

### Redaction committee

Inserm- Thematic Institute of Immunology,  
Inflammation, Infectiology, and Microbiology (I3M):  
Guia Carrara; Eric D'Ortenzio; Evelyne Jouvin-Marche;  
Boris Lacarra; Claire Madelaine; Inmaculada Ortega-  
Perez; Oriane Puéchal; Erica Telford; Renaud Vatrinet

### With a precious help from:

- Inserm- Department of Partnerships and External Relations (DPRE)
- Inserm- USA office

### Additional links:

Rapid Evidence Reviews Group: <https://isaric.tghn.org/covid-19-rapid-evidence-reviews-group/>

Biblioovid: <https://biblioovid.org>

CORD19 Publication Dashboard: <https://france-science.com/en/homepage-english-2/>

Journal and date	Title	Authors and link	Field of expertise	Key facts
EBioMedicine 17AUG2020	<b>SARS-CoV2 vertical transmission with adverse effects on the newborn revealed through integrated immunohistochemical, electron microscopy and molecular analyses of Placenta</b>	Facchetti F et al Italy <a href="#">gotopaper</a>	Clinic	<p><b>AIM:</b> debate around the occurrence of trans-placental transmission of SARS-CoV-2 infection</p> <p>Screened for SARS-CoV spike protein expression placentas from 101 women: 15 tested positive for SARS-CoV-2 + 34 tested negative + 52 not evaluated Immunostatin for SARS-CoV-2 nucleocapsid was performed in the placentas of all COVID-19 positive women.</p> <p><b>Results:</b></p> <ul style="list-style-type: none"> <li>- SARS-CoV-2 antigens, RNA and/or particles morphologically consistent with coronavirus were identified in villous syncytiotrophoblast, endothelial cells, fibroblasts, in maternal macrophages, and in Hofbauer cells and fetal intravascular mononuclear cells</li> <li>- Absence of villitis was associated with an increase in the number of Hofbauer cells, which expressed PD-L1 □ may have prevented immune cell-driven placenta damage.</li> </ul> <p>□ We provide first-time evidence for maternal-fetal transmission of SARS-CoV-2, likely propagated by circulating virus-infected fetal mononuclear cells</p> <p>□ permissiveness of trans-placental SARS-CoV-2 transmission is rare.</p> <p>□ depend on host genetic factor, the infection with a unique SARS-CoV-2 genetic variant and/or the production of a high load of maternal antibody-virus immune complexes</p>
JAMA 14AUG2020	<b>Association Between Number of In-Person Health Care Visits and SARS-CoV-2 Infection in Obstetrical Patients</b>	Reale S.C., et al. USA <a href="#">gotopaper</a>	Public Health / Epidemio	<p><b>AIM:</b> to examine whether the number of in-person health care visits of obstetrical patients from 4 hospitals in Boston (USA) between April and June 2020 was associated with the risk of SARS-CoV-2 infection.</p> <ul style="list-style-type: none"> <li>- Of 2968 deliveries, 111 patients (3.7%) tested positive for SARS-CoV-2 infection, of which 45 tested positive for SARS-CoV-2 infection antenatally and 66 tested positive at the time of admission for labour and delivery.</li> <li>- Performing a nested case-control study, 93 cases were matched with 372 control observations. The mean number of in-person visits was 3.1 for cases and 3.3 for controls. For the association between the number of in-person health care visits and SARS-CoV-2 infection, the odds ratio was 0.93 per additional visit.</li> </ul> <p>There was no meaningful association between the number of in-person health care visits and the rate of SARS-CoV-2 infection in this sample of obstetrical patients, despite the high infection rate in Boston. Necessary, in-person care should be therefore safely performed.</p>
Cell 14AUG2020	<b>Robust T cell immunity in convalescent individuals with asymptomatic or mild COVID-19</b>	Sekine et al., Sweden <a href="#">gotopaper</a>	Immuno	<p>SARS-CoV-2-specific memory T cells will likely prove critical for long-term immune protection against COVID-19. Method: Systematic mapping of the functional and phenotypic landscape of SARS-CoV-2-specific T cell responses in unexposed individuals, exposed family members, and individuals with acute or convalescent COVID-19.</p> <p><b>Results:</b></p> <ul style="list-style-type: none"> <li>- Acute phase SARS-CoV-2-specific T cells display an activated cytotoxic phenotype</li> <li>- Broad and polyfunctional SARS-CoV-2-specific T cell responses in convalescent phase</li> <li>- Detection of SARS-CoV-2-specific T cell responses also in seronegative individuals</li> </ul> <p><b>Conclusions:</b> SARS-CoV-2 elicits robust, broad and highly functional memory T cell responses, suggesting that natural exposure or infection may prevent recurrent episodes of severe COVID-19.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 13AUG2020	<b>Effect of an Inactivated Vaccine Against SARS-CoV-2 on Safety and Immunogenicity Outcomes</b> Interim Analysis of 2 Randomized Clinical Trials	Xia et al., China <a href="#">gotopaper</a>	Vaccine	<p>Interim analysis of 2 randomized placebo-controlled trials</p> <p>In 96 healthy adults in a phase 1 trial of patients randomized to aluminum hydroxide (alum) only and low, medium, and high vaccine doses on days 0, 28, and 56:</p> <ul style="list-style-type: none"> <li>- 7-day adverse reactions occurred in 12.5%, 20.8%, 16.7%, and 25.0%, respectively</li> <li>- geometric mean titers of neutralizing antibodies at day 14 after the third injection were 316, 206 and 297 in the low-, medium-, and high-dose groups, respectively</li> </ul> <p>In 224 healthy adults randomized to the medium dose :</p> <ul style="list-style-type: none"> <li>- 7-day adverse reactions occurred in 6.0% and 14.3% of the participants who received injections on days 0 and 14 vs alum only, and 19.0% and 17.9% who received injections on days 0 and 21 vs alum only, respectively</li> <li>- geometric mean titers of neutralizing antibodies in the vaccine groups at day 14 after the second injection were 121 vs 247, respectively.</li> </ul> <p>-&gt; This inactivated COVID-19 vaccine had a low rate of adverse reactions and demonstrated immunogenicity, but longer-term assessment of safety and efficacy will require phase 3 trials.</p>
JAMA 13AUG2020	<b>Outcomes Associated with Use of a Kinin B2 Receptor Antagonist Among Patients With COVID-19</b>	Van de Veerdonk. et al Netherlans <a href="#">gotopaper</a>	Therapeutic	<p>AIM: Loss of ACE2 might lead to plasma leakage + activation of the plasma kallikrein-kinin system that could contribute to pulmonary angioedema via stimulation of bradykinin 2 receptors.</p> <p>Case-control study: 10 patients - 3 doses of 30 mg of icatibant SC injection at 6-hour intervals (bradykinin 2 receptor antagonist).</p> <p>Nine cases were matched to 18 controls for sex, age, body mass index and day of illness.</p> <p><b>Results:</b> 90% were men - mean age was 55 years for case and 58 years for controls.</p> <ul style="list-style-type: none"> <li>- After 3 injection: 44% cases were no longer oxygen dependent</li> <li>- 55% with substantial decrease of oxygen supplementation,</li> <li>- Controls: 17% showed a spontaneous reduction in oxygen supplementation,</li> <li>- Icatibant was well tolerated – no adverse events</li> </ul> <ul style="list-style-type: none"> <li><input type="checkbox"/> association between receipt of icatibant and improved oxygenation</li> <li><input type="checkbox"/> might be beneficial especially in early stages of disease</li> <li><input type="checkbox"/> icatibant's short half life</li> <li><input type="checkbox"/> randomized trial?</li> </ul>
The Lancet Haematology 13AUG2020	<b>Clinical characteristics and risk factors associated with COVID-19 severity in patients with haematological malignancies in Italy: a retrospective, multicentre, cohort study</b>	Passamonti et al., Italy <a href="#">gotopaper</a>	Clinic	<p>The Italian Hematology Alliance on COVID-19 : Collection of data from adult patients with haematological malignancies who required hospitalisation for COVID-19.</p> <p>Multicentre, retrospective, cohort study included adult patients (aged ≥18 years) with diagnosis of a WHO-defined haematological malignancy admitted to 66 Italian hospitals between Feb 25 and May 18, 2020, with laboratory-confirmed and symptomatic COVID-19.</p> <ul style="list-style-type: none"> <li>-&gt; 198 (37%) of 536 patients died</li> <li>-&gt; When compared with the general Italian population with COVID-19, the standardised mortality ratio was 2.04 in the whole study cohort and 3.72 (2.86–4.64) in individuals younger than 70 years.</li> <li>-&gt; When compared with the non-COVID-19 cohort with haematological malignancies, the standardised mortality ratio was 41.3</li> </ul> <p>Were associated with worse overall survival :</p> <ul style="list-style-type: none"> <li>Older age - Progressive disease status - Diagnosis of acute myeloid leukaemia - Indolent non-Hodgkin lymphoma - Aggressive non-Hodgkin lymphoma - Plasma cell neoplasms - Severe or critical COVID-19</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Clin Inf Dis 12AUG2020	<b>Methylprednisolone as Adjunctive Therapy for Patients Hospitalized With COVID-19 (Metcovid): A Randomised, Double-Blind, Phase IIb, Placebo-Controlled Trial</b>	Prado Jeronimo CM et al Brazil <a href="#">gotopaper</a>	Therapeutic	<p><b>AIM:</b> evaluating the efficacy of methylprednisolone (MP) among hospitalized patients with suspected COVID-19</p> <p>Double blind, placebo-controlled, randomized (1:1), phase IIb Intravenous MP or placebo twice daily for 5 days. 416 randomized patients and 393 analyzed as mITT: 194 MP &amp; 199 placebo.</p> <p><b>Results:</b></p> <ul style="list-style-type: none"> <li>- No patients received anti IL-6, anti IL-1, remdesivir or convalescent plasma therapy,</li> <li>- No difference on mortality at day 28 between groups: 37,1% in the MP group vs 38,2% in the placebo group (p= 0,629),</li> <li>- Subgroup analysis: patient &gt;60 years in the MP group had a lower mortality rate at day 28: 46,6% (MP group) vs 61,9% (placebo) – p=0,039</li> <li>- Patients &gt; 60 years had higher CRP values: 81,3 vs 74,7 (p=0,0028)</li> <li>- No difference in virus clearance in respiratory secretion until day 7.</li> </ul> <p>□ no evidence of improved survival in the overall population with a short course of intravenous MP in patients hospitalized with COVID-19.</p> <p>□ lower mortality in patients over 60 years who received MP: more pronounced systemic inflammatory status (higher CRP).</p>
Nature 12AUG2020	<b>Phase 1/2 study of COVID-19 RNA vaccine BNT162b1 in adults</b>	Mulligan M J et al, USA <a href="#">gotopaper</a>	Vaccine	<p>Safety, tolerability and immunogenicity data - 45 healthy adults.</p> <p>Placebo-controlled, observer-blinded dose escalation study. Randomized to receive 2 doses, separated by 21 days of 10 µg, 30 µg, or 100 µg of BNT162b1.</p> <p>à 12 participants per dose level (10 and 30 µg) were vaccinated with BNT162b1 on D1 &amp; D21</p> <p>à 12 participants received a 100 µg dose on D1.</p> <p><b>BNT162b1:</b> nucleoside-modified mRNA vaccine that encodes trimerized SARS-CoV-2 spike glycoprotein receptor-binding domain (RBD).</p> <p><b>Tolerability and safety profile:</b></p> <ul style="list-style-type: none"> <li>- Consistent with those previously observed for mRNA-based vaccines.</li> <li>- AE: 50% of participant who received either 10 or 30 µg of BNT162b1 – 58,3% of those who received 100 µg – 11,1% of placebo recipients.</li> <li>- SAE: 2 patients: Grade 3 fever 2 days after vaccination in the 30 µg group, and sleep disturbance 1 day after vaccination in the 100 µg group.</li> </ul> <p><b>Immunogenicity:</b></p> <ul style="list-style-type: none"> <li>- Robust immunogenicity was observed after vaccination.</li> <li>- RDB-binding IgG concentrations and SARS-CoV-2 neutralizing titers in sera increased with dose level and after a second dose.</li> <li>- Geometric mean neutralizing titers reached 1.9- to 4.6-fold that of a panel of COVID-19 convalescent human sera at least 14 days after a positive SARS-CoV-2 PCR.</li> </ul> <p>Encouraging and strongly support accelerated clinical development.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Lancet Global Health 10AUG2020	<b>Effect of the COVID-19 pandemic response on intrapartum care, stillbirth, and neonatal mortality outcomes in Nepal: a prospective observational study</b>	Ashish KC et al Nepal/Sweden/... <a href="#">gotopaper</a>	Epidemiology	<p><b>Maternal and neonatal health service all over the world are affected by the pandemic.</b></p> <p>Prospective observational study between Jan 1 and May 30 2020, from 9 hospital in Nepal. à 12,5 weeks before the national lockdown and 9,5 weeks during the lockdown</p> <p>Exclusion: gestational age &lt; 22 weeks, no fetal heart sound at admission, multiple birth 21763 women and 20354 gave birth</p> <p>The mean weekly number of institutional birth decrease: 1261,1 before lockdown to 651,4 birth during lockdown (-52,4%) The institutional stillbirth rate increased from 14 per 1000 total birth before lockdown to 21 per 1000 during lockdown (p=0,0002)</p> <p>The institutional neonatal mortality increased from 13 per 1000 livebirths to 40 per 1000 (p=0,0022) Intrapartum fetal rate monitoring decrease by 13,4%</p> <p>Practice skin-to-skin with their mother increased by 13,2% and health workers' hand hygiene practices during childbirth increased by 12,9% during lockdown.</p> <ul style="list-style-type: none"> <li>- <b>Institutional childbirth reduced by more than half during lockdown</b></li> <li>- <b>increases in institutional stillbirth rate and neonatal mortality</b></li> <li>- <b>decrease in quality of care</b></li> <li>- <b>urgent need exists to protect access to high quality intrapartum care and prevent excess death</b></li> </ul>
Clinical Infectious Disease 09AUG2020	<b>Clinical Outcomes Associated with Methylprednisolone in Mechanically Ventilated Patients with COVID-19</b>	Nelson B C et al USA <a href="#">gotopaper</a>	Clinic	<p><b>Evaluation of the association between use or methylprednisolone and key clinical outcomes</b></p> <p>Case-control study – a subset of patients also underwent propensity-score matching. Primary outcome: ventilator-free day by 28 days after admission Characteristics: 117 patients – propensity matching yielded a cohort of 42 well-matched pairs. Median age: 63y – IQR [52 – 71] – 67% male Median BMI: 30 kg/m<sup>2</sup> Median SOFA score: 11 [8 – 12] 33 patients died by hospital day 28 Results: - 48 patients received methylprednisolone – median dose: 80 mg/d - The matched group were comparable at baseline - Patients who received steroids were less likely to have received HQC or Azithro - Ventilator-free days was significantly higher in steroid group (6,2 vs 3,14, p=0,044) - Probability of extubation by day 28 was significantly higher in patients who received steroid (45% vs 21%) No significant differences in mortality (19% vs 36%, p=0,087)</p> <p>Multivariable linear regression: - Only methylprednisolone use was associated with higher number of ventilator-free day</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nat Med 7AUG2020	Using a real-world network to model localized COVID-19 control strategies	Firth et al., UK <a href="#">gotopaper</a>	Public Health / Epidemiology	<p><b>Simulation of control strategies for SARS-CoV-2 transmission in a real-world social network generated from high-resolution GPS data that were gathered in the course of a citizen-science experiment.</b></p> <p>-&gt; Tracing the contacts of contacts reduced the size of simulated outbreaks more than tracing of only contacts, but this strategy also resulted in almost half of the local population being quarantined at a single point in time.</p> <p>-&gt; Testing and releasing non-infectious individuals from quarantine led to increases in outbreak size, suggesting that contact tracing and quarantine might be most effective as a 'local lockdown' strategy when contact rates are high.</p> <p>-&gt; Combining physical distancing with contact tracing could enable epidemic control while reducing the number of quarantined individuals.</p> <p><b>Conclusion : Our findings suggest that targeted tracing and quarantine strategies would be most efficient when combined with other control measures such as physical distancing.</b></p>
Nature 05AUG2020	SARS-CoV-2 mRNA vaccine design enabled by prototype pathogen preparedness	Corbett K S. et al USA <a href="#">gotopaper</a>	Vaccine	<p><b>The release of SARS-CoV-2 sequences triggered immediate rapid manufacturing of an mRNA vaccine expressing the prefusion-stabilized SARS-CoV-2 spike trimer (mRNA-1273)</b></p> <p>Prior to vaccination of the first human subject, expression and antigenicity of the S-2P antigen delivered by mRNA was confirmed in vitro and immunogenicity of mRNA-1273 was documented:</p> <ul style="list-style-type: none"> <li>- mRNA-1273 induced dose-dependent S-specific binding antibodies after prime and boost in all mouse strains,</li> <li>- The level of pseudovirus neutralizing activity induced by 1 µg of mRNA-1273 in mice is similar in magnitude to that induced in human subjects by 100 µg</li> <li>- 1 µg of mRNA-1273 was sufficient to induce robust pseudovirus neutralizing activity and CD8 T cell responses, balanced Th1/Th2 antibody isotype responses, and protection from viral replication for more than 3 months following a prime/boost regimen</li> </ul> <p>Protection against SARS-CoV-2 infection in the lungs:</p> <ul style="list-style-type: none"> <li>- Efficacy of mRNA-1273 was dose-dependent, with two 0.1 µg mRNA-1273 doses reducing lung viral load by ~100-fold and two 0.01 µg mRNA-1273 doses reducing lung viral load by ~3-fold</li> </ul> <p><b>mRNA-1273 is currently in Phase 3 efficacy evaluation</b></p>
Cell 5AUG2020	Elevated calprotectin and abnormal myeloid cell subsets discriminate severe from mild COVID-19	Silvin et al., France <a href="#">gotopaper</a>	Immuno	<p><b>High dimensional flow cytometry and single cell RNA sequencing of COVID-19 patient peripheral blood cells.</b></p> <p>-&gt; Detection of the disappearance of non-classical CD14<sup>Low</sup>CD16<sup>High</sup> monocytes, the accumulation of HLA-DR<sup>Low</sup> classical monocytes, and the release of massive amounts of calprotectin (S100A8/S100A9) in severe cases.</p> <p>-&gt; Immature CD10<sup>Low</sup>CD101<sup>-</sup>CXCR4<sup>+</sup>/ - neutrophils with an immuno-suppressive profile accumulated as well in blood and lungs, suggesting emergency myelopoiesis</p> <p>-&gt; Calprotectin plasma level and a routine flow cytometry assay detecting decreased frequencies of non-classical monocytes could discriminate patients who develop a severe COVID-19 form, suggesting a predictive value that deserves prospective evaluation.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Science 04AUG2020	<b>Selective and cross-reactive SARS-CoV-2 T cell epitopes in unexposed humans</b>	Mateus et al., USA and Australia <a href="#">gotopaper</a>	Immuno	<p><b>SARS-CoV-2 reactive CD4+ T cells have been reported in unexposed individuals, suggesting pre-existing cross-reactive T cell memory in 20-50% of people.</b></p> <p>-&gt; We demonstrate a range of pre-existing memory CD4+ T cells that are cross-reactive with comparable affinity to SARS-CoV-2 and the common cold coronaviruses HCoV-OC43, HCoV-229E, HCoV-NL63, or HCoV-HKU1. Thus, variegated T cell memory to coronaviruses that cause the common cold may underlie at least some of the extensive heterogeneity observed in COVID-19 disease.</p> <p>-&gt; Variegated T cell memory to coronaviruses that cause the common cold may underlie at least some of the extensive heterogeneity observed in COVID-19 disease.</p>
Science 4AUG2020	<b>Engineering human ACE2 to optimize binding to the spike protein of SARS coronavirus 2</b>	Chan et al., USA <a href="#">gotopaper</a>	Therapeutic	<p>The spike protein S of SARS coronavirus 2 (SARS-CoV-2) binds ACE2 on host cells to initiate entry, and soluble ACE2 is a therapeutic candidate that neutralizes infection by acting as a decoy.</p> <p>Using deep mutagenesis, mutations in ACE2 that increase S binding are found across the interaction surface, in the N90-glycosylation motif and at buried sites. The mutational landscape provides a blueprint for understanding the specificity of the interaction between ACE2 and S and for engineering high affinity decoy receptors. Combining mutations gives ACE2 variants with affinities that rival monoclonal antibodies. A stable dimeric variant shows potent SARS-CoV-2 and -1 neutralization in vitro. The engineered receptor is catalytically active and its close similarity with the native receptor may limit the potential for viral escape.</p>
The Lancet Infectious Diseases 04AUG2020	<b>SeroTracker: a global SARS-CoV-2 seroprevalence dashboard</b>	Arora, R.K. et al., Canada <a href="#">gotopaper</a>	Public Health / Epidemiology	<p>Despite the value of antibody testing, there is no unified resource for seroprevalence estimates. The platform SeroTracker has therefore been created, a custom-built dashboard that systematically monitors and synthesises findings from global SARS-CoV-2 serological studies. The dashboard allows users to visualise seroprevalence estimates on a world map and compare estimates between regions, population groups, and testing modalities. SeroTracker integrates evidence from serosurveillance studies through a live systematic review of published articles (MEDLINE, Embase, Web of Science, and Cochrane), preprints (medRxiv and bioRxiv), government reports, and news articles. SeroTracker has proven useful to researchers, policy makers, and public health officials. SeroTracker will be hosted throughout the COVID-19 outbreak to support evidence-based decision making.</p> <p><b><a href="https://serotracker.com/Dashboard">https://serotracker.com/Dashboard</a></b></p>
Nature Medicine 03AUG2020	<b>Immune complement and coagulation dysfunction in adverse outcomes of SARS-CoV-2 infection</b>	Ramlall, V. et al., USA, <a href="#">gotopaper</a>	Public Health / Epidemiology	<p>Aim: retrospective observational study to determine whether conditions associated with dysregulated complement or coagulation systems impact Covid-19 disease.</p> <p>- <b>History of macular degeneration and history of coagulation disorders</b> are risk factors for SARS-CoV-2-associated morbidity and mortality, independently from age, sex or history of smoking.</p> <p>- In addition to type-I interferon and interleukin-6-dependent inflammatory responses, infection results in <b>robust engagement of the complement and coagulation pathways</b>.</p> <p>- Putative complement and coagulation-associated loci related to severe disease were identified, including missense, eQTL and sQTL variants of critical complement and coagulation regulators. These data provide evidence that <b>complement function modulates SARS-CoV-2 infection outcome, and point to putative transcriptional genetic markers</b> of susceptibility.</p> <p>The results highlight the value of using a multimodal analytical approach to reveal determinants and predictors of immunity, susceptibility and clinical outcome associated with infection.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Clinical infectious diseases 03AUG2020	<b>Longitudinal dynamics of the neutralizing antibody response to SARS-CoV-2 infection</b>	Wang, K. et al., China <a href="#">gotopaper</a>	Public Health / Epidemiology	<p><b>AIM: studying the longitudinal dynamics of SARS-CoV-2-specific neutralizing antibodies (NABs) and the levels of IgG and proinflammatory cytokines in COVID-19 patients, through analysis of 173 blood samples from 30 patients.</b></p> <ul style="list-style-type: none"> <li>- SARS-CoV-2-specific NAB titers were <b>low for the first 7-10 d</b> after symptom onset and <b>increased after 2-3 weeks</b>. The median peak time for NABs was 33 d after symptom onset.</li> <li>- NAB titers in 93.3% of the patients <b>declined gradually over the 3-month study period</b>, with a median decrease of 34.8%.</li> <li>- NAB titers increased over time in parallel with the <b>rise in IgG antibody levels</b>, correlating well at week 3.</li> <li>- The NAB titers also demonstrated a significant <b>positive correlation with levels of plasma proinflammatory cytokines</b>, including SCF, TRAIL, and M-CSF.</li> </ul>
Journal of Inf Dis 01AUG2020	<b>Prospective study comparing deep-throat saliva with other respiratory tract specimens in the diagnosis of novel coronavirus disease (COVID-19)</b>	Lai et al., China <a href="#">gotopaper</a>	Diagnostic	<p><b>Prospective study in two regional hospitals in Hong Kong</b></p> <ul style="list-style-type: none"> <li>- 563 serial samples collected during the virus shedding periods of 50 patients: 150 deep-throat saliva (DTS), 309 pooled-nasopharyngeal (NP) and throat swabs, and 104 sputum</li> <li>- DTS had the lowest overall RT-PCR positive rate (68.7% vs. 89.4% [sputum] and 80.9% [pooled NP and throat swabs]), and the lowest viral RNA concentration (mean log copy/mL 3.54 vs. 5.03 [sputum] and 4.63 [pooled NP and throat swabs])</li> <li>- Analyses with respect to time from symptom onset and severity also revealed similar results. Virus yield of DTS correlated with that of sputum (Pearson correlation index [95% CI]: 0.76 [0.62 – 0.86]). We estimated the overall false-negative rate of DTS could be 31.3%, and increased 2.7 times among patients without sputum.</li> </ul> <p><b>Conclusion : DTS produced the lowest viral RNA concentration and RT-PCR positive rate compared to conventional respiratory specimens in all phases of illness. Self-collect sputum should be the choice for patients with sputum.</b></p>
The Lancet Public Health 31JUL2020	<b>Risk of COVID-19 among front-line health-care workers and the general community: a prospective cohort study</b>	Nguyen, L.H. et al., USA / UK, <a href="#">gotopaper</a>	Public Health / Epidemiology	<p><b>Aim: assess risk of COVID-19 among front-line health-care workers compared with the general community and the effect of personal protective equipment (PPE) on risk, through a prospective, observational cohort study in the UK and the USA using self-reported data from the COVID Symptom Study smartphone application.</b></p> <ul style="list-style-type: none"> <li>- Among 2 035 395 community individuals and 99 795 front-line health-care workers, we recorded <b>5545 incident reports of a positive COVID-19 test</b> over 34 435 272 person-days.</li> <li>- Compared with the general community, <b>front-line health-care workers were at increased risk</b> for reporting a positive COVID-19 test.</li> <li>- Secondary and post-hoc analyses suggested <b>adequacy of PPE, clinical setting, and ethnic background</b> were also important factors.</li> </ul> <p>Health-care systems should ensure adequate availability of PPE and develop additional strategies to protect health-care workers from COVID-19, particularly those from Black, Asian, and minority ethnic backgrounds.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 30JUL2020	<b>Single-shot Ad26 vaccine protects against SARS-CoV-2 in rhesus macaques</b>	Mercadi N B et al USA <a href="#">gotopaper</a>	Vaccine	<p><b>A vaccine that requires only a single immunization would be optimal.</b> 52 rhesus macaques were immunized with Ad26 vectors encoding S variants or sham control and were challenged with SARS-CoV-2. Animals received a single immunization.</p> <p><b>Immunogenicity:</b> - Induced robust neutralizing antibody responses and provided complete or near-complete protection following SARS-CoV-2 challenge. Protective efficacy: - Vaccine-elicited neutralizing antibody titres correlated with protective efficacy, suggesting an immune correlate of protection.</p> <p><b>-&gt; Robust single-shot vaccine protection against SARS-CoV-2 in rhesus macaques.</b> <b>-&gt; Being evaluated in clinical trials</b></p>
SCIENCE 30JUL2020	<b>Adaptation of SARS-CoV-2 in BALB/c mice for testing vaccine efficacy</b>	Gu et al., China <a href="#">gotopaper</a>	Vaccine	<p><b>Clinical isolate of SARS-CoV-2 by serial passaging in the respiratory tract of aged BALB/c mice.</b></p> <p>-&gt; The resulting mouse-adapted strain at passage 6 (termed MASCP6) showed increased infectivity in mouse lung, and led to interstitial pneumonia and inflammatory responses in both young and aged mice following intranasal inoculation.</p> <p>-&gt; Deep sequencing revealed a panel of adaptive mutations potentially associated with the increased virulence. In particular, the N501Y mutation is located at the receptor binding domain (RBD) of the spike protein.</p> <p>-&gt; The protective efficacy of a recombinant RBD vaccine candidate was validated using this model.</p> <p><b>This mouse-adapted strain and associated challenge model should be of value in evaluating vaccines and antivirals against SARS-CoV-2.</b></p>
EBioMedicine 30JUL2020	<b>Serologic responses to SARS-CoV-2 infection among hospital staff with mild disease in eastern France</b>	Fafi-Kremer, S. et al., France, <a href="#">gotopaper</a>	Public Health / Epidemiology	<p>Aim: investigating the serologic response of 160 hospital staff who had recovered from mild forms of PCR-confirmed SARS-CoV-2 (no hospitalization required) using two assays for detection (rapid immunodiagnostic test and S-Flow assay), and an assay for neutralizing activity of the sera.</p> <p>- The median time from symptom onset to blood sample collection was 24. The <b>rapid immunodiagnostic test detected antibodies in 153 (95.6%)</b> of the samples and the <b>S-Flow assay in 159 (99.4%)</b>.</p> <p>- <b>Neutralizing antibodies (NABs) were detected in 79%, 92% and 98% of samples</b> collected 13–20, 21–27 and 28–41 days after symptom onset, respectively.</p> <p>This finding <b>supports the use of serologic testing for the diagnosis of individuals who have recovered from SARS-CoV-2 infection</b>. Future studies will help assess the persistence of the humoral response in recovered patients.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 29JUL2020	<b>Association Between Statewide School Closure and COVID-19 Incidence and Mortality in the US</b>	Auger et al., USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p><b>Analysis conducted between March 9, 2020, and May 7, 2020.</b></p> <p>-&gt; School closure was associated with a significant decline in both incidence of COVID-19 (adjusted relative change per week, -62%) and mortality (adjusted relative change per week, -58%).</p> <p>-&gt; In a model derived from this analysis, it was estimated that closing schools when the cumulative incidence of COVID-19 was in the lowest quartile compared with the highest quartile was associated with 128.7 fewer cases per 100 000 population over 26 days and with 1.5 fewer deaths per 100 000 population over 16 days.</p>
Nature 29JUL2020	<b>Papain-like protease regulates SARS-CoV-2 viral spread and innate immunity</b>	Shin et al., Germany <a href="#">gotopaper</a>	Therapeutic	<p><b>Papain-like protease PLpro : an essential coronavirus enzyme required for processing viral polypeptides to generate a functional replicase complex and enable viral spread</b></p> <p>-&gt; Biochemical, structural and functional characterization of the SARS-CoV-2 PLpro (SCoV2-PLpro) and outline differences to SARS-CoV PLpro (SCoV-PLpro) in controlling host interferon (IFN) and NF-κB pathways</p> <p>-&gt; Upon infection, SCoV2-PLpro contributes to the cleavage of ISG15 from interferon responsive factor 3 (IRF3) and attenuates type I interferon responses.</p> <p>-&gt; Inhibition of SCoV2-PLpro with GRL-0617 impairs the virus-induced cytopathogenic effect, fosters the anti-viral interferon pathway and reduces viral replication in infected cells.</p> <p><b>These results highlight a dual therapeutic strategy in which targeting of SCoV2-PLpro can suppress SARS-CoV-2 infection and promote anti-viral immunity.</b></p>
Nature 29JUL2020	<b>SARS-CoV-2-reactive T cells in healthy donors and patients with COVID-19</b>	Braun, J. et al, Germany, <a href="#">gotopaper</a>	Public Health / Epidemiology	<p>Aim: investigating <b>SARS-CoV-2 spike glycoprotein (S)-reactive CD4(+) T cells</b> in peripheral blood of patients with COVID-19 and SARS-CoV-2-unexposed healthy donors (HD).</p> <p>- SARS-CoV-2 S-reactive CD4(+) T cells were detected in <b>83% of patients with COVID-19</b> but also in <b>35% of HD</b>. S-reactive CD4(+) T cells in HD reacted primarily to C-terminal S epitopes, which show a higher homology to spike glycoproteins of human endemic coronaviruses, compared to N-terminal epitopes.</p> <p>- S-reactive T cell lines generated from SARS-CoV-2-naïve HD <b>responded similarly to C-terminal S of human endemic coronaviruses 229E and OC43 and SARS-CoV-2</b>, demonstrating the presence of S-cross-reactive T cells, probably generated during past encounters with endemic coronaviruses.</p> <p>The role of pre-existing SARS-CoV-2 cross-reactive T cells for clinical outcomes remains to be determined in larger cohorts. However, their presence in the general population may affect the dynamics of the current pandemic.</p>
Nature 29JUL2020	<b>Association of COVID-19 inflammation with activation of the C5a-C5aR1 axis</b>	Carvelli, J. et al, France, <a href="#">gotopaper</a>	Public Health / Epidemiology	<p>Aim: longitudinal analysis of immune responses linked to soluble factor C5a and cells expressing its receptor C5aR1 (CD88) in the blood and broncho-alveolar lavage fluid of patients at various stages of COVID-19 severity.</p> <p>- <b>An increase in soluble C5a levels proportional to COVID-19 severity and high levels of C5aR1 expression in blood and pulmonary myeloid cells</b> is reported, supporting a role for the C5a-C5aR1 axis in the pathophysiology of acute respiratory distress syndrome (ARDS).</p> <p>- <b>Anti-C5aR1 therapeutic monoclonal antibodies prevented C5a-mediated human myeloid cell recruitment and activation</b>, and inhibited acute lung injury in human C5aR1 knockin mice.</p> <p>These results suggest that C5a-C5aR1 axis blockade might be used to limit myeloid cell infiltration in damaged organs and prevent excessive lung inflammation associated with ARDS in COVID-19 patients.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
NEJM 28JUL2020	<b>Evaluation of the mRNA-1273 Vaccine against SARS-CoV-2 in Nonhuman Primates</b>	Kizzmekia S. Corbett et al., USA <a href="#">gotopaper</a>	Vaccine	<p><b>Method:</b> Nonhuman primates received 10 or 100 µg of mRNA-1273, a vaccine encoding the prefusion-stabilized spike protein of SARS-CoV-2, or no vaccine. Antibody and T-cell responses were assessed before upper- and lower-airway challenge with SARS-CoV-2. Active viral replication and viral genomes in bronchoalveolar-lavage (BAL) fluid and nasal swab specimens were assessed by PCR, and histopathological analysis and viral quantification were performed on lung-tissue specimens.</p> <p><b>Results:</b> - The mRNA-1273 vaccine candidate induced antibody levels exceeding those in human convalescent-phase serum, with live-virus reciprocal 50% inhibitory dilution (ID50) geometric mean titers of 501 in the 10-µg dose group and 3481 in the 100-µg dose group.-- Vaccination induced type 1 helper T-cell (Th1)–biased CD4 T-cell responses and low or undetectable Th2 or CD8 T-cell responses. - Viral replication was not detectable in BAL fluid by day 2 after challenge in seven of eight animals in both vaccinated groups. No viral replication was detectable in the nose of any of the eight animals in the 100-µg dose group by day 2 after challenge - Limited inflammation or detectable viral genome or antigen was noted in lungs of animals in either vaccine group.</p> <p><b>Conclusion:</b> The results reported here provide data on mRNA-1273 immunogenicity and protection of the upper and lower airways in nonhuman primates that complement the immunogenicity and safety data established by a phase 1 clinical study involving humans. Studies are now under way to determine the durability of immunity and protection over 1 year after vaccination.</p>
Cell Systems 27JUL2020	<b>Computationally Optimized SARS-CoV-2 MHC Class I and II Vaccine Formulations Predicted to Target Human Haplotype Distributions</b>	Ge Liu et al., USA <a href="#">gotopaper</a>	Vaccine	<p><b>Aim</b> Validation of combinatorial machine learning method to evaluate and optimize peptide vaccine formulations for SARS-CoV-2</p> <p><b>Method</b> By testing human HLA haplotype frequencies of three major populations and using recent advances in machine learning, the authors have evaluated completed designs models to provide a conservative evaluation of vaccine peptide presentation.</p> <p><b>Results:</b> - SARS-CoV-2 MHC class I vaccine formulations provide 93.21% predicted population coverage with at least five vaccine peptide-HLA average hits per person (<math>\geq 1</math> peptide: 99.91%), - MHC class II vaccine formulations provide 97.21% predicted coverage with at least five vaccine peptide-HLA average hits per person with all peptides having an observed mutation probability of <math>\leq 0.001</math>. - OptiVax can be used to augment S protein vaccine designs to increase their population coverage.</p> <p><b>Conclusion</b> EvalVax can be used for vaccine designs that are focused on the expression of entire viral proteins or their subunits to evaluate the level of viral peptide-HLA presentation.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 27JUL2020	Longitudinal analyses reveal immunological misfiring in severe COVID-19	Lucas, Carolina et al., USA <a href="#">gotopaper</a>	Immunology	<p>Immune profiling in 113 COVID-19 patients with moderate (non-ICU) and severe (ICU) disease revealed:</p> <ul style="list-style-type: none"> <li>- <b>Association between early elevated cytokines and worse disease outcomes.</b></li> </ul> <p>Following early increase in cytokines:</p> <ul style="list-style-type: none"> <li>- <b>moderate disease COVID-19 patients displayed a progressive reduction in type-1 (antiviral) and type-3 (antifungal) responses.</b></li> <li>- <b>severe disease patient maintained elevated type-1 and 3 responses throughout the course of disease, accompanied by increase in multiple type 2 (anti-helminths) effectors (IL-5, IL-13, IgE, eosinophils).</b></li> </ul> <p>-&gt; <b>Identified 4 immune signatures correlating with distinct disease trajectories of patients:</b> (A) enriched growth factors, (B) type-2/3 cytokines, (C) mixed type-1/2/3 cytokines, and (D) chemokines.</p> <p>-&gt; <b>Patients who recovered with moderate disease = immune signature A (enriched tissue reparative growth factors).</b></p> <p>-&gt; <b>Patients with worsened disease trajectory = elevated levels of all 4 signatures.</b></p>
Nature 24JUL2020	Discovery of SARS-CoV-2 antiviral drugs through large-scale compound repurposing	Riva, Laura et al. USA/China <a href="#">gotopaper</a>	Therapeutics	<p><b>High-throughput reprofiling screen</b> using the ReFRAME (Repurposing, Focused Rescue, and Accelerated Medchem) drug library, a <b>comprehensive open-access library of ~12,000</b> that have been either <b>FDA-approved or registered outside the US, entered clinical trials, or undergone significant pre-clinical characterization</b>, to identify existing drugs that harbor <b>antiviral activity against SARS-CoV-2 in a cell-based assay.</b></p> <p><b>Results:</b> Identification of <b>100 molecules that inhibit SARS-CoV-2 replication in mammalian cells</b>, including 21 known drugs that exhibit <b>dose response relationships</b>. Of these, 13 were found to harbor effective concentrations likely commensurate with <b>achievable therapeutic doses in patients</b>, including the PIKfyve kinase inhibitor apilimod2–4, and the cysteine protease inhibitors MDL-28170, Z LVG CHN2, VBY-825, and ONO 5334. Notably, MDL-28170, ONO 5334, and apilimod were found to antagonize viral replication in <b>human iPSC-derived pneumocyte-like cells</b>, and the PIKfyve inhibitor also demonstrated antiviral efficacy in a <b>primary human lung explant model</b>.</p> <p>Publicly disclosed and relevant preclinical and clinical properties of the most advanced among these molecules will enable prioritization of known drugs for in vivo preclinical and clinical evaluation for the treatment of SARS-CoV-2.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature Communications 24JUL2020	<b>Structural basis of RNA cap modification by SARS-CoV-2</b>	Viswanathan, Thiruselvam et al USA <a href="#">gotopaper</a>	Therapeutics	<p>Report of the high-resolution structure of a ternary complex of SARS-CoV-2 nsp16 and nsp10 in the presence of cognate RNA substrate analogue and methyl donor, S-adenosyl methionine (SAM).</p> <p><b>Results</b></p> <ul style="list-style-type: none"> <li>- This enzyme is specifically adapted to bind and methylate the RNA cap.</li> <li>- The structure provides a snapshot of pre-catalytic state of methyl transfer from SAM to 2'-OH of ribose of the first transcribing nucleotide of the mRNA cap.</li> <li>- Large conformational changes associated with substrate binding are observed as the enzyme transitions from a binary to a ternary state. This induced fit model provides mechanistic insights into the 2'-O methylation of the viral mRNA cap.</li> <li>- A distant (25 Å) ligand-binding site unique to SARS-CoV-2 was discovered in nsp16 with distinct capability to accommodate small molecule ligands. It could alternatively be targeted, in addition to RNA cap and SAM pockets, for antiviral development.</li> <li>- The acquired mutations in SARS-CoV-2 nsp16 were mapped. One of these mutation hotspots showed high frequency in COVID-19 strains associated with New York City outbreak.</li> </ul> <p><b>Conclusion</b></p> <p>Solid framework from which therapeutic modalities may be designed by targeting different ligand-binding sites of nsp16, including RNA cap and SAM pockets, for the treatment of COVID-19 and emerging coronavirus illnesses.</p>
Clinical infectious diseases 24JUL2020	<b>Remdesivir for Severe COVID-19 versus a Cohort Receiving Standard of Care</b>	Olender, Susan A. et al. <a href="#">gotopaper</a>	Therapeutics	<p><b>Comparison of the efficacy of remdesivir versus standard-of-care</b> treatment in adults with <b>severe COVID-19</b> using data from a phase 3 remdesivir trial (GS-US-540-5773) and a <b>retrospective cohort</b> of patients with severe COVID-19 <b>treated with standard-of-care</b> (GS-US-540-5807), using the stabilized inverse probability of treatment weighting (IPTW) method. Inclusion criteria were similar between studies: confirmed SARS-CoV-2 infection, hospitalized patients, with oxygen saturation 94% or lower on room air or required supplemental oxygen, and pulmonary infiltrates.</p> <p><b>Primary endpoint:</b> proportion of patients with <b>recovery on day 14</b>, dichotomized from a 7-point clinical status ordinal scale. A key secondary endpoint was mortality.</p> <p><b>Results</b></p> <p><b>312 and 818 patients</b> were counted in the remdesivir- and non-remdesivir-cohorts, respectively.</p> <p><b>At day 14, 74.4% of patients in the remdesivir-cohort had recovered versus 59.0% in the non-remdesivir-cohort</b> (adjusted odds ratio 2.03: 95% confidence interval 1.34–3.08, p&lt;0.001).</p> <p><b>At day 14, 7.6% of patients in the remdesivir-cohort had died versus 12.5% in the non-remdesivir-cohort</b> (adjusted odds ratio 0.38, 95% confidence interval: 0.22–0.68, p=0.001).</p> <p><b>Conclusions</b></p> <p>In this comparative analysis, by day 14, <b>remdesivir was associated with significantly greater recovery and 62% reduced odds of death versus standard-of-care treatment in patients with severe COVID-19.</b></p> <p><b>Limitations</b></p> <p>Comparison was not randomized ; open-label design of Study 5773</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Cell 23JUL2020	<b>A thermostable mRNA vaccine against COVID-19</b>	Zhang, Na-Na et al. China <a href="#">gotopaper</a>	Vaccine	<p>Development of a lipid-nanoparticle-encapsulated mRNA (mRNA-LNP) encoding the receptor binding domain (RBD) of SARS-CoV-2 as a vaccine candidate (ARCoV).</p> <p><b>Results:</b></p> <ul style="list-style-type: none"> <li>- Intramuscular immunization of ARCoV mRNA-LNPs elicited robust neutralizing antibodies against SARS-CoV-2 as well as Th1-biased cellular response in mice and non-human primates.</li> <li>- Two doses of ARCoV immunization in mice conferred complete protection against the challenge of a SARS-CoV-2 mouse adapted strain.</li> <li>- ARCoV was manufactured in liquid formulation and can be stored at room temperature for at least one week.</li> </ul> <p><b>Conclusion:</b></p> <p>Thermostable mRNA vaccine candidate against SARS-CoV-2, with first line evidence of immunogenicity and efficacy in multiple animal models, and currently being evaluated in phase 1 clinical trials.</p>
Science 23JUL2020	<b>Structural basis for neutralization of SARS-CoV-2 and SARS-CoV by a potent therapeutic antibody</b>	Lv, Zhe et al. China <a href="#">gotopaper</a>	Vaccine/ Therapeutics	<p>The RBDs of SARS-CoV and SARS-CoV-2 have an amino-acid sequence identity of around 75%, raising the possibility that RBD-targeting cross-neutralizing NABs could be possibly identified. Using phage display technique, an antibody library was generated from RNAs extracted from peripheral lymphocytes of mice immunized with recombinant SARS-CoV RBD. SARS-CoV-2 RBD was used as the target for screening the phage antibody library for potential hits.</p> <p><b>Results:</b></p> <ul style="list-style-type: none"> <li>- Humanized monoclonal antibody, H014, efficiently neutralizes SARS-CoV-2 and SARS-CoV pseudoviruses as well as authentic SARS-CoV-2 at nM level by engaging the S receptor binding domain (RBD).</li> <li>- H014 administration reduced SARS-CoV-2 titers in the infected lungs and prevented pulmonary pathology in hACE2 mouse model.</li> <li>- Cryo-EM characterization of the SARS-CoV-2 S trimer in complex with the H014 Fab fragment unveiled a novel conformational epitope, which is only accessible when the RBD is in open conformation.</li> <li>- Biochemical, cellular, virological and structural studies demonstrated that H014 prevents attachment of SARS-CoV-2 to its host cell receptors.</li> </ul> <p><b>Conclusion:</b></p> <p>The molecular features of H014 epitopes facilitate the discovery of broad cross-neutralizing epitopes within lineage B and pose interesting targets for structure-based rational vaccine design, as well as for the promise of antibody-based therapeutic interventions for the treatment of COVID-19.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
NEJM 23JUL2020	<b>Hydroxychloroquine with or without Azithromycin in Mild-to-Moderate Covid-19</b>	Cavalcanti, Alexandre B. et al., Brazil <a href="#">gotopaper</a>	Therapeutics	<p><b>Multicenter, randomized, open-label, 3-group, controlled trial</b> in Brazil involving <b>hospitalized patients with suspected or confirmed Covid-19</b> who were receiving either <b>no supplemental oxygen or a maximum of 4 liters per minute of supplemental oxygen</b>. Patients were randomly assigned in a 1:1:1 ratio to receive <b>standard care</b>, <b>standard care + HCQ</b> (400 mg twice daily), or <b>standard care + HCQ (400 mg twice daily) + azithromycin</b> (500 mg once daily) for 7 days.</p> <p><b>Primary outcome:</b> clinical status at 15 days as assessed with the use of a seven-level ordinal scale in the modified intention-to-treat population (patients with a confirmed diagnosis of Covid-19). Safety was also assessed.</p> <p>A total of 667 patients underwent randomization; <b>504 patients had confirmed Covid-19 and were included in the modified intention-to-treat analysis</b>.</p> <p>- As compared with standard care, the proportional odds of having a higher score on the seven-point ordinal scale at 15 days was <b>not affected by either hydroxychloroquine alone</b> (odds ratio, 1.21; 95% confidence interval [CI], 0.69 to 2.11; P=1.00) <b>or hydroxychloroquine plus azithromycin</b> (odds ratio, 0.99; 95% CI, 0.57 to 1.73; P=1.00). - Prolongation of the corrected QT interval and elevation of liver-enzyme levels were <b>more frequent in patients receiving hydroxychloroquine, alone or with azithromycin</b>, than in those who were not receiving either agent.</p> <p><b>=&gt; Among patients hospitalized with mild-to-moderate Covid-19, the use of hydroxychloroquine, alone or with azithromycin, did not improve clinical status at 15 days as compared with standard care.</b></p>
Science 23JUL2020	<b>Evolution and epidemic spread of SARS-CoV-2 in Brazil</b>	Candido, Darlan S. et al. UK-Brazil <a href="#">gotopaper</a>	Public Health/ Epidemiology	<p>Brazil currently has one of the fastest growing SARS-CoV-2 epidemics in the world.</p> <p>- A mobility-driven transmission model showed non-pharmaceutical interventions reduced reproduction number from &gt;3 to 1–1.6 in São Paulo and Rio de Janeiro.</p> <p>Sequencing of 427 new genomes identified:</p> <ul style="list-style-type: none"> <li>- <b>Over 100 international virus introductions in Brazil.</b></li> <li>- <b>Most (76%) of Brazilian strains fell in 3 clades introduced from Europe</b> between 22 February and 11 March 2020.</li> <li>- During the <b>early epidemic phase, SARS-CoV-2 spread mostly locally</b> and within-state borders. <b>After this period, despite sharp decreases in air travel, multiple exportations from large urban centers coincided with a 25% increase in average travelled distances in national flights.</b></li> </ul> <p>-&gt; Study sheds light on the epidemic transmission and evolutionary trajectories of SARS-CoV-2 lineages in Brazil, and provide <b>evidence that current interventions remain insufficient to keep virus transmission under control in the country.</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature structural & molecular biology 22JUL2020	<b>Controlling the SARS-CoV-2 spike glycoprotein conformation</b>	Henderson, Rory et al. USA <a href="#">gotopaper</a>	Vaccine	<p>To better understand S-protein mobility, a structure-based vector analysis of available <math>\beta</math>-CoV S-protein structures was implemented.</p> <ul style="list-style-type: none"> <li>- S-proteins from different <math>\beta</math>-CoVs display distinct configurations.</li> <li>- Two soluble ectodomain constructs for the SARS-CoV-2 S-protein were developed, in which the highly immunogenic and mobile receptor binding domain (RBD) is either locked in the all-RBDs 'down' position or adopts 'up' state conformations more readily than the wild-type S-protein.</li> </ul> <p>=&gt; The conformation of the S-protein can be controlled via rational design and can provide a framework for the development of engineered CoV S-proteins for vaccine applications.</p>
Nature 22JUL2020	<b>Hydroxychloroquine use against SARS-CoV-2 infection in non-human primates</b>	Maisonnasse, Pauline et al. France <a href="#">gotopaper</a>	Therapeutics	<p>Evaluation of the <b>antiviral activity of HCQ both in vitro and in SARS-CoV-2-infected macaques.</b></p> <ul style="list-style-type: none"> <li>- HCQ showed <b>antiviral activity in African green monkey kidney cells (VeroE6)</b> but <b>not in a model of reconstituted human airway epithelium.</b></li> <li>- <b>Cynomolgus macaques</b> provides a <b>relevant model</b> for studying the <b>early stages of SARS-Cov-2 infection in humans.</b></li> <li>- <b>In macaques, no antiviral activity nor clinical efficacy of HCQ treatment, alone or in combination with azithromycin (AZTH), was shown, regardless of the timing of treatment initiation,</b> either before infection, early after infection (before viral load peak) or late after infection (after viral load peak). This was in spite of high HCQ concentration in blood and lung and plasma exposure similar to that observed in COVID-19 patients.</li> <li>- When the drug was used as a <b>pre-exposure prophylaxis (PrEP), HCQ did not confer protection against acquisition of infection.</b></li> </ul> <p>=&gt; <b>These findings do not support the use of HCQ, either alone or in combination with AZTH, as an antiviral treatment for COVID-19 in humans.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 22JUL2020	<b>Potent neutralizing antibodies directed to multiple epitopes on SARS-CoV-2 spike</b>	Liu, Lihong et al. USA/China <a href="#">gotopaper</a>	Therapeutics	<p>Isolation of 61 <b>SARS-CoV-2-neutralizing monoclonal antibodies</b> from 5 infected patients hospitalized with severe disease. Among these are <b>19 antibodies that potently neutralized the authentic SARS-CoV-2 in vitro</b>, 9 of which exhibited <b>exquisite potency</b>, with 50% virus-inhibitory concentrations of 0.7 to 9 ng/mL.</p> <p>Epitope mapping showed this collection of 19 antibodies to be about <b>equally divided between those directed to the receptor-binding domain (RBD) and those to the N-terminal domain (NTD)</b>, indicating that both of these <b>regions at the top of the viral spike are immunogenic</b>.</p> <p>In addition, 2 other powerful neutralizing antibodies <b>recognized quaternary epitopes that overlap with the domains at the top of the spike</b>.</p> <p><b>Cryo-electron microscopy reconstructions</b> of one antibody targeting RBD, a second targeting NTD, and a third bridging two separate RBDs revealed <b>recognition of the closed, “all RBD-down” conformation of the spike</b>.</p> <p>=&gt; Collection of SARS-CoV-2-neutralizing mAbs that are <b>not only potent but also diverse</b>. The potency and diversity of our SARS-CoV-2-neutralizing mAbs are <b>likely attributable to patient selection</b> (infected individuals with severe disease developed a more robust virus-neutralizing antibody response). The diversity of these antibodies is also attributable, in part, to the choice of <b>using the S trimer to sort from memory B cells</b>, while most groups focused on the use of RBD. RBD and NTD are, no doubt, quite immunogenic. <b>Neutralizing antibodies to the stem region of the S trimer remain to be discovered</b>.</p>
Nature 22JUL2020	<b>Chloroquine does not inhibit infection of human lung cells with SARS-CoV-2</b>	Hoffmann, Markus et al. Germany <a href="#">gotopaper</a>	Therapeutics	<p>Comparison of <b>chloroquine and hydroxychloroquine-mediated inhibition of SARS-2-S-mediated entry into Vero (kidney), Vero-TMPRSS2 and Calu-3 (lung) cells</b>.</p> <p>Calu-3 cells, like airway epithelium, express low amounts of cathepsin L (CatL) and SARS-CoV-2 entry into these cells is TMPRSS2-dependent. In contrast, Vero cell entry of SARS-CoV-2 is CatL-dependent while both CatL and TMPRSS2 support entry into Vero-TMPRSS2 cells. As control, camostat mesylate, which inhibits TMPRSS2-dependent entry, was used.</p> <ul style="list-style-type: none"> <li>- Chloroquine and hydroxychloroquine <b>can block SARS-2-S-driven entry</b> but <b>inhibition is cell line-dependent</b> and <b>efficient inhibition is not observed with TMPRSS2+ lung cells</b>.</li> <li>- Chloroquine efficiently <b>blocked SARS-CoV-2 infection of Vero kidney cells</b>, as expected, but <b>failed to efficiently inhibit SARS-CoV-2 infection of Calu-3 lung cells</b>.</li> <li>- Chloroquine <b>failed to efficiently block Calu-3 cell infection with SARS-2-S-bearing pseudotypes and authentic SARS-CoV-2</b>, indicating that in these cells chloroquine does not appreciably interfere with viral entry or the subsequent steps of the viral replication cycle.</li> </ul> <p>Confirmation of the results with primary respiratory epithelium is pending.</p> <p>=&gt; These results indicate that <b>chloroquine targets a pathway for viral activation that is not operative in lung cells and is unlikely to protect against SARS-CoV-2 spread in and between patients</b>.</p> <p>=&gt; Moreover, they highlight that <b>cell lines mimicking important aspects of respiratory epithelial cells should be used when analyzing the antiviral activity of drugs targeting host cell functions</b>.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Cell Reports Medicine 21JUL2020	<b>Characterization and Treatment of SARS-CoV-2 in Nasal and Bronchial Human Airway Epithelia</b>	Pizzorno, Andrés et al. France <a href="#">gotopaper</a>	Therapeutics	<p>Use of <b>reconstituted human airway epithelia</b> to <b>isolate</b> and then <b>characterize</b> the <b>viral infection kinetics</b>, <b>tissue-level remodeling of the cellular ultrastructure</b>, and <b>transcriptional early immune signatures</b> induced by SARS-CoV-2 in a physiologically relevant model.</p> <p>Results emphasize <b>distinctive transcriptional immune signatures between nasal and bronchial HAE</b>, both in terms of kinetics and intensity, hence suggesting <b>putative intrinsic differences in the early response to SARS-CoV-2 infection</b>.</p> <p>Most important, <b>evidence in human-derived tissues was provided on the antiviral efficacy of remdesivir monotherapy</b> and the <b>potential of the remdesivir-diltiazem combination</b> was explored as an option worthy of further investigation to respond to the still-unmet COVID-19 medical need.</p> <p>=&gt; Results expected to provide a benchmark for future studies aimed at further characterizing the local pathophysiology and immune response to SARS-CoV-2 infection, particularly in the lower respiratory tract, with the ultimate objective of <b>providing insight in terms of putative prognostic biomarkers and/or patient management</b>.</p> <p>=&gt; The <b>HAE model of SARS-CoV-2 infection</b> described in this study also constitutes an <b>advantageous physiologic model to evaluate candidate therapeutic approaches</b>, provided that in many cases the inhibitory effects observed in classic reductionist models of immortalized cell lines do not necessarily translate into a real clinical setting.</p>
Science 21JUL2020	<b>Serial interval of SARS-CoV-2 was shortened over time by nonpharmaceutical interventions</b>	Ali, Sheikh Taslim et al. UK-France-USA-China <a href="#">gotopaper</a>	Public Health/ Epidemiology	<p>A database of 677 COVID-19 transmission pairs (in mainland China) (symptom onset dates and social relationships available for both the infector and infectee) show that:</p> <ul style="list-style-type: none"> <li>- <b>Mean serial intervals of COVID-19 have shortened substantially from 7.8 days to 2.6 days within a month (January 9 to February 13, 2020). This change is driven by enhanced non-pharmaceutical interventions, in particular reduction of case isolation delay period.</b></li> <li>- Using <b>real-time estimation of serial intervals allowing for variation over time, provides more accurate estimates of reproduction numbers</b> than using conventionally fixed serial interval distributions.</li> </ul> <p>-&gt; Findings could improve assessment of transmission dynamics, forecasting future incidence, and estimating the impact of control measures.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
PLoS Med 21JUL2020	<b>Impact of self-imposed prevention measures and short-term government-imposed social distancing on mitigating and delaying a COVID-19 epidemic: A modelling study</b>	Teslya, Alexandra et al. Netherlands-Portugal <a href="#">gotopaper</a>	Public Health/ Epidemiology	<p>Effectiveness of self-imposed measures and short-term government-imposed social distancing in mitigating COVID-19 epidemic (transmission model in population stratified by disease status and disease awareness status). Principles: Self-imposed measures assumed in disease-aware individuals (handwashing, mask-wearing, social distancing). Government-imposed social distancing reduced contact rate of individuals irrespective of disease or awareness status.</p> <p>- <b>For fast awareness spread in population</b>, self-imposed measures can significantly reduce the attack rate and diminish and postpone the peak number of diagnoses. <b>Estimated: large epidemic preventable if efficacy of these measures exceeds 50%.</b></p> <p>- <b>For slow awareness spread</b>, self-imposed measures <b>reduce peak number of diagnoses and attack rate but do not affect the timing of the peak.</b></p> <p>- <b>Early implementation of short-term government-imposed social distancing alone is estimated to delay</b> (by at most 7 months for a 3-month intervention) <b>but not reduce the peak. Delay can be longer and height of peak further reduced if combined with self-imposed measures that continue after government-imposed distancing is lifted.</b></p> <p>Limitations: analyses do not account for stochasticity, demographics, heterogeneities in contact patterns or mixing, spatial effects, imperfect isolation of individuals with severe disease, and reinfection.</p> <p>-&gt; <b>Suggests that information dissemination and disease awareness, which mobilise people to adopt effective self-imposed control measures, can be an effective strategy to mitigate and delay the epidemic.</b></p> <p>-&gt; Early initiated short-term government-imposed social distancing can buy extra time.</p>
Science Translational Medicine 20JUL2020	<b>An alphavirus-derived replicon RNA vaccine induces SARS-CoV-2 neutralizing antibody and T cell responses in mice and nonhuman primates</b>	Erasmus, Jesse H et al. USA <a href="#">gotopaper</a>	Vaccine	<p>Development of an alphavirus-derived replicon RNA vaccine candidate, repRNA-CoV2S, encoding the SARS-CoV-2 spike (S) protein. The RNA replicons were formulated with Lipid InOrganic Nanoparticles (LION) that were designed to enhance vaccine stability, delivery, and immunogenicity.</p> <p><b>Results:</b></p> <p>- A single intramuscular injection of the LION/repRNA-CoV2S vaccine in mice elicited robust production of anti-SARS-CoV-2 S protein IgG antibody isotypes indicative of a Type 1 T helper cell response.</p> <p>- A prime/boost regimen induced potent T cell responses in mice including antigen-specific responses in lung and spleen. Prime-only immunization of aged (17-month old) mice induced smaller immune responses compared to young mice, but this difference was abrogated by booster immunization.</p> <p>- In nonhuman primates, prime-only immunization in one intramuscular injection site or prime/boost immunizations in 5 intramuscular injection sites elicited modest T cell responses and robust antibody responses. The antibody responses persisted for at least 70 days and neutralized SARS-CoV-2 at titers comparable to those in human serum samples collected from individuals convalescing from COVID-19.</p> <p><b>Conclusion:</b></p> <p>Potential for LION/repRNA-CoV2S, which will enter clinical development under the name HDT-301, to induce rapid immune protection from SARS-CoV-2 infection.</p> <p><b>Limitations:</b></p> <p>Absence of challenge data ; limited number of animals ; time points after vaccination where observed responses, especially T-cell responses, exhibited great variability</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Gut 20JUL2020	Depicting SARS-CoV-2 faecal viral activity in association with gut microbiota composition in patients with COVID-19	Zuo, Tao et al. China <a href="#">gotopaper</a>	Gut Microbiota	<p>Longitudinal faecal microbiome alterations in patients with COVID-19 (RNA metagenomics sequencing on faecal viral extractions from 15 hospitalised patients with COVID-19):</p> <ul style="list-style-type: none"> <li>- <b>7 of 15 COVID-19 patients (46.7%) had SARS-CoV-2 positive stool</b> (viral RNA). Even in absence of GI manifestations, all 7 patients showed strikingly <b>higher 3' vs 5' end SARS-CoV-2 genome coverage and density</b> in their faecal viral metagenome profile.</li> <li>- 3 patients continued to display active fecal viral infection signature up to 6 days after clearance of SARS-CoV-2 from respiratory samples.</li> <li>- Faecal samples with <b>high SARS-CoV-2 infectivity signature -&gt; higher abundances of bacterial species <i>Collinsella aerofaciens</i>, <i>Collinsella tanakaei</i>, <i>Streptococcus infantis</i>, <i>Morganella morganii</i>, and higher capacity for nucleotide <i>de novo</i> biosynthesis</b>, amino acid biosynthesis and glycolysis.</li> <li>- Faecal samples with <b>low-to-no SARS-CoV-2 infectivity signature -&gt; higher abundances of short-chain fatty acid producing bacteria</b>, <i>Parabacteroides merdae</i>, <i>Bacteroides stercoris</i>, <i>Alistipes onderdonkii</i> and <i>Lachnospiraceae bacterium</i>.</li> </ul> <p>-&gt; Gut microbiota of patients with active SARS-CoV-2 GI infection was characterised by enrichment of opportunistic pathogens, loss of salutary bacteria and increased functional capacity for nucleotide and amino acid biosynthesis and carbohydrate metabolism.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
LANCET 20JUL2020	Immunogenicity and safety of a recombinant adenovirus type-5-vectored COVID-19 vaccine in healthy adults aged 18 years or older: a randomised, double-blind, placebo controlled, phase 2 trial	Feng-Cai Zhu et al.- China <a href="#">gotopaper</a>	Vaccine	<p><b>PROTOCOLE DESIGN (NCT04341389):</b></p> <ul style="list-style-type: none"> <li>&gt; Vaccine: Ad5-vectored COVID 19 vaccine from CanSino</li> <li>&gt; Phase 2 clinical trial; randomized, double blinded, placebo controlled.</li> <li>&gt; 508 participants &gt; 18 years of age (seronegative for SARS-CoV-2 infection). Recruited in one single site (Wuhan)</li> <li>&gt; Doses: either 1x10exp11 vp/mL, 5x10exp10vp/mL or placebo (2:1:1). One single administration IM</li> </ul> <p><b>Primary endpoints:</b></p> <ul style="list-style-type: none"> <li>&gt; Geometric mean titers of specific ELISA antibody responses to the receptor binding domain (RBD)</li> <li>&gt; Neutralising antibody responses at day 28</li> <li>&gt; incidence of adverse reactions within 14 days.</li> </ul> <p><b>RESULTS:</b></p> <p><b>Cohort characteristics:</b> 50% male; mean age 39.7 years, SD 12.5)</p> <p><b>RDB specific antibodies induction et seroconversion rates at day 28:</b></p> <ul style="list-style-type: none"> <li>&gt; Dose 1x10exp11 vp/mL: 656,5 (95% CI 575,2–749,2); 96% (95% CI 93–98)</li> <li>&gt; Dose 5x10exp10vp/mL: 571,0 (467,6–697,3); 97% (92–99)</li> </ul> <p><b>Neutralizing antibodies induction:</b></p> <ul style="list-style-type: none"> <li>&gt; Dose 1x10exp11 vp/mL: GMTs of 19,5 (95% CI 16,8–22,7)</li> <li>&gt; Dose 5x10exp10vp/mL: GMTs of 18,3 (14,4–23,3)</li> </ul> <p><b>Specific interferon <math>\gamma</math> enzyme-linked immunospot assay responses post vaccination observed in:</b></p> <ul style="list-style-type: none"> <li>&gt; 227/253 participants (90%, 95% CI 85–93) (Dose 1x10exp11vp/mL)</li> <li>&gt; 113/129 participants (88%, 81–92) (Dose 5x10exp10vp/mL)</li> </ul> <p><b>Solicited adverse reactions were reported by:</b></p> <ul style="list-style-type: none"> <li>&gt; 183/253 participants (72%) (Dose 1x10exp11 vp/mL)</li> <li>&gt; 96/129 participants (74%) (Dose 5x10exp10vp/mL)</li> </ul> <p><b>Severe adverse reactions were reported by:</b></p> <ul style="list-style-type: none"> <li>&gt; 24/253 participants (9%) (Dose 1x10exp11 vp/mL)</li> <li>&gt; 1/129 participant (1%) (Dose 5x10exp10vp/mL)</li> </ul> <p><b>No severe adverse reactions documented</b></p>
LANCET 20JUL2020	Safety and immunogenicity of the ChAdOx1 nCoV-19 vaccine against SARS-CoV-2: a preliminary report of a phase 1/2, single-blind, randomised controlled trial	Folegatti P et al., <a href="#">gotopaper</a>	Vaccine	<p><b>PROTOCOLE DESIGN (NCT04324606):</b></p> <ul style="list-style-type: none"> <li>&gt; Vaccine: ChAdOx1-vectored COVID 19 vaccine from Oxford/AstraZeneca</li> <li>&gt; Phase 1/2 clinical trial; randomized, single blinded, -controlled.</li> <li>&gt; 1077 participants between 18 and 55 years of age (seronegative for SARS-CoV-2 infection). Recruited in 5 sites across UK</li> <li>&gt; Dose: 5x10exp10vp/mL of ChAdOx1-nCov19 . MenACWY used as control (1:1). One single administration IM</li> <li>-10 participants received ChAdOx1 nCoV-19 end prime boost (administration at 28d)</li> <li>&gt; Prophylactic paracetamol administered in 2/3 sites</li> </ul> <p><b>Primary outcomes:</b></p> <ul style="list-style-type: none"> <li>&gt; Measurement of humoral (Total and NABs) and cellular responses</li> <li>&gt; Assessment of efficacy and safety</li> </ul> <p><b>RESULTS:</b></p> <ul style="list-style-type: none"> <li>&gt; Local and systemic reactions were more common in the ChAdOx1 nCoV-19 group and reduced by use of prophylactic paracetamol</li> <li>&gt; In ChAdOx1 nCoV-19 vaccinated participants: <ul style="list-style-type: none"> <li>i. spike-specific T-cell responses peaked on day 14 (median 856 spot-forming cells per million peripheral blood mononuclear cells, IQR 493–1802; n=43).</li> <li>ii. Anti-spike IgG responses rose by day 28 (median 157 ELISA units [EU], 96–317; n=127), and were boosted following a second dose (639 EU, 360–792; n=10).</li> <li>iii. Neutralising antibody responses were detected in 32/35 participants (91%) or 35/35 participants (100%) depending on the measuring method after a single dose. After a booster dose, all participants had neutralising activity</li> </ul> </li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Cell host & microbe 18JUL2020	<b>Retrospective Multicenter Cohort Study Shows that Early Interferon Therapy is Associated with Favorable Clinical Responses in COVID-19 Patients</b>	Wang, Nan et al. China/USA <a href="#">gotopaper</a>	Therapeutics	<p><b>Retrospective multicenter cohort study of 446 COVID-19 patients</b> in China to examine the association between the <b>use and timing of IFN-<math>\alpha</math>2b</b> and clinical outcomes.</p> <ul style="list-style-type: none"> <li>- Early IFN use with significantly reduced in-hospital mortality.</li> <li>- No significant clinical benefit of IFNs was observed in moderately ill COVID-19 patients</li> <li>- Late administration of IFN could be associated with longer hospital stay and slower recovery of lung function.</li> <li>- Using early IFNs with LPV/r is associated with more favorable clinical responses than by using LPV/r alone in COVID-19 patients.</li> </ul> <p>=&gt; Early administration of IFN-<math>\alpha</math>2b was associated with reduced in-hospital mortality among severe to critical COVID-19 patients. In contrast, late interferon therapy increased mortality and delayed recovery, suggesting the <b>timing of interferon therapy is crucial for favorable responses</b> in COVID-19 patients.</p> <p><b>Limitations:</b> retrospective design and non randomized assignment of therapies ; detailed virologic data were not included in the study that precluded comparison with randomized controlled trials of IFNs ; regression models did not include location of care as a confounder because of multicollinearity with therapy choices between IFNs and LPV/r ; adjunctive and supportive therapies were not included in the analyses but could influence the length of hospital stay.</p>
N. Engl. J. Med. 17JUL2020	<b>Dexamethasone in Hospitalized Patients with Covid-19 - Preliminary Report</b>	Horby, Peter et al. UK <a href="#">gotopaper</a>	Therapeutics	<p><b>Preliminary results</b> oral or intravenous <b>dexamethasone</b> vs usual care alone in <b>RECOVERY, controlled, open-label trial</b> comparing a range of possible treatments in patients <b>hospitalized with Covid-19</b>. Primary outcome: 28-day mortality.</p> <p>2104 patients assigned to receive dexamethasone and 4321 to receive usual care.</p> <ul style="list-style-type: none"> <li>- 482 patients (22.9%) in the dexamethasone group and 1110 patients (25.7%) in the usual care group died within 28 days after randomization (age-adjusted rate ratio, 0.83; 95% confidence interval [CI], 0.75 to 0.93; <math>P &lt; 0.001</math>).</li> <li>- The proportional and absolute between-group differences in mortality <b>varied considerably according to the level of respiratory support</b> that the patients were receiving at the time of randomization. In the dexamethasone group, the incidence of death was lower than that in the usual care group among patients receiving invasive mechanical ventilation (29.3% vs. 41.4%; rate ratio, 0.64; 95% CI, 0.51 to 0.81) and among those receiving oxygen without invasive mechanical ventilation (23.3% vs. 26.2%; rate ratio, 0.82; 95% CI, 0.72 to 0.94) but not among those who were receiving no respiratory support at randomization (17.8% vs. 14.0%; rate ratio, 1.19; 95% CI, 0.91 to 1.55).</li> </ul> <p>=&gt; <b>In patients hospitalized with Covid-19, the use of dexamethasone resulted in lower 28-day mortality among those who were receiving either invasive mechanical ventilation or oxygen alone at randomization but not among those receiving no respiratory support and results were consistent with possible harm in this subgroup</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Eurosurveillance 16JUL2020	<b>Convalescent plasma treatment for SARS-CoV-2 infection: analysis of the first 436 donors in England, 22 April to 12 May 2020</b>	Harvala, Heli et al. UK <a href="#">gotopaper</a>	Therapeutics	<p><b>Serological reactivity</b> was analysed in <b>plasma from 436 individuals with a history of disease compatible with COVID-19</b>, including 256 who had been <b>laboratory-confirmed</b> with SARS-CoV-2 infection.</p> <ul style="list-style-type: none"> <li>- <b>Over 99% of laboratory-confirmed cases developed a measurable antibody response</b> (254/256) and <b>88% harboured neutralising antibodies</b> (226/256).</li> <li>- <b>Antibody levels declined over 3 months following diagnosis</b>, emphasising the importance of the timing of convalescent plasma collections.</li> <li>- Finally, the study indicates that <b>commercial ELISA can perform effectively as surrogate assays for predicting neutralising antibody titres</b> and represent a stream-lined and rapid way to guide convalescent plasma donor selection.</li> </ul>
Ann Intern Med. 16JUL2020	<b>Hydroxychloroquine in Nonhospitalized Adults With Early COVID-19: A Randomized Trial</b>	Skipper, Caleb P. et al. USA/Canada <a href="#">gotopaper</a>	Therapeutics	<p><b>Internet-based multisite, international randomized, double blind, placebo-controlled trial</b> to investigate whether <b>HCQ</b> could <b>reduce COVID-19 severity</b> in <b>symptomatic, non hospitalized adults with laboratory-confirmed COVID-19 or probable COVID-19 and high-risk exposure</b> within 4 days of symptom onset.</p> <p>The primary end point was change in <b>overall symptom severity</b> over 14 days, because the pooled event rate of hospitalization or death was substantially lower than initial 10% expectation.</p> <p><b>423 patients contributed primary end point data.</b> Of these, <b>341 (81%) had lab-confirmed infection</b> with SARS-CoV-2 or <b>epidemiologically linked exposure</b> to a person with lab confirmed infection; 56% (236) were enrolled within 1 day of symptoms starting.</p> <ul style="list-style-type: none"> <li>- Change in symptom severity over 14 days <b>did not differ</b> between the HCQ and placebo groups (difference in symptom severity: relative, 12%; absolute, -0.27 points [95% CI, -0.61 to 0.07 points]; P = 0.117).</li> <li>- At 14 days, 24% (49 of 201) of participants receiving HCQ had ongoing symptoms compared with 30% (59 of 194) receiving placebo (P = 0.21).</li> <li>- Medication adverse effects occurred in 43% (92 of 212) of participants receiving HCQ versus 22% (46 of 211) receiving placebo (P &lt; 0.001). With placebo, 10 hospitalizations occurred (2 non-COVID-19-related), including 1 hospitalized death. With HCQ, 4 hospitalizations occurred plus 1 non hospitalized death (P = 0.29).</li> </ul> <p><b>=&gt; HCQ did not substantially reduce symptom severity in outpatients with early, mild COVID-19.</b></p> <p><b>Limitations:</b> Only 58% of participants received SARS-CoV-2 testing because of severe U.S. testing shortages.</p>
Clin. Infect. Dis. 16JUL2020	<b>Hydroxychloroquine for Early Treatment of Adults with Mild Covid-19: A Randomized Controlled Trial</b>	Mitja, Oriol et al. Spain <a href="#">gotopaper</a>	Therapeutics	<p><b>Multicenter, open label, randomized controlled trial in non hospitalized adult patients with recently confirmed SARS-CoV-2 infection</b> and less than five days of symptoms receiving <b>HCQ or no antiviral treatment</b> (not-placebo controlled). Study outcomes were the reduction of viral RNA load in nasopharyngeal swabs up to 7 days after treatment start, patient disease progression using the WHO scale up to 28 days, and time to complete resolution of symptoms. Adverse events were assessed up to 28 days.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 16JUL2020	<b>Reconstruction of the full transmission dynamics of COVID-19 in Wuhan</b>	Hao, Xingjie <i>et al.</i> China, USA <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Modelling approach to reconstruct the full-spectrum dynamics of COVID-19 (Wuhan China, 1 January to 8 March 2020) across 5 periods marked by events and non-pharmaceutical interventions (based on 32,583 laboratory-confirmed cases), accounting for presymptomatic infectiousness, time-varying ascertainment rates, transmission rates and population movements:</p> <ul style="list-style-type: none"> <li>- High covertness and high transmissibility were key features identified.</li> <li>- <b>Estimate 87%</b> (lower bound 53%) <b>of infections before March 8 were unascertained</b>, potentially including asymptomatic and mild-symptomatic cases, and <b>basic reproduction number R0 of 3.54 in the early outbreak</b> (much higher than for SARS and MERS).</li> <li>- Positive effects of <b>multi-pronged interventions</b> : <b>decreasing the reproduction number to 0.28, reducing total infections in Wuhan by 96.0%</b> as of March 8.</li> <li>- <b>Probability of resurgence following lifting of all interventions after 14 days of no ascertained infections, estimated at 0.32 and 0.06 based on models with 87% and 53% unascertained infections, respectively.</b></li> <li>-&gt; <b>Risk posed by unascertained cases when changing intervention strategies highlight the important implications for continuing surveillance and interventions to eventually contain COVID-19 outbreaks.</b></li> </ul>
Antiviral Research 15JUL2020	<b>Lower prevalence of antibodies neutralizing SARS-CoV-2 in group O French blood donors</b>	Gallian, Pierre <i>et al.</i> France <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Distribution of antibodies neutralizing SARS-CoV-2 according to age, sex or blood group in 998 samples from French blood donors (last week of March or first week of April 2020) :</p> <ul style="list-style-type: none"> <li>- At this stage of the outbreak, prevalence was low (2.7%) and criteria for blood donation imply that vast majority of seropositives had asymptomatic or pauci-symptomatic SARS CoV-2 infections.</li> <li>- Antibodies neutralizing SARS-CoV-2 found with similar prevalence in men and women.</li> <li>- Virus infection may occur at similar incidence in men and women (2.82% vs 2.69%), while severe forms are more frequent in men.</li> <li>- <b>Proportion of seropositives significantly lower in group O donors</b> (1.32% vs 3.86% in other donors, <math>p = 0.014</math>).</li> <li>-&gt; <b>Blood group O persons are less at risk of being infected by SARS-CoV-2 than other blood groups persons.</b></li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
BMJ 15JUL2020	<b>Physical distancing interventions and incidence of coronavirus disease 2019: natural experiment in 149 countries</b>	Islam, N et al. UK/USA <a href="#">gotopaper</a>	Public Health / Epidemio	<p>AIM: To evaluate the incidence of Covid-19 before and after physical distancing intervention using data of 149 countries that implemented one of the five physical distancing interventions (closures of schools, workplaces, and public transport, restrictions on mass gatherings and public events, and restrictions on movement (lockdowns)) between 1 January and 30 May 2020.</p> <p>--&gt; Implementation of any physical distancing intervention was associated with an overall reduction in covid-19 incidence of 13%;</p> <p>--&gt; Closure of public transport was not associated with any additional reduction in covid-19 incidence when the other four physical distancing interventions were in place;</p> <p>--&gt; Data from 11 countries also suggested similar overall effectiveness (pooled IRR 0.85, 0.81 to 0.89) when school closures, workplace closures, and restrictions on mass gatherings were in place;</p> <p>--&gt; Earlier implementation of lockdown was associated with a larger reduction in covid-19 compared with a delayed implementation of lockdown after other physical distancing interventions were in place.</p> <p>These findings might support policy decisions related to physical distancing measures in current or future epidemic waves.</p>
Journal of clinical medicine 14JUL2020	<b>Epidemiology and Clinical Presentation of Children Hospitalized with SARS-CoV-2 Infection in Suburbs of Paris</b>	Gaborieau, L et al. France <a href="#">gotopaper</a>	Public Health / Epidemio	<p>Prospective, observational, multicentre study aimed at identifying the epidemiological characteristics, clinical presentation, and prognosis of 192 children with Covid-19 hospitalized in Paris suburb hospitals from 23 March to 10 May 2020 (lockdown period).</p> <ul style="list-style-type: none"> <li>- Median age of children was one year old, sex ratio 1.3:1;</li> <li>- Fever was recorded in 76.6% children and poorly tolerated in 15.1%. Symptoms ranged from rhinorrhoea (34.4%) and gastrointestinal (35.5%) to respiratory distress (25%). Only 5.2% of children had anosmia and 2.6% had chest pain. An underlying condition was identified in almost 30% of cases;</li> <li>- 24 (12.5%) children were admitted to paediatric intensive care units, 12 required mechanical ventilation, and three died.</li> </ul> <p>Although most Covid-19 paediatric cases showed mild to moderate clinical expression, one-eighth of children were admitted to paediatric intensive care units and three died.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 15JUL2020	<b>Potently neutralizing and protective human antibodies against SARS-CoV-2</b>	Zost, Seth J. et al. USA <a href="#">gotopaper</a>	Therapeutics	<p>From a larger panel of human monoclonal antibodies (mAbs) targeting the spike (S) glycoprotein, several were identified that exhibited potent neutralizing activity and fully blocked the receptor-binding domain of S (SRBD) from interacting with human ACE2 (hACE2).</p> <ul style="list-style-type: none"> <li>- Competition-binding, structural, and functional studies allowed clustering of the mAbs into classes recognizing distinct epitopes on the SRBD as well as distinct conformational states of the S trimer.</li> <li>- Potent neutralizing mAbs recognizing non-overlapping sites, COV2-2196 and COV2-2130, bound simultaneously to S and synergistically neutralized authentic SARS-CoV-2 virus.</li> <li>- In two mouse models of SARS-CoV-2 infection, passive transfer of either COV2-2196 or COV2-2130 alone or a combination of both mAbs protected mice from weight loss and reduced viral burden and inflammation in the lung.</li> <li>- Passive transfer of each of two of the most potently ACE2 blocking mAbs (COV2-2196 or COV2-2381) as monotherapy protected rhesus macaques from SARS-CoV-2 infection.</li> </ul> <p>=&gt; These results identify protective epitopes on SRBD and provide a structure-based framework for rational vaccine design and the selection of robust immunotherapeutics.</p>
JAMA 14JUL2020	<b>Rapid implementation of SARS-CoV-2 sequencing to investigate cases of health-care associated COVID-19: a prospective genomic surveillance study</b>	Meredith et al, UK <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Prospective surveillance study Between March 13 and April 24, 2020, collection of clinical data and samples from 5613 patients with COVID-19 from across the East of England</p> <ul style="list-style-type: none"> <li>-&gt; Rapid SARS-CoV-2 nanopore sequencing from PCR-positive diagnostic samples, enabling sample-to-sequence in less than 24 h</li> <li>-&gt; Establishment of a weekly review and reporting system with integration of genomic and epidemiological data to investigate suspected health-care associated COVID-19 cases</li> </ul> <p>Results :</p> <p>Sequencing of 1000 samples producing 747 high-quality genomes. Combination of epidemiological and genomic analysis of the 299 patients from our hospital and identified 35 clusters of identical viruses involving 159 patients. 92 (58%) of 159 patients had strong epidemiological links and 32 (20%) patients had plausible epidemiological links. These results were fed back to clinical, infection control, and hospital management teams, leading to infection-control interventions and informing patient safety reporting.</p> <p>Conclusions :</p> <p>Benefit of combined genomic and epidemiological analysis for the investigation of health-care associated COVID-19 Possibility to detect cryptic transmission events and identify opportunities to target infection-control interventions to further reduce health-care associated infections. Our findings have important implications for national public health policy as they enable rapid tracking and investigation of infections in hospital and community settings.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 14JUL2020	<b>Association Between Universal Masking in a Health Care System and SARS-CoV-2 Positivity Among Health Care Workers</b>	Wang et al., USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Mass General Brigham (MGB) is the largest health care system in Massachusetts, with 12 hospitals and more than 75 000 employees.</p> <p>Using electronic medical records -&gt; identification of HCWs providing direct and indirect patient care who were tested for SARS-CoV-2 with RT PCR between March 1 and April 30, 2020.</p> <p>3 phases during the study period:</p> <ol style="list-style-type: none"> <li>1- A preintervention period before implementation of universal masking of HCWs (March 1-24, 2020)</li> <li>2- A transition period until implementation of universal masking of patients (March 25–April 5, 2020)</li> <li>3- An additional lag period to allow for manifestations of symptoms (April 6-10, 2020), as previously defined; and an intervention period (April 11-30, 2020)</li> </ol> <p><b>Results</b> : Of 9850 tested HCWs, 1271 (12.9%) had positive results for SARS-CoV-2 (median age, 39 years; 73% female; 7.4% physicians or trainees, 26.5% nurses or physician assistants, 17.8% technologists or nursing support, and 48.3% other).</p> <p>During the preintervention period, the SARS-CoV-2 positivity rate increased exponentially from 0% to 21.32%, with a weighted mean increase of 1.16% per day and a case doubling time of 3.6 days (95% CI, 3.0-4.5 days).</p> <p>During the intervention period, the positivity rate decreased linearly from 14.65% to 11.46%, with a weighted mean decline of 0.49% per day and a net slope change of 1.65% (95% CI, 1.13%-2.15%; <math>P &lt; .001</math>) more decline per day compared with the preintervention period.</p> <p><b>Conclusion</b> : Universal masking at MGB was associated with a significantly lower rate of SARS-CoV-2 positivity among HCWs. Randomized trials of universal masking of HCWs during a pandemic are likely not feasible. Nonetheless, these results support universal masking as part of a multipronged infection reduction strategy in health care settings.</p>
Nature Comm 14JUL2020	<b>Transplacental transmission of SARS-CoV-2 infection</b>	Vivanti A J. et al France <a href="#">gotopaper</a>	Clinic Case study	<p>23 years old G1P0: SARS-CoV-2 detected in blood, in nasopharyngeal and vaginal swabs (35+2 weeks of gestation) All routine tests were normal, ultrasound normal.</p> <p>At 35+5 weeks of gestation à cesarean section for pathological fetal heart rate tracing:</p> <ul style="list-style-type: none"> <li>- Amniotic fluid tested positive for SARS-CoV-2 prior to the rupture of membranes</li> <li>- RT-PCR on the placenta was positive for SARS-CoV-2 (viral load much higher)</li> <li>- No other pathogen agent was detected on the placenta</li> </ul> <p>Neonate:</p> <ul style="list-style-type: none"> <li>- Male, birth weight 2540g</li> <li>- Apgar scores 4 – 2 – 7 (1, 5 and 10 min) à intubation à intensive care unit</li> <li>- Extubated at H6, no sedative or analgesic drug</li> <li>- Echocardiography and lung ultrasound were normal</li> <li>- RT-PCR positive for SARS-CoV-2 on blood and non-bronchoscopic bronchoalveolar lavage fluid</li> <li>- Nasopharyngeal and rectal swabs were positive for SARS-CoV-2: 1h of life, at 2 and 18 days.</li> </ul> <p>On the third day:</p> <ul style="list-style-type: none"> <li>- Irritability, axial hypertonia &amp; opisthotonos</li> <li>- CSF was negative, blood culture was sterile, EEG normal, no signs suspected for metabolic disease</li> <li>- Improved slowly, mild hypotonia and feeding difficulty persisted</li> </ul> <p>MRI at 11 day: bilateral gliosis of the deep white periventricular and subcortical matter</p> <p>2 months of life: improved neurological examination an MRI. Clinical exam normal</p> <p>-&gt; Description of an actual neonatal infection</p> <p>-&gt; A case of congenital infection associated with neurological manifestations following neonatal viremia</p> <p>-&gt; Demonstration that the transplacental transmission of SARS-CoV-2 infection is possible</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature Communications 14JUL2020	<b>Exploring the SARS-CoV-2 virus-host-drug interactome for drug repurposing</b>	Sadegh, Sepideh et al. Germany <a href="#">gotopaper</a>	Therapeutics	CoVex, a web-based platform for the interactive exploration and network-based analysis of virus–host interactions, aimed towards drug repurposing for the treatment of COVID-19. It integrates virus-human protein interactions, human protein-protein interactions, and drug-target interactions. It allows visual exploration of the virus-host interactome and implements systems medicine algorithms for network-based prediction of drug candidates. Thus, CoVex is a resource to understand molecular mechanisms of pathogenicity and to prioritize candidate therapeutics.
NEJM 14JUL2020	<b>An mRNA Vaccine against SARS-CoV-2 — Preliminary Report</b>	Jackson LA et al, USA <a href="#">gotopaper</a>	Vaccine	<p>&gt; Phase 1 clinical Trial with mRNA-1273 vaccine from Moderna</p> <ul style="list-style-type: none"> <li>- dose-escalation, open-label trial including</li> <li>- 45 healthy adults, 18 to 55 years of age</li> <li>- two vaccinations, 28 days apart</li> <li>- Doses: 25 µg, 100 µg, or 250 µg.</li> </ul> <p><b>RESULTS:</b></p> <p>&gt; After the first vaccination, antibody responses were higher with higher dose. GMTs (measured by ELISA d29):</p> <ul style="list-style-type: none"> <li>- 40,227 in the 25-µg group,</li> <li>- 109,209 in the 100-µg group</li> <li>- 213,526 in the 250-µg group.</li> </ul> <p>&gt; After the second vaccination, the titers increased. GMT(measured by ELISA d57):</p> <ul style="list-style-type: none"> <li>- 299,751 in the 25-µg group,</li> <li>- 782,719 in the 100-µg group</li> <li>- 1,192,154 in the 250-µg group.</li> </ul> <p>- Serum neutralizing activity was detected in all participants evaluated</p> <p>&gt; Solicited AE occurred in more than half the participants (fatigue, chills, headache, myalgia, and pain at the injection site)</p> <p>&gt; Systemic AE adverse events were more common after the second vaccination, particularly with the highest dose.</p> <p>&gt; 3 participants in the 250-µg dose group reported one or more severe AE</p> <p><b>CONCLUSIONS</b></p> <p>The mRNA-1273 vaccine induces anti-SARS-CoV-2 immune responses in all participants, and no trial-limiting safety concerns were identified.</p>
Nature structural & molecular biology 13JUL2020	<b>Neutralizing nanobodies bind SARS-CoV-2 spike RBD and block interaction with ACE2</b>	Huo, Jiangdong et al. UK <a href="#">gotopaper</a>	Therapeutics	<p>Using a naive llama single-domain antibody library and PCR-based maturation, 2 closely related nanobodies, H11-D4 and H11-H4, that bind RBD (KD of 39 and 12 nM, respectively) and block its interaction with ACE2 were produced.</p> <ul style="list-style-type: none"> <li>- Single-particle cryo-EM revealed that both nanobodies bind to all three RBDs in the spike trimer.</li> <li>- Crystal structures of each nanobody–RBD complex revealed how both nanobodies recognize the same epitope, which partly overlaps with the ACE2 binding surface, explaining the blocking of the RBD–ACE2 interaction.</li> <li>- Nanobody-Fc fusions showed neutralizing activity against SARS-CoV-2 (4–6 nM for H11-H4, 18 nM for H11-D4) and additive neutralization with the SARS-CoV-1/2 antibody CR3022. Such additive combinations are a well-known strategy to reduce the propensity of the virus to escape by mutating.</li> </ul> <p>=&gt; Nanobody maturation technology can be deployed to produce a highly neutralizing agent against an emerging viral threat in real time. The approach may be useful in identifying complementary epitopes to those identified by animal immunization approaches. The H11-H4 and H11-D4 nanobodies may find application in a cocktail of laboratory-synthesized neutralizing antibodies given for passive immunization of severely ill COVID-19 patients.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Science 13JUL2020	<b>Impaired type I interferon activity and inflammatory responses in severe COVID-19 patients</b>	Hadjadj J et al France <a href="#">gotopaper</a>	Therapeutic	<p>Integrated immune analysis – 50 COVID-19 patients and 18 healthy controls were included: mild-to-moderate (n=15) – severe (n=17) – critical (n=18)</p> <p>Immunological transcriptional signature:</p> <ul style="list-style-type: none"> <li>- Data suggested a severity grade-dependent increase in activation of innate and inflammatory pathways.</li> <li>- IFN response was high in mild-to-moderate patients while it was reduced in more severe patients.</li> <li>- IFN activity in serum was significantly lower in severe/critical patients as compared to mild-to-moderate patients.</li> <li>- Genes specifically up-regulated in severe or critical patients mainly belonged to the NF-κB pathway</li> </ul> <p>SARS-CoV-2 infection à an absence of circulating IFN-β in COVID-19 patients with all disease-severity grade</p> <ul style="list-style-type: none"> <li>&gt; Suggested that infected patients had no detectable circulating IFN-β and that an impaired IFN-α production characterized the most severe COVID-19 cases.</li> <li>&gt; Impaired type I IFN response in severe and critical COVID-19 patients, accompanied by high blood viral load and an excessive NF-κB driven inflammatory response associated with increase TNFα and IL-6.</li> <li>&gt; Type I IFN deficiency is a hallmark of severe COVID-19 à combined therapeutic approaches</li> </ul>
Cell 13JUL2020	<b>Longitudinal isolation of potent near-germline SARS-CoV-2-neutralizing antibodies from COVID-19 patients</b>	Kreer, Christoph et al. Germany <a href="#">gotopaper</a>	Vaccines	<p>Longitudinal analysis of the antibody response of 12 COVID-19 patients from 8 to 69 days after diagnosis.</p> <ul style="list-style-type: none"> <li>- By screening 4,313 SARS-CoV-2-reactive B cells, 255 antibodies were isolated from different time points as early as 8 days after diagnosis.</li> <li>- Of these, 28 potentially neutralized authentic SARS-CoV-2 with IC<sub>50</sub> as low as 0.04mg/mL, showing a broad spectrum of variable (V) genes and low levels of somatic mutations.</li> <li>- Interestingly, potential precursor sequences were identified in naive B cell repertoires from 48 healthy individuals who were sampled before the COVID-19 pandemic.</li> </ul> <p>=&gt; These results demonstrate that SARS-CoV-2-neutralizing antibodies are readily generated from a diverse pool of precursors, fostering hope for rapid induction of a protective immune response upon vaccination.</p>
Clin. Infect. Dis. 11JUL2020	<b>Tocilizumab for treatment of mechanically ventilated patients with COVID-19</b>	Somers, Emily C. et al. USA <a href="#">gotopaper</a>	Therapeutics	<p>Observational, single-center, controlled study of 154 patients with severe COVID-19 illness requiring mechanical ventilation to assess effectiveness and safety of IL-6 blockade with tocilizumab. Primary endpoint was survival probability post-intubation; secondary analyses included an ordinal illness severity scale integrating superinfections.</p> <ul style="list-style-type: none"> <li>- In propensity score inverse probability weighting (IPTW)-adjusted models, tocilizumab was associated with a 45% reduction in hazard of death [hazard ratio 0.55 (95% CI 0.33, 0.90)] and improved status on the ordinal outcome scale [odds ratio per 1-level increase: 0.58 (0.36, 0.94)].</li> <li>- Though tocilizumab was associated with an increased proportion of patients with superinfections (54% vs. 26%; p&lt;0.001), there was no difference in 28-day case fatality rate among tocilizumab-treated patients with versus without superinfection [22% vs. 15%; p=0.42]. Staphylococcus aureus accounted for ~50% of bacterial pneumonia.</li> </ul> <p>Limitations: observational study ; does not address the potential role of tocilizumab earlier in illness for preventing mechanical ventilation, the optimal dose of tocilizumab, the potential utility of multiple doses, or the role of IL-6 serum concentrations (which were not routinely available) in predicting tocilizumab response ; tocilizumab usage in this clinical care setting was not dictated by a firm study protocol, and therefore not completely standardized.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature Med 10JUL2020	<b>Rapid isolation and profiling of a diverse panel of human monoclonal antibodies targeting the SARS-CoV-2 spike protein</b>	Zost SJ et al USA <a href="#">gotopaper</a>	Therapeutic	<p>Use a rapid antibody discovery platform to isolate hundreds of human monoclonal antibodies (mAbs) against SARS-CoV-2 s protein</p> <p>à obtain human mAbs from the B cells of some patients in North America (4 patients)</p> <p>à used different workflows in parallel</p> <p>The antibodies could be grouped into five binding patterns on the basis of domain recognition and cross-reactivity.</p> <p>à Most neutralizing mAbs recognizing the receptor-binding domain (RBD) of S</p> <p>à the RBD= the principal site of vulnerability for SARS-CoV-2 neutralization</p> <p>à RBD of SARS-CoV-2 for vaccine design and therapeutic-antibody development</p>
PNAS 9 JULY2020	<b>BCG vaccine protection from severe coronavirus disease 2019 (COVID-19)</b>	Escobar LE et al, USA <a href="#">gotopaper</a>	Vaccine/Public Health	<p>&gt; Epidemiological explorations suggest a negative association between national BCG vaccination policy and the prevalence and mortality of COVID-19</p> <p>&gt; But comparisons are difficult due to broad differences between countries:</p> <ul style="list-style-type: none"> <li>- socioeconomic status,</li> <li>- demographic structure</li> <li>- rural vs. urban settings</li> <li>- number of diagnostic tests</li> <li>- national control strategies</li> </ul> <p>RESULTS:</p> <p>1. Strong correlation between the BCG index (estimation of the degree of universal BCG vaccination in a country), and COVID19 mortality in different socially similar European countries (<math>r^2 = 0.88</math>; <math>P = 8 \times 10^{-7}</math>)</p> <p>- every 10% increase in the BCG index was associated with a 10.4% reduction in COVID-19 mortality</p> <p>2. Results fail to confirm the null hypothesis of no association between BCG vaccination and COVID-19 mortality, and suggest that BCG could have a protective effect</p> <p>3. Nevertheless, the analyses should be considered with caution. BCG vaccination clinical trials are required to corroborate the patterns detected</p>
Eurosurveillance 09JULY2020	<b>International external quality assessment for SARS-CoV-2 molecular detection and survey on clinical laboratory preparedness during the COVID-19 pandemic, April/May 2020</b>	Matheeußen, V et al. International <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>External quality assessments (EQA) on molecular detection of RT-PCR SARS-CoV-2 tests : based on 521 datasets from 365 of 406 laboratories from 36 countries. 23% of the assays were in-house assays which performed as well or better than commercial ones. The overall qualitative performance of the participating laboratories was at an acceptable level. A lack of standardisation emerges, and extraction and amplification methods were significantly associated with correct classification of all samples. Laboratories should be aware of the limitations of their assays and perform their own validation and verification in line with ISO 15189 or equivalent requirements</p> <p>EU RECOVER project survey assessing the molecular testing capacity and throughput for SARS-CoV-2 detection of clinical laboratories : almost 80% of the participating laboratories E5(n = 360) are capable of generating a PCR result within 24 h after receiving the sample, 48% of the laboratories could analyse more than 250 samples per day.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 08JULY2020	<b>OpenSAFELY: factors associated with COVID-19 death in 17 million patients</b>	Williamson, EJ et al. <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>OpenSAFELY is a secure health analytics platform covering 40% of all patients in England. Primary care records of 17,278,392 adults were pseudonymously linked to 10,926 COVID-19-related deaths.</p> <ul style="list-style-type: none"> <li>- COVID-19-related death was associated with male sex, older age and deprivation, diabetes, severe asthma and other conditions / Compared with people with white ethnicity, Black and South Asian people were at higher risk.</li> </ul> <p>Retrospective cohort study on data from COVID-19, SARS and other HCoV patients from Hong Kong, and the association with Alanine aminotransferase (ALT)/aspartate aminotransferase (AST) levels as marker of liver injury.</p> <ul style="list-style-type: none"> <li>- Cohorts : 1040 COVID-19 patients (mean age 38, 54% men), 1670 SARS patients (mean age 44, 44% men) and 675 other HCoV patients (mean age 20, 57% men).</li> <li>- ALT/AST was elevated in 50.3% SARS patients, 22.5% COVID-19 patients and 36.0% other HCoV patients.</li> <li>- For COVID-19 patients, 53 (5.1%) were admitted to ICU, 22 (2.1%) received invasive mechanical ventilation and 4 (0.4%) died.</li> <li>- ALT/AST elevation was independently associated with primary end point (ICU admission, mechanical ventilation and/or death) after adjusted for albumin, diabetes and hypertension. Use of lopinavir-ritonavir +/- ribavirin + interferon and corticosteroids was independently associated with ALT/AST elevation.</li> </ul>
CLIN. INFECT. DIS. 08JULY2020	<b>Evaluating use cases for human challenge trials in accelerating SARS-CoV-2 vaccine development</b>	Nguyen LC et al USA, CANADA, UK, <a href="#">gotopaper</a>	Vaccine	<p>Human challenge trials (HCTs) have been proposed as a means to accelerate SARS-CoV-2 vaccine development.</p> <p>Potential scenarios where using HCT could generate useful data for rapid vaccine development:</p> <ol style="list-style-type: none"> <li>1. Evaluating efficacy: use of HCT alongside an expanded safety trial to replace Ph3 , or in parallel with Ph3 trial to give an early indicator of efficacy. HCT could take as little as two months to conduct and would require far fewer participants than a phase 3 trial due to viral exposure being guaranteed by the challenge</li> <li>2. Converging on correlates of protection: HCTs could be used to identify or verify CoPs against disease endpoints. These CoPs could then be used as surrogate endpoints in phase 3 trials (instead of clinical endpoints).</li> <li>3. Improving understanding of pathogenesis and the human immune response: A COVID-19 HCM would allow close observation of the participants prior to and from the point of vaccination and infection. This could help resolve the physiological basis for variation in disease severity, the disease's progression from infection, or the immune response upon re-infection.</li> </ol> <p><b>LIMITATIONS</b></p> <ul style="list-style-type: none"> <li>&gt; Timing of viral challenge relative to vaccination (same for all patients in HCT but highly variable in real world) / Method of administration can affect the nature of infection and the immune response</li> </ul>
Cell Reports 07JUL2020	<b>Remdesivir inhibits SARS-CoV-2 in human lung cells and chimeric SARS-CoV expressing the SARS-CoV-2 RNA polymerase in mice</b>	Pruijssers A et al USA <a href="#">gotopaper</a>	Therapeutic	<p>Rendesivir (RDV) showed both prophylactic and therapeutic efficacy in mouse models of SARS and MERS.</p> <p><b>In vitro:</b></p> <ul style="list-style-type: none"> <li>- RDV potently inhibits SARS-CoV-2 replication in human lung cells and primary human airway epithelial culture.</li> <li>- Different cell lines were tested: Vero E6, Vero CCL-8, Huh7 and Calu3 2B4.</li> <li>- Weaker activity is observed in Vero E6 cells (EC50 = 1,65 µ M) due to their low capacity to metabolize RDV.</li> </ul> <p><b>In vivo:</b></p> <ul style="list-style-type: none"> <li>- SARS-CoV-2 does not bind the murine ortholog of the human entry receptor to enter cells / Constructed a chimeric mouse-adapted SARS-CoV variant encoding the target of RDV antiviral activity, the RdRp, of SARS-CoV-2 / RDV is active against the SARS-CoV-2 RdRp in vivo / Therapeutic RDV administration diminishes lung viral load and improves pulmonary function compared to vehicle treated animals.</li> </ul> <p>Caution should be exercised when interpreting nucleoside prodrug potency experiments performed using Vero cell lineages.</p> <p>RDV is potently active against SARS-CoV-2 in vitro and in vivo</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Annals of Internal Medicine 06JULY2020	<b>Clinical Validity of Serum Antibodies to SARS-CoV-2</b>	Caturegli et al., USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Case Control Study Objective: To determine the clinical validity and utility of SARS-CoV-2 antibodies.</p> <p>Serum IgG and IgA antibodies against SARS-CoV-2 spike protein were detected by using ELISA</p> <p>Sensitivity and specificity of the SARS-CoV-2 IgG assay were 0.976 (95% CI, 0.928 to 0.995) and 0.988 (CI, 0.974 to 0.995), respectively, when performed 14 days or later after symptom onset, but sensitivity decreased at earlier time points. Immunoglobulin G developed rapidly and was sustained at high levels throughout follow-up (up to 58 days). Antibodies to SARS-CoV-2 predicted the odds of developing acute respiratory distress syndrome, which increased by 62% (CI, 48% to 81%; <math>P &lt; 0.001</math>) for every 2-fold increase in IgG. Of 11 066 NAAT-tested patients, 457 were repeatedly NAAT-negative, and serum samples were obtained for 18 such patients: 6 COVID-19 case patients and 12 non-COVID-19 control patients. Antibodies were present in 5 of 6 case patients and none of the 12 control patients (<math>P = 0.001</math>).</p> <p>Antibodies to SARS-CoV-2 demonstrate infection when measured at least 14 days after symptom onset, associate with clinical severity, and provide valuable diagnostic support in patients who test negative by NAAT but remain clinically suspicious for COVID-19.</p>
The Lancet 06JULY2020	<b>Prevalence of SARS-CoV-2 in Spain (ENE-COVID): a nationwide, population-based seroepidemiological study</b>	Pollan et al., Spain and USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>- 35 883 households were selected</p> <p>From April 27 to May 11, 2020, 61 075 participants answered a questionnaire on history of symptoms compatible with COVID-19 and risk factors, received a point-of-care antibody test, and, if agreed, donated a blood sample for additional testing with a chemiluminescent microparticle immunoassay.</p> <p>-&gt; Seroprevalence was 5.0% (95% CI 4.7–5.4) by the point-of-care test and 4.6% (4.3–5.0) by immunoassay, with a specificity–sensitivity range of 3.7% (3.3–4.0; both tests positive) to 6.2% (5.8–6.6; either test positive), with no differences by sex and lower seroprevalence in children younger than 10 years (&lt;3.1% by the point-of-care test).</p> <p>-&gt; Substantial geographical variability: higher prevalence around Madrid (&gt;10%) and lower in coastal areas (&lt;3%).</p> <p>-&gt; In 7273 individuals with anosmia or at least three symptoms, seroprevalence ranged from 15.3% (13.8–16.8) to 19.3% (17.7–21.0). Around a third of seropositive participants were asymptomatic, ranging from 21.9% (19.1–24.9) to 35.8% (33.1–38.5). Only 19.5% (16.3–23.2) of symptomatic participants who were seropositive by both the point-of-care test and immunoassay reported a previous PCR test.</p>
The Lancet ID 03JULY2020	<b>Asymptomatic SARS-CoV-2 infection in Belgian long-term care facilities</b>	Hoxha et al., Sweden and Belgium <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Nasopharyngeal and oropharyngeal swabs were collected from residents and staff in long-term care facilities and sent for real-time PCR testing</p> <p>-&gt; Data were reported for 2074 LTCFs</p> <p>-&gt; cross-sectional analysis of data received from the laboratories between April 8, and May 18, 2020</p> <p>-&gt; 280 427 people were tested, including 142 100 (51%) residents and 138 327 (49%) staff</p> <p>-&gt; 8343 (3.0%) people tested positive, including 2953 (2.1%) staff and 5390 (3.8%) residents.</p> <p>-&gt; No symptoms were reported for 6244 (74.8%, 95% CI 73.9–75.8) of 8343 people who tested positive</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Ann Rheum Dis 03JUL2020	<b>Interleukin-6 blockade with sarilumab in severe COVID-19 pneumonia with systemic hyperinflammation: an open-label cohort study</b>	Della-Torre E et al Italy <a href="#">gotopaper</a>	Therapeutic	<p>Open label study of sarilumab (IL-6 blockade) in severe COVID-19 patients with hyperinflammation</p> <p>2 groups: 28 patients sarilumab + standard of care matched with 28 patients standard of care</p> <p>At day 28 of follow-up</p> <ul style="list-style-type: none"> <li>- Survival rate: 93% (sarilumab) vs 82% (control): HR: 0,36 IC95%[0,08 – 1,68]</li> <li>- Median time to death: 19 days (sarilumab) vs 4 (control), p=0,006</li> <li>- Death: 7% (sarilumab) vs 18% (control), p=0,42</li> <li>- Clinical improvement: 60% (sarilumab) vs 64% (control), p=0,99</li> <li>- Adverse events: similar in both group, 43% (sarilumab) vs 36% (control)</li> </ul> <p>Independent factors of clinical improvement:</p> <ul style="list-style-type: none"> <li>- Baseline PaO<sub>2</sub>/FiO<sub>2</sub> &gt; 100mmHg</li> <li>- Lung consolidation &lt;17% at CT scan</li> </ul> <p>No difference for clinical improvement and mortality between two group</p> <p>Sarilumab was associated with faster recovery in a subset of patient showing minor lung consolidation at baseline</p>
Lancet Child Adolesc Health 02JUL2020	<b>Emergence of Kawasaki disease related to SARS-CoV-2 infection in an epicenter of the French COVID-19 epidemic: a time-series analysis</b>	Ouldali N et al France <a href="#">gotopaper</a>	Clinic	<p>Time series analysis over the past 15 years in a pediatric centre in Paris - Estimation of the number of Kawasaki disease (KD) cases over time (quasi-Poisson regression)</p> <p>230 patients with KD (dec 2005 to may 2020) à 1,2 case/month (quasi-Poisson model)</p> <p>2 peak of hospital admission due to KD</p> <p>April 2020:</p> <ul style="list-style-type: none"> <li>- Increase of KD (497% increase): 6 cases per month</li> <li>- Starting 2 weeks after the peak of COVID-19</li> <li>- 10 cases between April 15 and May 20</li> <li>- 80% of the cases were positive for SARS-CoV-2 (PCR or serology)</li> <li>- 6 cases required intensive care</li> <li>- Similar characteristics to the patients with KD in Bergamo</li> </ul> <p><b>December 2009:</b></p> <ul style="list-style-type: none"> <li>- Concomitant with the influenza A H1N1 pandemic – 1 – 3 weeks after the peak in Paris</li> <li>- 6 cases per month (365% increase)</li> </ul> <p>Characteristics of patients with KD in April 2020 appeared to be different from those diagnosed during the H1N1 epidemics:</p> <ul style="list-style-type: none"> <li>- Older (11,8 years vs 2,1 years, p=0,034)</li> <li>- Less inflammatory: CRP à 23,6 mg/dL vs 8,4 mg/dL, p=0,042</li> </ul> <p>SARS-CoV-2 : only respiratory virus with intense circulation in April 2020 - Viral respiratory infections could be triggers for KD</p>
Eurosurveillance 02JULY2020	<b>Rapid risk assessment from ECDC: Resurgence of reported cases of COVID-19 in the EU/EEA, the UK and EU candidate and potential candidate countries</b>	Eurosurveillance editorial team, Europe <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>From 31.12.2019 to 30.06.2020: 10,273,001 Covid-19 cases reported worldwide, 505,295 deaths. EU/EEA countries and the UK reported 1,556,709 cases (15% of total) and 176,800 deaths (35% of total).</p> <p>Decreasing trends in disease incidence are being observed in Europe (12% decrease in 14-day incidence of reported cases, 16-30 June), but community transmission is still reported, as well as a resurgence of observed cases or localised outbreaks. Reasons explaining apparent increase of resurgence vary (changes in case ascertainment, genuine increases in transmission, importation of cases).</p> <p>In this risk assessment, ECDC assesses the risks associated with the reported increases of incidence. National authorities should carefully analyse every increase in incidence to assess if these are associated with increases in transmission or involve populations at risk. ECDC does not consider travel restrictions within and to the Schengen area as an efficient way to reduce transmission within the EU.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Eurosurveillance 02JULY2020	<b>Introductions and early spread of SARS-CoV-2 in France, 24 January to 23 March 2020</b>	Gámbaro, F. et al., France <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>We sequenced SARS-CoV-2 genomes from clinical cases sampled through surveillance, gaining insights into the initial introductions and spread of the virus in France.</p> <ul style="list-style-type: none"> <li>- Imposed quarantine prevented local transmission. The first cases were imported from China.</li> <li>- The Clade V characteristics suggest convergent evolution or a reversion of the V-clade. Subsequent early cases detected in February in the West or East of France had recent history of travel to Italy but also do not appear to have seeded local transmission.</li> <li>- All other sequences from northern France fall in clade G. Phylogenetic analysis suggests that the French outbreak was mainly seeded by one or several variants of this clade, unlike what is observed for other European countries.</li> <li>- This surveillance allowed to capture one of the earliest representatives of clade G. The mutations in early and late sampling highlight the complexity and risk of inferences based on 1 or 2 nucleotide substitutions.</li> </ul> <p>It is impossible to infer with confidence how the virus was introduced to France, but these data suggest that SARS-CoV-2 might have been present before the first recorded local cases.</p>
Cellular & Molecular Immunology 01JULY2020	<b>Identification of druggable inhibitory immune checkpoints on Natural Killer cells in COVID-19</b>	Demaria, Olivier et al. France <a href="#">gotopaper</a>	Therapeutic	<p>Analysis of NK cells in blood from a cohort of 82 individuals: 10 healthy controls (HC), 10 paucisymptomatic COVID-19 patients (pauci), 34 patients with pneumonia (pneumo) and 28 patients with ARDS due to SARS-CoV-2 infection.</p> <ul style="list-style-type: none"> <li>- The absolute numbers of peripheral blood NK cells, B, CD4+, and CD8+ T lymphocytes were lower in the pneumonia and ARDS groups than in healthy controls.</li> <li>- Among CD45+CD3-CD56+ total NK cells the proportion of mature NK cells was markedly lower in patients with ARDS. Loss of mature NK cells may contribute to the pulmonary complications occurring in the most severe cases of COVID-19.</li> <li>- Presence of a CD39-expressing NK cell population observed in the blood of the COVID-19 patients of the pneumonia and ARDS groups that was absent in the HC and paucisymptomatic groups. Expression of CD39 on NK cells from COVID-19 patients may be explained by the levels of circulating IL-6 that rise with disease severity.</li> <li>- PD-1 receptor was upregulated on NK lymphocytes in COVID-19 patients, and several pneumonia and ARDS COVID-19 patients had a particularly large subset of NK cells expressing PD-1.</li> <li>- Bronchoalveolar lavage fluid (BALF) analysis showed a lack of CD16+CD57+ mature NK cells in the lungs of ARDS patients, suggesting that the decrease in mature NK cell levels observed in blood is not a consequence of their migration to infected lungs. In addition, high levels of CD39, PD-1, and NKG2A expression were also observed in NK cells isolated from the BALF of ARDS COVID-19 patients.</li> <li>- NK cells isolated from the blood of ARDS COVID-19 patients retained cytotoxic functions, and that incubation with monalizumab an anti-NKG2A mAb blocking the inhibitory interaction with HLA-E, was able to unleash their killing ability.</li> </ul> <p>These data suggest that NK cells do not participate in the exaggerated inflammatory response observed in ARDS. Thus, therapies targeting PD-(L)1, NKG2A and CD39 should be investigated as means of boosting NK cell antiviral immunity in patients at early stages of SARS-CoV-2 infection.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
J Intern Med 01JUL2020	<b>Treatment with proton pump inhibitors increases the risk of secondary infections and ARDS in hospitalized patients with COVID-19: coincidence or underestimated risk factor?</b>	Luxemburger, H. et al. Germany <a href="#">gotopaper</a>	Therapeutic	<p>Retrospective study, 152 hospitalized patients with confirmed SARS-Cov-2 infection were included in the analysis. 62 patients (40.8%) received regular treatment with proton pump inhibitors (PPI). 48 patients (31.6%) presented with a secondary infection during hospitalization.</p> <ul style="list-style-type: none"> <li>- PPI-treated patients with COVID-19 presented more often with secondary infections compared to patients without PPI treatment. This effect remained statistically significant after adjusting for other possible risk factors.</li> <li>- Secondary infections were strongly associated with the development of ARDS indicating an indirect negative impact of PPI treatment on the development of ARDS.</li> <li>- Index mortality was higher in patients with PPI treatment.</li> </ul> <p>Ø PPI treatment may be a negative predictive factor for development of secondary infections and consecutive ARDS in patients with COVID-19.</p> <p>Limitations: retrospective study, not able to analyse the effect of the duration of PPI treatment on the outcome of SARS-CoV2-infected patients.</p>
MICROBIOLOGY AND INFECTION 26JUN2020	<b>Childhood COVID-19: a multi-center retrospective study</b>	Chen Z et al- CHINA <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Description of clinical and epidemiological characteristics of pediatric patients with COVID19</p> <p>Cohort of 32 children from 3 months to 18 years old (China).</p> <ul style="list-style-type: none"> <li>&gt; Family aggregation occurred in 87.5% of infant and preschool/school-aged, but only 12.5% of adolescents.</li> <li>&gt; Most common symptoms (mild): fever, cough, fatigue (4/32).</li> <li>&gt; Average duration of viral RNA in respiratory samples 15.8 d</li> <li>&gt; Average duration of viral RNA in gastrointestinal samples: 28.9d</li> <li>&gt; 14 children developed pneumonia, but no statistical significance in the incidence between age groups.</li> </ul> <p>Most children with COVID-19 had a mild process and good prognosis. More attention should be paid to household contact history investigation</p>
LANCET PHSYCHIATRY 25JUN2020	<b>Neurological and neuropsychiatric complications of COVID-19 in 153 patients: a UK-wide surveillance study</b>	Varatharaj A et al UK and USA <a href="#">gotopaper</a>	Public Health/Epidemio	<p>The aim of this study was to investigate the breadth of complications of COVID-19 across the UK that affected the brain</p> <p><b>METHODS</b></p> <ul style="list-style-type: none"> <li>&gt; development of a rapid case report notifications portal across UK neurosciences bodies</li> <li>&gt; clinical syndromes associated with COVID-19 were classified as a cerebrovascular event, altered mental status, peripheral neurology, or other.</li> </ul> <p><b>RESULTS</b></p> <p>During the study period, platforms received notification of 153 cases that met the clinical case definitions (with an exponential growth in reported cases similar to overall COVID-19 data). Complete clinical datasets were available for 125 (82%) of 153 patients (median age 71 years)</p> <ul style="list-style-type: none"> <li>&gt; 77 (62%) of 125 patients presented with a cerebrovascular event,</li> <li>&gt; 39 (31%) of 125 patients presented with altered mental status,</li> <li>&gt; 9 (23%) with unspecified encephalopathy</li> <li>&gt; 7 (18%) with encephalitis.</li> <li>&gt; 23 (59%) fulfilled the clinical case definitions for psychiatric diagnoses (mostly new onset disorders)</li> </ul>



Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA network open 24JUN2020	<b>Effect of Colchicine vs Standard Care on Cardiac and Inflammatory Biomarkers and Clinical Outcomes in Patients Hospitalized With Coronavirus Disease 2019: The GRECCO-19 Randomized Clinical Trial</b>	Deftereos, Spyridon G. et al. Greece <a href="#">gotopaper</a>	Therapeutic	<p>Prospective, open-label, multicenter randomized clinical trial, involving 105 patients hospitalized with COVID-19 randomized in a 1:1 allocation to either standard medical treatment or colchicine with standard medical treatment. Primary endpoints were (1) maximum high-sensitivity cardiac troponin level; (2) time for C-reactive protein to reach more than 3 times the upper reference limit; and (3) time to deterioration by 2 points on a 7-grade clinical status scale, ranging from able to resume normal activities to death.</p> <p><b>Participants who received colchicine had statistically significantly improved time to clinical deterioration. There were no significant differences in high-sensitivity cardiac troponin or C-reactive protein levels.</b></p> <p>Limitations: study allowed for co-treatment with other investigational agents ; small sample size combined with low event rate make data underpowered and hypothesis generating.</p>
THE LANCET CHILD & ADOLESCENT HEALTH 25JUN2020	<b>COVID-19 in children and adolescents in Europe: a multinational, multicentre cohort study</b>	Göttinger Florian et al- 25 European countries. <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Factors associated with need for ICU admission and initiation of drug treatment for COVID-19 for children and adolescents with SARS-CoV-2.</p> <p>METHODS - Multicentre cohort (582 individuals). Children under 18 years of age (median 5 IQR 0.5–12.0) with confirmed SARS-CoV-2 Infection (PCR). 82 participating health-care institutions; 25 European countries (Paediatric Tuberculosis Network European Trials Group)</p> <p>RESULTS : Four children died.          &gt; 145 (25%) had pre-existing medical conditions.          &gt; 363 (62%) individuals were admitted to hospital.          &gt; 48 (8%) individuals required ICU admission,          &gt; 25 (4%) mechanical ventilation          &gt; 19 (3%) inotropic support,          &gt; 1 (&lt;1%) extracorporeal membrane oxygenation</p> <p>Significant risk factors for requiring ICU admission were:          &gt; being younger than 1 month          &gt; male sex          &gt; pre-existing medical conditions          &gt; presence of lower respiratory tract infection signs          &gt; symptoms at presentation</p>
Cell Metabolism 24JUN2020	<b>In-hospital Use of Statins is Associated with a Reduced Risk of Mortality among Individuals with COVID-19</b>	Zhang, Xiao-Jing et al. China <a href="#">gotopaper</a>	Therapeutic	<p>Retrospective multicenter study on 13,981 clinically confirmed cases of COVID-19, among which 1,219 received statins, to determine the association of in-hospital use of statins with clinical outcomes. In a subgroup analysis, the additional effects of combining ACEi/ARB with statins on the clinical outcomes of COVID-19 was investigated.</p> <ul style="list-style-type: none"> <li>- In-hospital use of statins was associated with a lower risk of all-cause mortality: the risk for 28-day all-cause mortality was 5.2% and 9.4% in the matched statin and non-statin groups, respectively, with an adjusted hazard ratio of 0.58.</li> <li>- The combination of statins and ACEi/ARB utilization was not significantly associated with the risk of all-cause mortality among individuals with COVID-19 and hypertension.</li> <li>- In-hospital usage of statins and combination usage of statins and ACE inhibitors or ARBs did not increase the risk of organ damage and other adverse effects.</li> </ul> <p>Limitation: retrospective study</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
American Journal of Obstetrics and Gynecology 23JUN2020	<b>Exposure and Seroconversion to SARS-CoV-2 Among Obstetric Healthcare Providers Following a Contained Outbreak</b>	KIEFER MK et al, USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Key question: Exposure and seroconversion to SARS-CoV-2 among obstetric HCWs (in a tertiary care center)</p> <p>&gt; Obstetric units are an underestimated hotspot in the pandemic, owing to an asymptomatic population, high patient turnover, integrated workstations, and frequent emergencies requiring response from multiple disciplines and expedient transfer to onsite surgical suites</p> <p>STUDY DESIGN - Prospective cohort of HCW: study of SARS-CoV-2 antibody levels Collection of Blood samples at two time points four weeks apart (IgM and IgG levels) RESULTS - 110 HCW recruited (females, median age 34). 68,2% nurses, 24,5% physicians</p> <p>&gt; 90 participants (82%) reported a SARS-CoV-2 exposure &gt; 66 (60%) reported one or more COVID-19 symptoms; &gt; 52 (47%) had nasopharyngeal PCR testing, of which 15 (29%) tested positive for the virus, 14 seroconverted &gt; at baseline, 3 participants (2,7%) had positive antibodies &gt; 5 participants (4.5%) who reported being asymptomatic, seroconverted</p>
Blood 23JUN2020	<b>Improved Clinical Symptoms and Mortality on Severe/Critical COVID-19 Patients Utilizing Convalescent Plasma Transfusion</b>	Xia, Xinyi et al. China <a href="#">gotopaper</a>	Therapeutic	<p>Evaluation of the effectiveness, safety, and indications of convalescent plasma transfusion (CPT) therapy for severe or critical COVID-19 patients, through analysis of the clinical, laboratory, and radiologic characteristics of 1,568 patients from a single center, in which 138 patients received ABO-compatible CPT and 1,430 patients received standard treatment.</p> <ul style="list-style-type: none"> <li>- 2.2% and 4.1% of cases died in the CPT group and in the standard-treatment group, respectively.</li> <li>- 2.4% and 5.1% of patients in the CPT and the standard-treatment group have been admitted to ICU eventually.</li> <li>- 70% of the patients who had severe respiratory symptoms got improved and removed oxygen supports within 7 days after CPT.</li> <li>- The viral loads and C-reactive protein (CRP) concentration significantly decreased (<math>P&lt;0.001</math>), and the percentage of lymphocytes increased (<math>P=0.006</math>), 76.8% of cases received radiological improvements within 14 days after CPT.</li> <li>- Patients with a higher percentage of lymphocytes and a lower percentage of neutrophils and CRP concentration respond better to CPT (<math>P&lt;0.05</math>).</li> <li>- Notably, for the patients who received CPT within 7 weeks after symptom onset, the median time from CPT to clinical improvements was approximately 10 days. But the time to clinical improvements was significantly prolonged for patients who received CPT later than 7 weeks after onset.</li> </ul> <p><b>The results suggest CCP, transfused even after two weeks of symptom onset, could improve the symptoms and mortality in severe or critical COVID-19 patients.</b></p> <p>Limitations: single center, retrospective study ; complete data on neutralizing antibody titers in CCP units were not available ; a stratified analysis of severe and critical patients could not be performed due to the low proportion of critical patients.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
CELL 23JUN2020	<b>A universal design of betacoronavirus vaccines against COVID-19, MERS and SARS</b>	Lianpan D et al, CHINA <a href="#">gotopaper</a>	Vaccine	<p>Design of a dimeric form of MERS-CoV RBD inducing high immunogenicity and significantly increased neutralizing antibody (NAb) titers compared to conventional monomeric form.</p> <p>&gt; Crystal structures of the RBD-dimer shows fully exposed dual receptor-binding motifs which are the targets of NABs</p> <p>Strategy (RBD dimers) to design a SARSCoV2 vaccine candidate</p> <p>&gt; Selected sequences: from R319 to K537</p> <p>&gt; Dimerization and binding to hACE was demonstrated</p> <p>&gt; Immunization of BABL/c mice with RBD +adjuvant induced significantly higher antigen-specific IgG compared to immunization with RBD-monomer</p> <p>&gt; RBD dimers elicited ~10-100-fold higher titer of NAb compare to monomer</p> <p>&gt; No differences in T-cell responses in of RBDdimers-vaccinated mice compared to the PBS-vaccinated ones.</p> <p>Expression on SARS CoV 2 RBD dimers in clinical grade CHO cells line reached expression levels of &gt; 1.5 g/L, with a final yield of 0.67 gram purified antigen per liter</p> <p>&gt; Highly scalable production</p>
Science 23JUN2020	<b>A mathematical model reveals the influence of population heterogeneity on herd immunity to SARS-CoV-2</b>	Britton et al., Sweden and UK <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Population heterogeneity can significantly impact disease-induced immunity as the proportion infected in groups with the highest contact rates is greater than in groups with low contact rates</p> <p>-&gt; if <math>R_0 = 2.5</math> in an age-structured community with mixing rates fitted to social activity then the disease-induced herd immunity level can be around 43%, which is substantially less than the classical herd immunity level of 60% obtained through homogeneous immunization of the population</p> <p>These estimates should be interpreted as an illustration of how population heterogeneity affects herd immunity, rather than an exact value or even a best estimate.</p>
J Infect Dis 19JUN2020	<b>Interferon-beta 1a inhibits SARS-CoV-2 in vitro when administered after virus infection</b>	Clementi, Nicola et al. Italy <a href="#">gotopaper</a>	Therapeutic	<p>Evaluation of anti-SARS-CoV-2 in vitro activity of Interferon-beta 1a.</p> <p>Vero E6 cells treated with 5,000 to 0.01 IU/mL of IFN-<math>\beta</math> 1a 1 h after inoculation with the SARS-CoV-2 (MOI of 0.001) and monitored for cytopathic effect and q-RT-PCR evaluation at 48, 72 and 96 hpi.</p> <p>- Inhibition of the SARS-CoV-2 by IFN-<math>\beta</math> 1a was dependent on both time and drug concentration. No altered cell morphology related to drug toxicity was observed on uninfected cells treated with IFN-<math>\beta</math> 1a at 5000 IU/mL.</p> <p>- EC50 calculations at different time points, resulting in 1.971 IU/mL (95% CI: 0.3969 to 4.891 IU/mL) at 48 hpi, 2.071 IU/mL (95% CI: 0.5982 to 5.819 IU/mL) at 72 hpi, and 4.682 IU/mL (95% CI: 3.505 to 6.018 IU/mL) at 96 hpi. Can be easily accessed in the clinical setting.</p> <p>=&gt; Shed light for the first time on antiviral activity of IFN-<math>\beta</math> 1a against SARS-CoV-2 when administered after the infection of cells, highlighting its possible efficacy in an early therapeutic setting. IFN-<math>\beta</math> 1a activity is retained up to 96 hours after its use on the infected cells.</p> <p>Preclinical evaluation of the antiviral activity of a drug, such as IFN-<math>\beta</math> 1a, is only a partial assessment of its possible clinical role in a disease such as COVID-19 in which the beneficial or detrimental effect of type I IFN is still to be established and in which immune-mediated damage is probably an extremely important factor in determining the development of the worst outcomes of the infection.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
VACCINE 19JUN2020	<b>Recombinant SARS-CoV-2 spike S1-Fc fusion protein induced high levels of neutralizing responses in nonhuman primates</b>	Ren W et al_China <a href="#">gotopaper</a>	Vaccine	<p>Immunogenicity of SARS-CoV-2 S1 Fc fusion proteins as potential vaccine candidate</p> <ul style="list-style-type: none"> <li>&gt; the S1 is located at the N-terminus of the S protein and contains the RBD</li> <li>&gt; the S1 was fused with the Fc region of human IgG1 and expressed as a recombinant protein in CHO-K1 cell line</li> <li>&gt; the S1-Fc protein was formulated with different adjuvants and tested as vaccine candidate</li> </ul> <p>Injection of S1 Fc protein in mice, rabbit and macaques elicited</p> <ul style="list-style-type: none"> <li>&gt; high levels of anti-S1 antibodies in all animal models</li> <li>&gt; high neutralizing activities against SARS-CoV-2 (in the antisera of macaques)</li> </ul> <p>These results indicate that the S1-Fc fusion protein can effectively induce humoral immune responses in various animals models and may be a good vaccine candidate.</p>
JAMA 19JUN2020	<b>Association of Angiotensin-Converting enzyme inhibitor or angiotensin receptor blocker use with COVID-19 diagnosis and mortality</b>	Fosbøl EL et al Danish <a href="#">gotopaper</a>	Clinic	<p>Use of ACEI/ARBs was associated with COVID-19 diagnosis and worse outcomes in patients with COVID-19?</p> <p>Retrospective cohort study (data from Danish national administrative registries)</p> <p>4480 patients included: 895 users of ACEI/ARBs and 3585 nonusers</p> <p>Nested case-control framework: association use of ACEI/ARBs vs other antihypertensive drugs and incidence rate of a COVID-19 diagnosis</p> <ul style="list-style-type: none"> <li>- 571 patients prior hypertension + COVID-19</li> <li>- 570 controls with prior hypertension no COVID-19</li> </ul> <p>Primary outcome: death</p> <ul style="list-style-type: none"> <li>- 18,1% in ACEI/ARBs group vs 7,3% within 30-days, HRa: 0,83 IC95%[0,67 – 1,03]</li> </ul> <p>Secondary outcome: death or severe COVID-19</p> <ul style="list-style-type: none"> <li>- 31,9% in ACEI/ARBs group vs 14,2% in nonusers, HRa: 1,04 IC95%[0,89 – 1,23]</li> </ul> <p>Nested case-control framework:</p> <ul style="list-style-type: none"> <li>- 86,5% of cases used ACEI/ARBs versus 85,4% of controls</li> <li>- ACEI/ARBs not associated with higher incidence of COVID-19, HRa: 1,05 IC95%[0,80 – 1,36].</li> </ul> <p>Limits: observational study – laboratory data were not available – exposure to ACEI/ARBs was defined by prescription fillings – confounding by indication treatment</p> <p>Prior use of ACEI/ARBs was not significantly associated with COVID-19 diagnosis among patients with hypertension or with mortality or severe disease</p>
Nature Medicine 18JUN2020	<b>Clinical and immunological assessment of asymptomatic SARS-CoV-2 infections</b>	Long, Quan-Xin, et al. China <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>AIM: to describe the clinical features of study 37 SARS-CoV-2 positive but asymptomatic individuals in the Wanzhou District.</p> <ul style="list-style-type: none"> <li>- Median duration of viral shedding in the asymptomatic group (19d) significantly longer than in the symptomatic group (14d in patients with mild symptoms).</li> <li>- Virus-specific IgG levels in the asymptomatic group significantly lower relative to the symptomatic group in the acute phase. In the early convalescent phase, 93.3% and 81.1% of asymptomatic individuals had reduction in IgG and neutralizing antibody levels respectively, as compared to 96.8% and 62.2% of symptomatic patients.</li> <li>- 40% of asymptomatic individuals and 12.9% of the symptomatic group became seronegative in the early convalescent phase. Asymptomatic individuals exhibited lower levels of 18 pro- and anti-inflammatory cytokines.</li> </ul> <p><b>These data suggest that asymptomatic individuals had a weaker immune response to SARS-CoV-2 infection, that might have implications for immunity strategy and serological surveys.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
EClinicalMedicine 18JUN2020	<b>Tocilizumab for treatment of patients with severe COVID-19: A retrospective cohort study</b>	Kewan T et al USA <a href="#">gotopaper</a>	Therapeutic	<p>Retrospective cohort study – hypoxic COVID-19 patients – patients with lung infiltrates and elevated inflammatory markers received a single dose of tocilizumab</p> <p>51 patients included: 28 received tocilizumab and 23 did not received</p> <p>At baseline:</p> <ul style="list-style-type: none"> <li>- More mechanical ventilation in tocilizumab cohort: 68% vs 22%</li> </ul> <p>Clinical improvement: tocilizumab vs non tocilizumab cohort</p> <ul style="list-style-type: none"> <li>- All patients: 6,5 days versus 7 days, HR:1,14 IC95%[0,55 – 2,38] / Mechanically ventilated patients: 8 days versus 13 days, HR:1,83 IC95%[0,57 – 5,84]</li> </ul> <p>Duration of vasopressor support</p> <ul style="list-style-type: none"> <li>- 2 days in tocilizumab cohort versus 5 days (p=0,039)</li> </ul> <p>Duration of mechanical ventilation / 7 days in tocilizumab cohort versus 10 days (p=0,11)</p> <p>Limits: retrospective / single center with small number of patients / tocilizumab cohort was younger / short follow-up / concomitant use of other drugs.</p> <p>-&gt; Tocilizumab was associated with shorter duration of vasopressor support and shorter median time to clinical improvement (not statistically significant)</p> <p>-&gt; Use of tocilizumab in select patients with severe COVID-19 ?</p>
New England Journal of Medicine 17JUN2020	<b>Genomewide Association Study of Severe Covid-19 with Respiratory Failure</b>	Ellinghaus, D., et al. Multicentre <a href="#">gotopaper</a>	Public Health/Epidemio	<p>AIM: performing a Genomewide association analysis on patients with severe disease and controls (Italy/Spain) to identify potential genetic factors involved in the development of Covid-19.</p> <ul style="list-style-type: none"> <li>- Significant cross-replicating associations with rs11385942 at locus 3p21.31 and with rs657152 at locus 9q34.2 were detected.</li> <li>- At locus 3p21.31, the association signal spanned six genes</li> <li>- The association signal at locus 9q34.2 coincided with the ABO blood group locus; a blood-group-specific analysis showed a higher risk in blood group A and a protective effect in blood group O, as compared with other blood groups.</li> </ul> <p>The 3p21.31 gene cluster identified as a genetic susceptibility locus in patients with severe Covid-19 and confirmed a potential involvement of the ABO blood-group system.</p>
The Lancet. Infectious diseases 16JUN2020	<b>Effectiveness of isolation, testing, contact tracing, and physical distancing on reducing transmission of SARS-CoV-2 in different settings: a mathematical modelling study</b>	Kucharski, AJ, et al. UK <a href="#">gotopaper</a>	Public Health/Epidemio	<p>AIM: to estimate the reduction in SARS-CoV-2 transmission under different control measures (testing, isolation, tracing, physical distancing) across settings (household, work, school, or other ) and the number of contacts to quarantine daily in different strategies for a given symptomatic case incidence.</p> <ul style="list-style-type: none"> <li>- Combined isolation and tracing strategies could reduce transmission more than mass testing or self-isolation alone: mean transmission reduction of 2% for weekly random testing of 5% of the population, 29% for self-isolation alone of symptomatic cases within the household, 35% for self-isolation alone outside the household, 37% for self-isolation plus household quarantine, 64% for self-isolation and household quarantine plus manual contact tracing of all contacts, 57% plus manual tracing of acquaintances only, 47% plus app-based tracing only.</li> <li>- If limiting gatherings outside of home, school, or work, manual contact tracing alone could reduce transmission similarly to detailed contact tracing.</li> <li>- If 1000 new symptomatic cases occurred daily, 15 000–41 000 contacts per day would be quarantined through contact tracing.</li> </ul> <p>Many cases would need to self-isolate and their contacts successfully traced to ensure a <math>R_0 &lt; 1</math> in the absence of other measures. If combined with moderate physical distancing, self-isolation and contact tracing would be more likely to be effective.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Lancet Rheumatol 16JUN2020	<b>GM-CSF blockade with mavrilimumab in severe COVID-19 pneumonia and systemic hyperinflammation: a single-centre, prospective cohort study</b>	De Lucas G et al Italy <a href="#">gotopaper</a>	Therapeutic	<p>Single center prospective cohort study, open label – Adults patients with severe COVID-19, hypoxemia &amp; systemic hyperinflammation</p> <p>2 groups:</p> <ul style="list-style-type: none"> <li>- Standard of care (26 patients)</li> <li>- Standard of care + mavrilimumab IV (13 patients)</li> </ul> <p>At inclusion time, none of the patient included was mechanically ventilated</p> <p>Main outcome: time to clinical improvement (ordinal scale of clinical status)</p> <p>At day 28</p> <ul style="list-style-type: none"> <li>- 100% clinical improvement in mavrilimumab group vs 65% control group (p=0,03)</li> <li>- Earlier improvement in mavrilimumab group (8 days vs 19 days, p=0,0001)</li> <li>- 0% death in mavrilimumab group vs 27% in control group (p=0,086)</li> <li>- 8% progressed to MV in mavrilimumab group vs 35% in control group (p=0,14)</li> </ul> <p>Fever resolution was faster in mavrilimumab group</p> <p>Mavrilimumab was well tolerated.</p> <p>Several limitations</p> <ul style="list-style-type: none"> <li>- Not randomly assigned</li> <li>- Other clinical variables besides mavrilimumab might have affected clinical outcomes</li> <li>- Patients in mavrilimumab group had a longer duration fever before enrolment</li> </ul> <p>Mavrilimumab was associated with clinical improvement compared with standard care in non-mechanically ventilated patients</p> <p>These results require placebo-controlled studies/multicenter/double-blind/randomised</p>
Nature Medicine 16JUN2020	<b>Age-dependent effects in the transmission and control of COVID-19 epidemics</b>	Davies et al., UK <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Age-structured mathematical model to epidemic data from China, Italy, Japan, Singapore, Canada and South Korea.</p> <p><b>Estimation:</b></p> <ul style="list-style-type: none"> <li>- Susceptibility to infection in individuals under 20 years of age is approximately half that of adults aged over 20 years</li> <li>- Clinical symptoms manifest in 21% (95% credible interval: 12–31%) of infections in 10- to 19-year-olds, rising to 69% (57–82%) of infections in people aged over 70 years</li> </ul> <p>-&gt; interventions aimed at children might have a relatively small impact on reducing SARS-CoV-2 transmission, particularly if the transmissibility of subclinical infections is low</p> <p><b>Conclusion:</b></p> <p>-&gt; In countries with younger population structures such as many low-income countries the expected per capita incidence of clinical cases would be lower than in countries with older population structures, although it is likely that comorbidities in low-income countries will also influence disease severity.</p> <p>-&gt;Without effective control measures, regions with relatively older populations could see disproportionately more cases of COVID-19, particularly in the later stages of an unmitigated epidemic.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
CELL 15JUN2020	<b>Structures of human antibodies bound to SARS-CoV-2 spike reveal common epitopes and recurrent features of antibodies</b>	Christopher O et al, USA/SWITZERLAND <a href="#">gotopaper</a>	Vaccine/Therapeutics	<p>Characterization of polyclonal IgGs and Fabs from COVID-19 convalescent individuals for recognition of coronavirus spikes.</p> <p>Analysis of purified IgG and Fabs from the plasmas of 10 COVID-19 convalescent individuals showed</p> <ul style="list-style-type: none"> <li>&gt; binding to trimeric S and monomeric RBD/S1B domains of six human coronaviruses</li> <li>&gt; neutralizing activity against SARS-CoV-2 pseudoviruses</li> </ul> <p>By using EM approaches it was shown that:</p> <ul style="list-style-type: none"> <li>&gt; Fabs recognize both S1A and RBD epitopes on SARS-CoV-2 S protein</li> <li>&gt; the monoclonal Fab-spike neutralizing complex passes through a specific epitope that blocks ACE2 receptor binding</li> </ul> <p>Overall, these studies structurally define a recurrent anti-SARSCoV-2 antibody class derived from VH3-53/VH3-66 and similarity to a SARS-CoV VH3-30 antibody, providing criteria for evaluating vaccine-elicited antibodies.</p>
Science 15JUN2020	<b>Antibody cocktail to SARS-CoV-2 spike protein prevents rapid mutational escape seen with individual antibodies</b>	Baum, Alina et al. USA <a href="#">gotopaper</a>	Therapeutic	<p>Investigate the <b>development of resistance against 4 antibodies to the spike protein</b> that potentially neutralize SARS-CoV-2, <b>individually</b> as well as when <b>combined into cocktails</b>.</p> <ul style="list-style-type: none"> <li>- Efficacy of antiviral antibodies against the breadth of spike RBD variants represented in publicly available SARS-CoV-2 sequences was assessed using VSV pseudoparticle system expressing the SARS-CoV-2 spike variants =&gt; <b>antibodies remain effective against spike variants that have arisen in the human population</b>.</li> <li>- Escape mutants were selected under pressure of single antibodies, as well as of antibody combinations, by using a replicating VSV-SARS-CoV-2-S virus =&gt; <b>novel spike mutants rapidly appeared following in vitro passaging in the presence of individual antibodies, resulting in loss of neutralization</b>; such escape <b>also occurred with combinations of antibodies binding diverse but overlapping regions of the spike protein</b>.</li> <li>- Escape following treatment with an antibody cocktail (REGN10987+REGN10933) was assessed, cocktail being rationally designed to avoid escape through inclusion of two antibodies that bind distinct and non-overlapping regions of the RBD, and which can thus simultaneously bind and block RBD function =&gt; <b>attempts to grow VSV-SARS-CoV-2-S virus in the presence of this non-competing antibody cocktail did not result in the outgrowth of escape mutants</b>.</li> </ul> <p>⇒ <b>Cocktail therapy may provide a powerful way to minimize mutational escape by SARS-CoV-2</b>; in particular when two antibodies were chosen so as to bind to distinct and non-overlapping regions of the viral target, and thus require the unlikely occurrence of simultaneous mutations at two distinct genetic sites for viral escape</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Science 15JUN2020	<b>Potent neutralizing antibodies from COVID-19 patients define multiple targets of vulnerability</b>	Brouwer, Philip J. M. et al. Netherlands/USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Isolation of monoclonal antibodies from 3 convalescent COVID-19 patients</b> using a SARS-CoV-2 stabilized prefusion spike protein. These antibodies had <b>low levels of somatic hypermutation</b> and showed a strong enrichment in VH1-69, VH3-30-3 and VH1-24 gene usage.</p> <ul style="list-style-type: none"> <li>- 19 NAb that target a diverse range of antigenic sites on the S protein, of which 2 showed picomolar neutralizing activities (<b>IC50s of 0.007 and 0.009 µg/mL</b> or 47 and 60 pM) against <b>authentic SARS-CoV-2 virus</b>.</li> <li>- Large-scale SPR-based competition assays allowed to define NAb that <b>target multiple sites of vulnerability on the RBD</b> as well as additional previously <b>undefined non-RBD epitopes</b> on SARS-CoV-2.</li> </ul> <p>⇒ Subsequent structural characterization of these potent NAb will <b>guide vaccine design</b>, while simultaneous targeting of <b>multiple non-RBD and RBD epitopes with mAb cocktails</b> paves the way for safe and effective COVID-19 prevention and treatment.</p>
Science 15JUN2020	<b>Studies in humanized mice and convalescent humans yield a SARS-CoV-2 antibody cocktail</b>	Hansen, Johanna et al. USA <a href="#">gotopaper</a>	Therapeutic	<p>Describe parallel efforts using both <b>humanized mice and convalescent patients to generate antibodies against the SARS-CoV-2 spike protein</b>, yielding a large <b>collection of fully-human antibodies</b> that were characterized for <b>binding, neutralization and three dimensional structure</b>.</p> <ul style="list-style-type: none"> <li>- screened for binding affinity to RBD monomer and dimer, epitope diversity, ability to block ACE2 receptor binding to RBD, and neutralize VSV-based SARS-CoV-2 spike pseudoparticles (pVSV-SARS-CoV-2-S(mNeon)). =&gt; Screening yielded over 200 neutralizing mAbs with broad potency ranges, dozens of which displayed neutralization potency in the pM range.</li> <li>- A smaller collection of 4 antibodies was chosen for further analyses to determine whether the above binding data to RBD reflected binding to trimeric spike protein, whether neutralization potencies noted in the above assays were consistent with those seen in other assays including with SARS-CoV-2, and whether these antibodies retained neutralization activity against pseudoparticles with mutations in the S1/S2 cleavage site.</li> <li>- examined 9 most potent neutralizing antibodies in cross-competition binding assays, identifying several pairs of non-competing mAbs with pM neutralization potency that could potentially be combined to form antibody cocktails.</li> </ul> <p>=&gt; Selection of pairs of highly-potent individual antibodies that simultaneously bind the receptor-binding domain of the spike protein, providing ideal partners for a therapeutic antibody cocktail that may deliver optimal antiviral potency while minimizing odds of virus escape.</p> <p>Such an <b>antibody cocktail is now being tested in human trials</b> (clinicaltrials.gov NCT04426695 and NCT04425629).</p>
Science 15JUN2020	<b>Broad neutralization of SARS-related viruses by human monoclonal antibodies</b>	Wec, Anna Z. et al. USA <a href="#">gotopaper</a>	Therapeutic	<p>Mined the memory B cell repertoire of a <b>convalescent SARS donor</b> and identified <b>200 SARS-CoV-2 binding antibodies</b> that <b>target multiple conserved sites on the spike (S) protein</b>.</p> <ul style="list-style-type: none"> <li>- A large proportion of the <b>non-neutralizing antibodies</b> display <b>high levels of somatic hypermutation</b> and <b>cross-react with circulating HCoVs</b>, suggesting <b>recall of pre-existing memory B cells (MBCs)</b> elicited by <b>prior HCoV infections</b>.</li> <li>- Several antibodies <b>potently cross-neutralize SARS-CoV, SARS-CoV-2, and the bat SARS-like virus WIV1</b> by blocking receptor attachment and inducing S1 shedding.</li> </ul> <p>⇒ These antibodies represent <b>promising candidates for therapeutic intervention</b> and reveal a target for the <b>rational design of pan-sarbecovirus vaccines</b>.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet GH 15JUN2020	Global, regional, and national estimates of the population at increased risk of severe COVID-19 due to underlying health conditions in 2020: a modelling study	Clark et al., UK, China, USA <a href="#">gotopaper</a>	Public Helath/Epidemio	<p>Aim: Understanding the number of individuals at increased risk of severe COVID-19 and how this varies between countries</p> <p><b>Estimation:</b></p> <ul style="list-style-type: none"> <li>-&gt; 1.7 billion people, comprising 22% of the global population, have at least one underlying condition that puts them at increased risk of severe COVID-19 if infected</li> <li>-&gt; 349 million people (4% of the global population) are at high risk of severe COVID-19 and would require hospital admission if infected</li> <li>-&gt; 6% of males to be at high risk compared with 3% of females.</li> </ul> <p>The share of the population at increased risk was highest in countries with older populations, African countries with high HIV/AIDS prevalence, and small island nations with high diabetes prevalence.</p> <p>Estimates of the number of individuals at increased risk were most sensitive to the prevalence of chronic kidney disease, diabetes, cardiovascular disease, and chronic respiratory disease.</p> <p><b>-&gt; About one in five individuals worldwide could be at increased risk of severe COVID-19</b></p>
Science 15JUN2020	Isolation of potent SARS-CoV-2 neutralizing antibodies and protection from disease in a small animal model	Rogers, Thomas F. et al. USA <a href="#">gotopaper</a>	Therapeutic	<p>Cohort of SARS-CoV-2-recovered participants, neutralization assays to interrogate antibody responses, high-throughput antibody generation pipeline to rapidly screen over 1800 antibodies, and animal model to test protection.</p> <p>Only a small fraction of these Abs was neutralizing, highlighting the value of deep mining of responses to access the most potent Abs.</p> <ul style="list-style-type: none"> <li>- <b>Potent neutralizing antibodies (nAbs) to two epitopes on the receptor binding domain (RBD) and to distinct non-RBD epitopes</b> on the spike (S) protein isolated.</li> <li>- <b>Passive transfer of a nAb provides protection against disease</b> in high-dose SARS-CoV-2 challenge <b>in Syrian hamsters</b>, as revealed by maintained weight and low lung viral titers in treated animals.</li> </ul> <p>⇒ The study suggests a <b>role for nAbs in prophylaxis, and potentially therapy</b>, of COVID-19. The nAbs <b>define protective epitopes to guide vaccine design</b>.</p>
Cell research 15JUN2020	Boceprevir, GC-376, and calpain inhibitors II, XII inhibit SARS-CoV-2 viral replication by targeting the viral main protease	Ma, Chunlong et al. USA <a href="#">gotopaper</a>	Therapeutic	<p>This study identified several <b>potent SARS-CoV-2 Mpro inhibitors with potent enzymatic inhibition as well as cellular antiviral activity</b>.</p> <ul style="list-style-type: none"> <li>- <b>Boceprevir</b>, an FDA-approved HCV drug, inhibits the enzymatic activity of Mpro with <b>IC50 of 4.13 µM</b>, and has an <b>EC50 of 1.90 µM</b> against the SARS-CoV-2 virus in the cellular viral CPE assay.</li> <li>- <b>GC-376</b>, an investigational veterinary drug, showed promising antiviral activity against the SARS-CoV-2 virus (<b>EC50 = 3.37 µM</b>). It has the highest enzymatic inhibition against the Mpro with an <b>IC50 value of 0.03 µM</b>. The <b>X-ray crystal structure</b> of SARS-CoV-2 Mpro in complex with GC-376 provides a molecular explanation of the <b>high binding affinity</b> of aldehyde-containing compounds as they can adopt two configurations R and S.</li> <li>- 3 calpain/cathepsin inhibitors, <b>MG-132, calpain inhibitors II and XII</b>, are potent inhibitors of Mpro with <b>single-digit to submicromolar efficacy</b> in the enzymatic assay. Calpain inhibitors II and XII also inhibit SARS-CoV-2 in the CPE assay with <b>EC50 values of 2.07 and 0.49 µM</b>, respectively.</li> <li>- <b>Ritonavir and lopinavir failed to inhibit the SARS-CoV-2 Mpro</b> (IC50 &gt; 20 µM), which might explain why they lack efficacy in clinical trials for COVID-19.</li> <li>- <b>Camostat has no inhibition against the SARS-CoV-2 Mpro</b> (IC50 &gt; 20 µM).</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
SCIENCE 12JUN2020	<b>The impact of COVID-19 and strategies for mitigation and suppression in low- and middle-income countries</b>	Walker et al., UK, USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Combining data on demography, contact patterns, disease severity, and health care capacity and quality to understand its impact and inform strategies for its control</p> <p>-&gt; Younger populations in lower income countries may reduce overall risk</p> <p>-&gt; Limited health system capacity coupled with closer inter-generational contact largely negates this benefit.</p> <p>-&gt; Mitigation strategies that slow but do not interrupt transmission will still lead to COVID-19 epidemics rapidly overwhelming health systems, with substantial excess deaths in lower income countries due to the poorer health care available.</p>
The Lancet 11JUN2020	<b>Seroprevalence of anti-SARS-CoV-2 IgG antibodies in Geneva, Switzerland (SEROCoV-POP): a population-based study</b>	Stringhini, S. et al. Switzerland <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>In SEROCov-POP, a population-based study, we planned a series of 12 consecutive weekly serosurveys of anti-SARS-CoV-2-IgG antibodies among randomly selected participants and their household members in Geneva, Switzerland (2766 participants from 2339 households), and estimated the seroprevalence in the population. We present results from the first 5 weeks of the study.</p> <p>- In the first week, we estimated a seroprevalence of 4.8%. The estimate increased to 8.5% in the second week, to 10.9% in the third week, 6.6% in the fourth week, and 10.8% in the fifth week.</p> <p>- Individuals aged 5–9 years and those older than 65 years had a significantly lower risk of being seropositive than those aged 20–49 years. We estimated that for every reported confirmed case, there were 11.6 infections in the community.</p> <p>These results suggest that most of the population of Geneva remained uninfected during this wave of the pandemic. Assuming that the presence of IgG antibodies is associated with immunity, these results highlight that the epidemic is far from coming to an end by means of fewer susceptible people in the population.</p>
The Journal of clinical investigation 11JUN2020	<b>Early safety indicators of COVID-19 convalescent plasma in 5,000 patients</b>	Joyner, Michael J et al. USA <a href="#">gotopaper</a>	Therapeutic	<p>Analysis of key safety metrics after <b>transfusion of ABO compatible human COVID-19 convalescent plasma</b> in <b>5,000 hospitalized adults with severe or life threatening COVID-19</b>, with 66% in the intensive care unit, as part of the US FDA Expanded Access Program for COVID-19 convalescent plasma.</p> <p>- <b>Incidence of all serious adverse events (SAEs) in the first four hours after transfusion was &lt;1% (n=36)</b>, including mortality rate (0.3%).</p> <p>- Of the 36 reported SAEs, there were 25 reported incidences of related SAEs, including mortality (n = 4), transfusion-associated circulatory overload (TACO; n = 7), transfusion-related acute lung injury (TRALI; n = 11), and severe allergic transfusion reactions (n = 3).</p> <p>- Only 2 (of 36) SAEs were judged as definitely related to the convalescent plasma transfusion by the treating physician.</p> <p>- <b>The seven-day mortality rate was 14.9%.</b></p> <p>⇒ no signal of toxicity beyond what is expected from plasma use in severely ill patients</p> <p>⇒ mortality rate does not appear excessive</p> <p>These early indicators <b>suggest that transfusion of convalescent plasma is safe in hospitalized patients with COVID-19.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
PSYCHOLOGICAL MEDICINE 10JUN2020	Health-protective behaviour, social media usage, and conspiracy belief during the COVID-19 public health emergency	Allington D et al. UK <a href="#">gotopaper</a>	SHS/SciPo	<p>&gt; Social media platforms are major disseminators of health misinformation</p> <p>&gt; Spread of COVID-19 is subject of conspiracy theories on social media.</p> <p><b>Methods</b></p> <p>&gt; 3 questionnaire surveys of social media use, conspiracy beliefs, and health-protective behaviours with regard to COVID-19 (N = 949, 2250 N=2254; UK) <b>Results</b></p> <p>&gt; negative relationship between COVID-19 conspiracy beliefs and COVID-19 health-protective behaviours,</p> <p>&gt; positive relationship between COVID-19 conspiracy beliefs and use of social media as a source of information about COVID-19.</p> <p><b>Conclusions</b></p> <p>Unregulated social media may present a health risk partly reducible to their role as disseminators of health-related conspiracy beliefs.</p>
Nature 09JUN2020	Clinical benefit of remdesivir in rhesus macaques infected with SARS-CoV-2	Williamson BN et al USA <a href="#">gotopaper</a>	Therapeutic	<p>Rhesus macaque model of SARS-CoV-2 infection → develop mild to moderate disease</p> <p>2 group of 6 macaque were inoculated:</p> <ul style="list-style-type: none"> <li>- 12 hours after 1 received remdesivir IV and 1 received vehicle solution</li> <li>- Assigned a daily clinical score sheet in a blinded fashion, 12 hours after inoculation</li> </ul> <p><u>Animal treated with remdesivir → 12 hours after inoculation:</u></p> <ul style="list-style-type: none"> <li>- <b>Clinical scores were significantly lower than in control animals</b></li> <li>- <b>Reduced pulmonary infiltrate on radiographs</b></li> <li>- <b>Reduced virus titers in bronchoalveolar lavage</b></li> <li>- <b>Reduction in damage to the lung significantly at necropsy</b></li> </ul> <p>Absence of resistance mutation in all remdesivir-treated animals</p> <p>No reduction in virus shedding</p> <p>→ <b>The first antiviral treatment with proven efficacy against SARS-CoV-2 in an animal model</b></p> <p>→ Early remdesivir treatment initiation in COVID-19 patients prevent progression to pneumonia</p>
European journal of emergency medicine 08JUN2020	COVID-19 epidemic in the Seine-Saint-Denis Department of Greater Paris: one month and three waves for a tsunami	Lapostolle, F. et al. France <a href="#">gotopaper</a>	Public Health/Epidemio	<p>It is proposed that the chronology of epidemic spread gives a window of time in which hospitals can act to prevent reaching capacity. The number of patients managed, of patients transferred to emergency departments (ED), and of mobile intensive care units (MICUs) dispatched in the department of Seine-Saint-Denis between February 17<sup>th</sup> and March 28<sup>th</sup> was compared to a reference period of 5 years (2015-2019). The alert threshold to indicate a public health crisis was defined as a 20% increase compared to the 5-year mean.</p> <ul style="list-style-type: none"> <li>- The daily numbers of patients managed crossed the threshold on February 25<sup>th</sup> and increased until the end of the study period.</li> <li>- The daily number of patients transferred to ED crossed the threshold on March 16<sup>th</sup> and increased until the end of the period.</li> <li>- The daily number of MICUs dispatched crossed the threshold on March 15<sup>th</sup> and increased until the end of the period.</li> </ul> <p>The COVID-19 epidemic reached the department in three consecutive waves, and the first wave preceded by 30 days the massive arrival of critical patients. Health care systems must take advantage of this delay to prepare for the next wave.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Intensive Care Medicine 08JUN2020	<b>Hydroxychloroquine pharmacokinetic in COVID-19 critically ill patients: an observational cohort study</b>	Painvin B et al France <a href="#">gotopaper</a>	Therapeutic	<p>HCQ pharmacokinetic in COVID-19 critically ill patients Oral loading dose: 400 mg or 200 mg if AKI and after 200mg t.i.d or 200mg b.i.d if AKI Measure daily blood trough and peak concentration of HCQ → evaluation of PK parameters Target HCQ concentration: 1 to 2 mg/L</p> <p><b>8 patients:</b> median age was 59 (IQR:53 – 70) – 5 with AKI and 1 with ECLS – all under MV – SOFA score at admission was 5 [3 – 7] <b>Analysis of 212 HCQ levels</b> Median HCQ peak concentration after loading dose: 0,5mg/L or 0,22mg/L (AKI) Median time to obtain target concentration: 4 days [3 – 7] Median duration in therapeutic range: 3,3 days Toxic levels were noted after day 5 of treatment → drug accumulation and high volume of distribution</p> <p>→ prescribing HCQ in COVID-19 patients if unsafe → <b>monitoring of electrocardiogram and blood concentration daily</b></p>
Chest 08JUN2020	<b>Tocilizumab treatment for cytokine release syndrome in hospitalized COVID-19 patients: survival and clinical outcomes</b>	Price C et al USA <a href="#">gotopaper</a>	Therapeutic	<p>Observational study – Patient were treated with tocilizumab using an algorithm that targeted cytokine release syndrome (CRS) Algorithm developed by a multidisciplinary team <b>239 patients included</b>– age median: 64 years</p> <p><b>153 were treated with tocilizumab</b> Patient with severe disease were significantly more likely to:</p> <ul style="list-style-type: none"> <li>- Have higher admission hsCRP levels (120 vs 71 mg/L, <math>p&lt;0,001</math>)</li> <li>- Have Abnormal chest radiographs</li> <li>- Received tocilizumab (90% vs 44%, <math>p&lt;0,001</math>)</li> <li>- Received tocilizumab sooner (2 vs 3 days, <math>p&lt;0,01</math>)</li> </ul> <p>14-days survival was 84% for patients treated with tocilizumab:</p> <ul style="list-style-type: none"> <li>- Not differ by disease severity (83% vs 91%, <math>p=0,11</math>)</li> <li>-</li> </ul> <p>Mechanical ventilation (MV) occurred in 31% of patients treated with tocilizumab</p> <p><u>After tocilizumab</u></p> <ul style="list-style-type: none"> <li>- Improve of inflammatory biomarkers (hsCRP, IL-6) and oxygenation</li> <li>- D-dimer increased significantly</li> <li>- Few adverse events</li> </ul> <p>→ Tocilizumab to target CRS may influence MV and survival outcomes → <b>Need for randomized trials</b></p>
Intensive Care Medecine 8JUN2020	<b>Severe COVID-19 is associated with deep and sustained multifaceted cellular immunosuppression</b>	Jeannet et al., France <a href="#">gotopaper</a>	Immuno	<p>This French study conducted on 13 consecutively recruited patients infected with SARS-CoV-2 virus during their first week of ICU stay with 10 healthy donors used as controls confirms that patients uniformly exhibited deep global and persisting T, NK and B cell lymphopenia from ICU admission to day 7 Interestingly the report indicated few CD4 T cells transiently expressed CTLA-4 during the first 3 days and expression of PD-1 observed at admission day increased until day 7 whereas CD8 T cells significantly expressed PD-1 from admission day to day 7 and CTLA-4 expression remained unchanged.</p> <p>Conclusion: These data strongly suggest a vicious effect of the virus to cause an upregulation of potent T cell killing and immunosuppressive mechanisms in critically-ill COVID-19 patients. This call into question therapies (e.g., anti-IL-6, corticosteroids, JAK inhibitors) that aim to block the ability of the patient to mount an effective immune response to the invading SARS-CoV-2.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
PLOS ONE 08 JUN2020	Quantify the role of superspreaders -opinion leaders- on COVID-19 information propagation in the Chinese Sina-microblog	Yin F et al CHINA AND CANADA <a href="#">gotopaper</a>	SHS/SciPo	<p><b>Background:</b></p> <ul style="list-style-type: none"> <li>&gt; the success of control of COVID-19 outbreak relies on the resilience of people to follow public health interventions</li> <li>➤ evidence shows that opinion leaders paly a significant role in the propagation of epidemic information and public health policy and implementations</li> </ul> <p><b>Methods:</b></p> <ul style="list-style-type: none"> <li>&gt; “Opinion-leader susceptible-forwarding-immune (OL-SFI)” mathematical model to quantify the roles of information “superspreaders”</li> <li>➤ analysis of the information propagation dynamics in the Chinese Sina -microblog.</li> </ul> <p><b>Results:</b></p> <p>The earlier opinion leaders get into the pubic heath intervention, the greater their influence will be on the population.</p>
PLoS biology 8JUN2020	A unifying structural and functional model of the coronavirus replication organelle: Tracking down RNA synthesis	Snijder, Eric J. <i>et al.</i> Netherlands <a href="#">gotopaper</a>	Fundamental research	<p>Which replication organelle (RO) element(s) of infected cells accommodate CoV RNA synthesis remains unclear.</p> <ul style="list-style-type: none"> <li>- 2D and 3D analyses of CoV ROs showed that <b>diverse CoVs induce the same membrane modifications</b>, including the small open double-membrane spherules (DMSs) (previously thought to be restricted to gamma- and delta-CoV infections). But <b>RNA synthesis could not be linked to DMSs</b> or any other cellular or virus-induced structure.</li> <li>- <b>Abundant association of newly synthesized viral RNA with double-membrane vesicles (DMVs) detected</b> in cells infected with the beta-CoV MERS-CoV and SARS-CoV, and gamma-CoV infectious bronchitis virus (Metabolic labelling by quantitative EM autoradiography).</li> </ul> <p><b>-&gt; Provides a unifying model of the CoV replication organelle and establish DMVs as the central hub for viral RNA synthesis and a potential drug target in CoV infection.</b></p>
Lancet 5JUN2020	<b>Retraction:</b> Hydroxychloroquine or chloroquine with or without a macrolide for treatment of COVID-19: a multinational registry analysis	Mandeep R Mehra et al. USA <a href="#">gotopaper</a>	Therapeutic	<p>Unable to conduct an independent and private peer review to evaluate the origination of the database elements, to confirm the completeness of the database, and to replicate the analyses presented in the paper “Hydroxychloroquine or chloroquine with or without a macrolide for treatment of COVID-19: a multinational registry analysis”</p> <p>The raw data could not be made available to an independent third-party peer review</p> <p><b>Withdrawal of the article at the request of the authors</b></p>
NEJM 04JUN2020	<b>Retraction:</b> Cardiovascular Disease, Drug Therapy, and Mortality in Covid-19. N Engl J Med. DOI: 10.1056/NEJMoa2007621	Mehra MR et al USA <a href="#">gotopaper</a>	Clinic	<p><b>Unable to validate the primary data source underlying the article “Cardiovascular Disease, Drug Therapy, and Mortality in Covid-19”</b></p> <p>The raw data could not be made available to a third-party auditor</p> <p><b>Withdrawal of the article at the request of the authors</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 8JUN2020	<b>Estimating the effects of non-pharmaceutical interventions on COVID-19 in Europe</b>	Flaxman, Seth et al. UK <a href="#">gotopaper</a>	Therapeutic	<p>Study of the <b>impact of major non-pharmaceutical interventions</b> across 11 European countries for the period from the start of COVID-19 until the 4th of May 2020 when lockdowns started to be lifted. These large-scale non-pharmaceutical interventions vary between countries but include social distancing (such as banning large gatherings), border closures, school closures, measures to isolate symptomatic individuals and their contacts, and large-scale lockdowns of populations with all but essential internal travel banned.</p> <ul style="list-style-type: none"> <li>- Estimate that, for all the countries considered, <b>current interventions have been sufficient to drive the reproduction number <math>R_t</math> below 1</b> (probability <math>R_t &lt; 1.0</math> is 99.9%) and <b>achieve epidemic control</b>.</li> <li>- Estimate that, across all 11 countries, between 12 and 15 million individuals have been infected with SARS-CoV-2 up to 4th May, representing <b>between 3.2% and 4.0% of the population</b>.</li> </ul> <p>=&gt; Major non-pharmaceutical interventions and lockdown in particular have had a large effect on reducing transmission. Continued intervention should be considered to keep transmission of SARS-CoV-2 under control.</p>
Blood 08JUN2020	<b>COVID and Coagulation: Bleeding and Thrombotic Manifestations of SARS-CoV2 Infection</b>	Al-Samkari H et al USA <a href="#">gotopaper</a>	Clinic	<p>Multicenter study – rate and severity of hemostatic and thrombotic complications  <b>400 patients COVID-19</b> receiving standard dose prophylactic anticoagulation  <u>Overall population:</u> <ul style="list-style-type: none"> <li>- Radiographically-confirmed VTE rate: 4,8% [2,9 – 7,3] / Thrombotic complications rate: 9,5% [6,8 – 12,8] / Major bleeding rate: 2,3% [1,0 – 4,2]</li> </ul> <u>Critically ill patients:</u> <ul style="list-style-type: none"> <li>- Radiographically-confirmed VTE rate: 7,6% [3,9 – 13,3] / Major bleeding rate: 5,6% [2,4 – 10,7]</li> </ul> <u>Predictive of coagulation-associated complications during hospitalization</u> <ul style="list-style-type: none"> <li>- Elevated D-dimer at admission / Platelet count &gt; 450x10<sup>9</sup>/L / CRP &gt; 100mg/L / Erythrocyte sedimentation rate &gt;40mm/h</li> </ul> <p>→ <b>Randomized trials are needed to determine any potential benefit of intensified anticoagulation prophylaxis</b></p> </p>
JAMA 08JUN2020	<b>Clinical Characteristics of 58 Children with a Pediatric Inflammatory Multisystem Syndrome Temporally Associated With SARS-CoV-2</b>	Whittaker E et al UK <a href="#">gotopaper</a>	Clinic	<p><b>58 children</b> in 8 hospitals in England with PIMS-TS  → Comparison with patients with KD (n=1132), KD shock syndrome (n=45) and toxic shock syndrome (n=37) admitted in US and Europe from 2002 to 2019</p> <p>Median age: 9 years [5,7 – 14] – 57% female  <u>Symptoms</u> <ul style="list-style-type: none"> <li>- Fever with non-specific symptoms: vomiting – abdominal pain – diarrhea - .... / 52% rash &amp; 45% conjunctival injection / 22% children met definition of KD (AHA)</li> </ul> <u>Biology</u> <ul style="list-style-type: none"> <li>- PCR SARS-CoV-2 positive 15/58 &amp; SARS-CoV-2 IgG positive in 40 of 46</li> <li>- <b>78% had evidence of current or prior infection of SARS-CoV-2</b></li> <li>- High level of inflammatory markers (CRP, ferritin)</li> </ul> <u>Outcomes</u> <ul style="list-style-type: none"> <li>- 50% developed shock with evidence of left ventricular dysfunction whom 79% received mechanical ventilation / 14% developed coronary artery dilatation or aneurysm / <b>No death</b></li> </ul> <u>Comparison with KD and KD shock syndrome</u> <ul style="list-style-type: none"> <li>- Older (9 vs 2,7 vs 3,8 respectively)</li> <li>- Greater elevation of inflammatory markers</li> <li>- Lower platelet count – higher fibrinogen levels and greater elevation of troponin</li> </ul> <p>→ <b>the comparison suggests this disorder differ from other pediatric inflammatory entities</b></p> </p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Lancet Infectious Disease 08JUN2020	<b>Pulmonary post-mortem findings in a series of COVID-19 cases from northern Italy: a two-centre descriptive study</b>	Carsana L et al Italy <a href="#">gotopaper</a>	Clinic	<p>Analysis of lung tissue samples from 38 patients who died from COVID-19</p> <p><b>Diffuse alveolar damage:</b></p> <ul style="list-style-type: none"> <li>- Capillary congestion (100%)</li> <li>- Necrosis of pneumocytes (100%)</li> <li>- Type 2 pneumocytes hyperplasia (100%)</li> <li>- Necrosis of hyaline membranes (87%)</li> <li>- Interstitial &amp; intra-alveolar edema (97%)</li> <li>- Platelet-fibrin thrombi (87%)</li> </ul> <p>→ <b>presence of platelet–fibrin thrombi in small arterial vessels is consistent with coagulopathy</b></p> <p>→ <b>same lesions in patients infected with SARS and MERS-CoV</b></p>
Science 08JUN2020	<b>Genomic surveillance reveals multiple introductions of SARS-CoV-2 into Northern California</b>	Deng, X. et al, USA <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Investigation of the genomic epidemiology of SARS-CoV-2 in Northern California from late January to mid-March 2020, using samples from 36 patients spanning 9 counties and the Grand Princess cruise ship.</p> <ul style="list-style-type: none"> <li>- Phylogenetic analyses revealed the cryptic introduction of at least 7 SARS-CoV-2 lineages into California, including epidemic WA1 strains associated with Washington State, with lack of a predominant lineage and limited transmission between communities.</li> <li>- Lineages associated with outbreak clusters in 2 counties were defined by a single base substitution in the viral genome.</li> </ul> <p>These findings support contact tracing, social distancing, and travel restrictions to contain SARS-CoV-2 spread.</p>
Nature 08JUN2020	<b>The effect of large-scale anti-contagion policies on the COVID-19 pandemic</b>	Hsiang S. et al, USA <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Reduced-form econometric methods are applied to empirically evaluate the effect of policies on the growth rate of infections, using new data on 1,717 local, regional, and national non-pharmaceutical interventions deployed in the Covid-19 pandemic across China, South Korea, Italy, Iran, France, and the US.</p> <ul style="list-style-type: none"> <li>- In the absence of policy actions, early infections exponential growth rates are estimated at roughly 38% per day.</li> <li>- Anti-contagion policies have significantly and substantially slowed this growth: interventions prevented or delayed on the order of 62 million confirmed cases, corresponding to averting roughly 530 million total infections.</li> </ul> <p>These findings may help inform whether or when these policies should be deployed, intensified, or lifted</p>
Journal of the American College of Cardiology 08JUN2020	<b>Prevalence and Impact of Myocardial Injury in Patients Hospitalized with COVID-19 Infection</b>	Lala A. et al, USA <a href="#">gotopaper</a>	Public Health/Epidemio	<p>AIM: Describing the degree of myocardial injury and associated outcomes in a hospitalized cohort with COVID-19 (n=2,736), based on troponin-I measures (normal value &lt;0.03ng/mL).</p> <ul style="list-style-type: none"> <li>- Cardiovascular disease (CVD; coronary artery disease, atrial fibrillation, and heart failure), hypertension and diabetes were more prevalent in patients with higher troponin concentrations.</li> <li>- 506 (18.5%) patients died during hospitalization. In all, 985 (36%) patients had elevated troponin concentrations.</li> <li>- After adjusting for disease severity and relevant clinical factors, even small amounts of myocardial injury (troponin I 0.03-0.09ng/mL) were significantly associated with death, while greater amounts (troponin I &gt;0.09 ng/dL) were significantly associated with higher risk.</li> </ul> <p>Myocardial injury is prevalent among patients hospitalized with COVID-19 however troponin concentrations were generally low.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
J Med Virol 5JUN2020	Identification of nsp1 gene as the target of SARS-CoV-2 real-time RT-PCR using nanopore whole genome sequencing	Chan, Wan-Mui et al. Hong Kong <a href="#">gotopaper</a>	Virology	<p>There is an <b>increasing number of SARS-CoV-2 viruses with mutations</b> at the primer or probe binding sites. <b>These mutations may affect the sensitivity</b> of RT-PCR) assays targeting the N, E, and ORF1a/b genes.</p> <p>=&gt; Using sequence-independent single-primer amplification (SISPA) and nanopore whole-genome sequencing, we have found that the nsp1 gene, located at the 5' end of the SARS-CoV-2 genome, was highly expressed in the nasopharyngeal or saliva specimens of 9 COVID-19 patients of different clinical severity.</p> <p>=&gt; we have developed <b>a novel nsp1 real-time RT-PCR assay</b>. The primers and probes are highly specific for SARS-CoV-2.</p> <p><b>Results:</b> Validation with 101 clinical specimens showed that our nsp1 RT-PCR assay has a sensitivity of 93.1% (95% confidence interval, 86.2-97.2%), which was similar to those of N and E gene RT-PCR assays. The diagnostic specificity was 100% (95% CI, 92.9-100%)</p> <p><b>Conclusion:</b> The <b>addition of nsp1 for multi-target detection of SARS-CoV-2 can avoid false negative results due to mutations</b> at the primers/probes binding sites of currently available RT-PCR assays.</p> <p><b>Limitations:</b> There can still be bias during the library preparation which may affect the coverage of each position in the viral genome + our nanopore sequencing did not encompass phylogenetic type C strains, which is mainly found in Europe and in America.</p>
Med 5JUN2020	Outcomes of hydroxychloroquine usage in United States veterans hospitalized with COVID-19	Magagnoli, Joseph et al. USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Retrospective study</b> of electronic health records of <b>807 US veterans patients hospitalized</b> with confirmed SARS-CoV-2 infection to analyze the associations between hydroxychloroquine and azithromycin use and clinical outcome.</p> <p>The primary outcomes were <b>mortality and use of mechanical ventilation</b>.</p> <p>- Compared to the no HC group, after propensity score adjustment for clinical characteristics, the <b>risk of death from any cause was higher in the HC group</b> (adjusted hazard ratio (aHR), 1.83; 95% CI, 1.16 to 2.89; P=0.009) but <b>not in the HC+AZ group</b> (aHR, 1.31; 95% CI, 0.80 to 2.15; P=0.28).</p> <p>- Both the <b>propensity score-adjusted risks of mechanical ventilation and death after mechanical ventilation were not significantly different</b> in the HC group (aHR, 1.19; 95% CI, 0.78 to 1.82; P=0.42 and aHR, 2.11; 95% CI, 0.96 to 4.62; P=0.06, respectively) or in the HC+AZ group (aHR, 1.09; 95% CI, 0.72 to 1.66; P=0.69 and aHR, 1.25; 95% CI, 0.59 to 2.68; P=0.56, respectively), compared to the no HC group.</p> <p><b>Limitations:</b> non-randomization of treatments; demographic composition of patients (US veterans) with a majority of men and majority of black.</p>
HEART FAILURE JUN2020	Incidence of New-Onset and Worsening Heart Failure Before and After the COVID-19 Epidemic Lockdown in Denmark	Andersson C et al, DENMARK <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Consequences of the lockdown for patients with heart failure (Cohort Study in Denmark)</p> <p>&gt; Incidence of new onset HF and hospitalizations for worsening HF before and after the lockdown (2019 vs 2020, same period)</p> <p><b>Results:</b></p> <p><b>1. Before lockdown:</b></p> <p>&gt; Rates of new-onset HF before lockdown were comparable for 2020 and 2019 / Hospitalizations for worsening HF were slightly higher in 2020 versus 2019 in</p> <p><b>2. During lockdown:</b></p> <p>&gt; Rates of new-onset HF diagnoses and of hospitalizations for worsening HF significantly lower in 2020 versus 2019 / Similar mortality before and after lockdown for the population with HF.</p> <p><b>Conclusions:</b></p> <p>&gt; These data raise concerns for a potential undertreatment of HF currently that may impact prognosis in the longer term</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JOURNAL OF MEDICAL INTERNET RESEARCH 05JUN2020	<b>COVID-19-Related Information Sources and the Relationship With Confidence in People Coping with COVID-19: Facebook Survey Study in Taiwan</b>	Wang PW et al. TAIWAN <a href="#">gotopaper</a>	SHS/SciPo	<p><b>Objectives:</b> to examine major COVID-19 information sources of people in coping with COVID-19 in Taiwan.</p> <p><b>Methods:</b></p> <ul style="list-style-type: none"> <li>➤ 1904 participants (1270 non–health-care workers (HCW) and 634 HCW) recruited from Facebook advertisement.</li> <li>➤ surveyed sources of information: internet (blogs, internet news, and social media: Facebook, Line, Twitter, and Plurk), friends, traditional media (television, newspapers, and radio broadcasting) and formal lesson on COVID19</li> <li>➤ Participants rated their frequency for each source, and responded to question on self-confidence and worries in coping with COVID19</li> </ul> <p><b>Results:</b></p> <ul style="list-style-type: none"> <li>➤ Sources of information by order of acces: internet (80%) followed by followed by traditional media, family members, coworkers, friends, formal lessons, and medical staff. &gt; 50% of participant consulted one or two sources. 10% zero.</li> <li>➤ for HCW: the use of formal lessons as an information source was associated with better self-confidence in coping with COVID-19</li> <li>➤ association between receiving information from more sources and greater self-confidence found in HCW, but not in other groups</li> </ul> <p><b>Conclusions:</b></p> <ul style="list-style-type: none"> <li>&gt; internet is a popular and accessible information source, but misinformation on COVID-19 is rife</li> <li>➤ severe worry is associated with using more information sources.</li> </ul> <p>Medical professionals should consider that when delivering information online.</p>
Sci. Total Environ. 5JUN2020	<b>Comparison of virus concentration methods for the RT-qPCR-based recovery of murine hepatitis virus, a surrogate for SARS-CoV-2 from untreated wastewater</b>	Ahmed, Warish et al. Australia <a href="#">gotopaper</a>	Public Health	<p>There is currently a clear benefit for many <b>countries to utilize wastewater-based epidemiology (WBE)</b> as part of ongoing measures to manage the COVID-19 global pandemic. It is <b>imperative to determine the efficiency of the most commonly used methods</b> for the enveloped SARS-CoV-2.</p> <p>⇒ Municipal wastewater seeded with a human coronavirus (CoV) surrogate, murine hepatitis virus (MHV), was used <b>to test the efficiency of seven wastewater virus concentration methods:</b> (A–C) adsorption-extraction with three different pre-treatment options, (D–E) centrifugal concentration device methods with two different devices, (F) polyethylene glycol (PEG 8000) concentration, and (G) ultrafiltration. MHV was quantified by reverse-transcription quantitative polymerase chain reaction and the concentration efficiency was calculated for each method.</p> <p><b>Results:</b> the most efficient methods were (B) Adsorption-extraction methods, with MgCl<sub>2</sub> pre-treatment.</p> <p><b>CI<sup>®</sup>:</b> absorption-extraction methods with minimal pretreatment or without manipulation can provide suitably rapid, cost-effective and relatively straightforward recovery of enveloped viruses in wastewater. <b>The MHV is a promising process control for SARS-CoV-2 surveillance and can be used as a quality control measure to support community-level epidemic mitigation and risk assessment.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
CELL 4JUN2020	Development of an inactivated vaccine candidate, BBIBP-CorV, with potent protection against SARS-CoV-2	Wuang H et al_China <a href="#">gotopaper</a>	Vaccin	<p><b>BBIBP-CorV inactivated SARS-CoV-2 vaccine candidate:</b> 3 viral strains isolated from BAL or throat swabs from 3 hospitalized patients in Vero cells. &gt; The strain showing the most optimal replication rates and highest virus yield was selected.</p> <p><b>Immunogenicity tests:</b>  <b>In mice:</b> BalbC injected with 2, 4, 8 ug/dose of BBIBP-CorV +aluminium hydroxide adjuvant.            &gt; One/Two/three dose programmes: High seroconversion rate in the three groups at D7. Better NAb production in three dose program as evaluated at D28  <b>In rabbits, guinea pigs, rats and NHP:</b> good immunogenicity and seroconversion rates in a two-dose vaccination program.  <b>Protection in NHP</b>            Immunization of NHP with BBIBP-CoV-2 after intratracheal challenge: 2 doses ; 2ug/dose            &gt; highly efficient protection against SARS-CoV-2            &gt; viral clearance in lungs, throat and and swabs            &gt; no side effects on serum biochemical parameters            &gt;no ADE</p> <p>Safety evaluation in rat showed no specific concerns            BBIBP-CorV exhibits efficient productivity and good genetic stability for vaccine manufacture</p>
Allergy 4JUN2020	Distribution of ACE2, CD147, CD26 and other SARS-CoV-2 associated molecules in tissues and immune cells in health and in asthma, COPD, obesity, hypertension, and COVID-19 risk factors	Radzikowska, U. <i>et al.</i> Switzerland-Poland-China-Norway-Japan-Germany-USA-Ireland <a href="#">gotopaper</a>	Fundamental research	<p>Gene expression analysis (RNA sequencing and RNA-Seq databases) of SARS-CoV-2 receptors and related molecules in collection of primary human cells and tissues from healthy (children and adults) and adult patients with risk factors and known comorbidities of COVID-19: (ACE2, CD147 (<i>BSG</i>), CD26 (<i>DPP4</i>), and their direct and indirect molecular partners in primary human bronchial epithelial cells, bronchial and skin biopsies, bronchoalveolar lavage fluid, whole blood, PBMCs, monocytes, neutrophils, DCs, NK cells, ILC1, ILC2, ILC3, CD4+ and CD8+ T cells, B cells and plasmablasts)            - ACE2 and TMPRSS2 co-expressed at epithelial sites of lung and skin,            - CD147, cyclophilins (<i>PPIA</i> and <i>PPIB</i>), CD26 and related molecules expressed in both epithelium and in immune cells.            - <b>Distinct age-related expression profile of these genes in the PBMCs and T cells from healthy children and adults.</b>            - <b>Higher expression of ACE2- and CD147-related genes generally in asthma, COPD, hypertension, smoking, obesity, and male gender</b> in the bronchial biopsy, BAL or blood.            - <b>CD147-related genes correlated positively with age and BMI.</b>            -&gt; Different receptor repertoire potentially involved in the SARS-CoV-2 infection at the epithelial barriers and in the immune cells. <b>Altered expression of these receptors related with age, gender, obesity and smoking, as well as with the disease status might contribute to COVID-19 morbidity and severity patterns.</b></p>
The Lancet Rheumatology 4JUN2020	Canakinumab in a subgroup of patients with COVID-19	Ucciferri, Claudio et al. Italie/USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Retrospective analysis of 10 hospitalized patients</b> with confirmed SARS-CoV-2 infection, bilateral pneumonia, hyperinflammation (defined as serum C-reactive protein <math>\geq 50</math> mg/L), and respiratory failure (requiring supplemental oxygen without invasive ventilation). These patients were treated with <b>canakinumab</b>, a human <b>monoclonal antibody against IL-1<math>\beta</math></b>. All patients also received hydroxychloroquine and lopinavir–ritonavir.            Canakinumab was safe, <b>well tolerated</b>, and associated with a <b>rapid reduction in the systemic inflammatory response</b> and an <b>improvement in oxygenation</b> compared to controls. Notably, none of the patients developed neutropenia or bacterial sepsis.  <b>Limitations:</b> small sample size and absence of a random comparison.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 3JUN2020	Effect of Convalescent Plasma Therapy on Time to Clinical Improvement in Patients With Severe and Life-threatening COVID-19: A Randomized Clinical Trial	Li, Ling et al. China <a href="#">gotopaper</a>	Therapeutic	<p><b>Open-label, multicenter, randomized clinical trial</b> including <b>103 participants</b> with laboratory-confirmed COVID-19 <b>severe</b> (respiratory distress and/or hypoxemia) or <b>life-threatening</b> (shock, organ failure, or requiring mechanical ventilation) treated with <b>Convalescent plasma</b> in addition to standard treatment vs standard treatment alone (stratified by disease severity).</p> <p>The trial was <b>terminated early</b> after 103 of a planned 200 patients were enrolled.</p> <p>Primary outcome was <b>time to clinical improvement</b> within 28 days. Secondary outcomes included 28-day mortality, time to discharge, and the rate of viral PCR results turned from positive at baseline to negative at up to 72 hours.</p> <ul style="list-style-type: none"> <li>- Clinical improvement occurred within 28 days in 51.9% of the convalescent plasma group vs 43.1% in the control group (difference, 8.8% [95% CI, -10.4% to 28.0%]; hazard ratio [HR], 1.40 [95% CI, 0.79-2.49]; P = .26).</li> <li>Among those with severe disease, the primary outcome occurred in 91.3% (21/23) of the convalescent plasma group vs 68.2% (15/22) of the control group (HR, 2.15 [95% CI, 1.07-4.32]; P = .03); among those with life-threatening disease the primary outcome occurred in 20.7% (6/29) of the convalescent plasma group vs 24.1% (7/29) of the control group (HR, 0.88 [95% CI, 0.30-2.63]; P = .83) (P for interaction = .17).</li> <li>- There was no significant difference in 28-day mortality (15.7% vs 24.0%; OR, 0.65 [95% CI, 0.29-1.46]; P = .30) or time from randomization to discharge (51.0% vs 36.0% discharged by day 28; HR, 1.61 [95% CI, 0.88-2.93]; P = .12).</li> <li>- Convalescent plasma treatment was associated with a negative conversion rate of viral PCR at 72 hours in 87.2% of the convalescent plasma group vs 37.5% of the control group (OR, 11.39 [95% CI, 3.91-33.18]; P &lt; .001).</li> <li>- Two patients in the convalescent plasma group experienced adverse events within hours after transfusion that improved with supportive care.</li> </ul> <p><b>=&gt; Among patients with severe or life-threatening COVID-19, convalescent plasma therapy added to standard treatment, compared with standard treatment alone, did not result in a statistically significant improvement in time to clinical improvement within 28 days.</b></p> <p><b>Interpretation is limited by early termination</b> of the trial, which may have been underpowered to detect a clinically important difference.</p>
Biosensors and Bioelectronics JUN2020	Homogeneous circle-to-circle amplification for real-time optomagnetic detection of SARS-CoV-2 RdRp coding sequence	Tian, Bo et al. Denmark <a href="#">gotopaper</a>	Diagnostic	<p><b>Circle-to-circle amplification (C2CA)</b> is a specific and precise cascade nucleic acid amplification method consisting of more than one round of padlock probe ligation and rolling circle amplification (RCA). We herein demonstrate a <b>homogeneous and isothermal nucleic acid quantification strategy based on C2CA and optomagnetic analysis of magnetic nanoparticle (MNP) assembly</b>. -&gt; eliminates the need for additional monomerization and ligation steps after the first round of RCA, and combines two amplification rounds in a one-pot reaction.</p> <p>-&gt; Applied for the <b>detection of a synthetic complementary DNA of SARS-CoV-2 RdRp</b> (RNA-dependent RNA polymerase) coding sequence, achieving a detection limit of 0.4 fM with a dynamic detection range of 3 orders of magnitude and a total assay time of <i>ca.</i> 100 min. A mathematical model was set up and validated to predict the assay performance.</p> <p><b>Conclusion:</b> Reaches a sub-femtomolar level detection limit and significantly simplified the operation by eliminating the labor-intensive and time-consuming operation steps requiring different reaction temperatures. <b>Capability of target quantification in 10% FBS samples was demonstrated</b> with an acceptable loss of sensitivity.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
J. Infect. 3JUN2020	<b>Detection of SARS-CoV-2 antibodies using commercial assays and seroconversion patterns in hospitalized patients</b>	Tuaillon, E. et al. France <a href="#">gotopaper</a>	Diagnostic	<p>SARS-CoV-2 antibody assays are needed for serological surveys and as a complement to molecular tests to confirm COVID-19. However, the <b>kinetics of the humoral response against SARS-CoV-2 remains poorly described and relies on the performance of the different serological tests.</b> =&gt; Evaluation of performance of six CE-marked point-of-care tests (POC) and three ELISA assays for the diagnosis of COVID-19 by <b>exploring seroconversions in hospitalized patients</b> who tested positive for SARS-CoV-2 RNA.</p> <p><b>Results:</b> our study is one of the very first to evaluate the performance of commercial SARS-CoV-2 serologic assays. The second week of COVID-19 seems to be the best period for assessing the sensitivity of commercial serological assays.</p> <p><b>CI*:</b> <b>serological assays may be useful in the diagnosis of patients with acute respiratory distress syndrome and a negative PCR assay</b> <b>Seroconversions occur during the second week of the disease.</b> To achieve an early diagnosis of COVID-19 based on antibody detection, a dual challenge must be met: <b>the immunodiagnostic window period must be shortened and an optimal specificity must be conserved</b></p> <p><b>Limitations:</b> relatively small number of plasma samples -&gt; the estimation of sensitivity and specificity values are relatively imprecise. Patients with moderate to severe COVID-19 -&gt; intensity of the humoral response to SARS-CoV-2 N or S proteins may be lower in asymptomatic or paucity symptomatic cases.</p>
NEJM 03JUN2020	<b>A Randomized Trial of Hydroxychloroquine as Postexposure Prophylaxis for Covid-19</b>	David R. Boulware et al. USA/Canada <a href="#">gotopaper</a>	Therapeutic	<p><b>Randomized, double-blind, placebo-controlled trial</b> to evaluate <b>postexposure prophylaxis with hydroxychloroquine</b> after exposure to Covid-19. Participants had known exposure (by participant report) to a person with laboratory-confirmed Covid-19, whether as a <b>household contact</b>, a <b>health care worker</b>, or a person with other <b>occupational exposures</b>. The primary outcome was the incidence of either laboratory-confirmed Covid-19 or illness compatible with Covid-19 within 14 days.</p> <p><b>821 asymptomatic adult</b> participants were randomly assigned to the hydroxychloroquine group (414 participants) or the placebo group (407 participants). Overall, 87.6% of the participants (719 of 821) had high-risk exposures without eye shields and surgical masks or respirators.</p> <p>- The <b>incidence of new illness</b> compatible with Covid-19 <b>did not differ significantly</b> between participants receiving hydroxychloroquine (49 of 414 [11.8%]) and those receiving placebo (58 of 407 [14.3%]); the absolute difference was -2.4 percentage points (95% confidence interval, -7.0 to 2.2; P=0.35).</p> <p>- Side effects were more common with hydroxychloroquine than with placebo (40.1% vs. 16.8%), but no serious adverse reactions were reported.</p> <p><b>Conclusion:</b> <b>high doses of hydroxychloroquine did not prevent illness compatible with Covid-19 when initiated within 4 days after a high-risk or moderate-risk exposure.</b></p> <p><b>Limitation:</b> a priori symptomatic case definition used because majority of the participants, including health care workers, were unable to access testing ; Internet-based approach used to rapidly recruit participants in the context of a pandemic, data were obtained by means of participant report ; predictive power of the case definition is unknown, particularly in the younger populations studied.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet Public Health 02JUN2020	<b>Effects of non-pharmaceutical interventions on COVID-19 cases, deaths, and demand for hospital services in the UK: a modelling study</b>	Davies, N.G., et al. UK <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Assessment of the potential impact of control measures for COVID-19 in the UK, using a stochastic age-structured transmission model tracking 66.4 million people. Interventions modelled were school closures, physical distancing, shielding of people aged &gt;70, and self-isolation of symptomatic cases, a combination of the four, as well as lockdown. Estimations of new cases, patients requiring ICU admission, death and R0 were calculated.</p> <ul style="list-style-type: none"> <li>- The median unmitigated burden was of 23 million clinical cases and 350 000 deaths in the UK by December 2021.</li> <li>- The four base interventions were each likely to decrease R0, but not sufficiently to prevent ICU demand from exceeding health service capacity. Only lockdown periods were sufficient to bring R0 near or below 1.</li> <li>- The most stringent lockdown scenario resulted in a projected 120 000 cases and 50 000 deaths.</li> </ul> <p><b>Intensive interventions with lockdown periods would need to be in place for a large proportion of the coming year to prevent health-care demand exceeding availability</b></p>
European journal of heart failure 2JUN2020	<b>ACE-inhibitors and Angiotensin-2 Receptor Blockers are not associated with severe SARS-COVID19 infection in a multi-site UK acute Hospital Trust</b>	Bean, Daniel M. et al. UK <a href="#">gotopaper</a>	Clinic	<p>Hypothesis: ACE-inhibitors (ACEi) and Angiotensin-2 Blockers (ARB), commonly used in patients with hypertension or diabetes and may raise tissue ACE2 levels, could increase the risk of severe COVID19 infection.</p> <p>Consecutive cohort of 1200 acute COVID19 inpatients (2 hospitals) with multi-ethnic catchment population in London:</p> <ul style="list-style-type: none"> <li>- Mean age: 68 ± 17 years (57% male); 74% of patients with at least 1 comorbidity.</li> <li>- 34.6% reached the primary endpoint of death or transfer to a critical care unit for organ support within 21-days of symptom onset.</li> <li>- 33.3% were taking ACEi or ARB (significantly older and more comorbidities). - <b>→ Odds ratio for the primary endpoint in patients on ACEi and ARB, after adjustment for age, sex and co-morbidities, was 0.63 (CI 0.47-0.84, p &lt; 0.01).</b></li> <li>- <b>→ No evidence for increased severity of COVID19 disease in hospitalised patients on chronic treatment with ACEi or ARB.</b></li> </ul> <p>A trend towards a beneficial effect of ACEi/ARB requires further evaluation in larger meta-analyses and randomised clinical trials.</p>
Cell Systems 02JUN2020	<b>Ultra-high-throughput clinical proteomics reveals classifiers of COVID-19 infection</b>	Messner et al., UK, Germany, Sweden <a href="#">gotopaper</a>	Diagnostic	<ul style="list-style-type: none"> <li>- A standardized, ultra-highthroughput clinical platform for serum and plasma proteomics</li> <li>- Platform enables high precision quantification of 180 patient samples/day at low cost</li> <li>- 27 biomarkers are differentially expressed between WHO severity grades for COVID-19</li> <li>- Biomarkers include proteins not previously associated with COVID-19 infection</li> </ul>
The Lancet Psy 2JUN2020	<b>Global mental health and COVID-19</b>	Lola Kola, Nigeria <a href="#">gotopaper</a>	Psy	<p>Disruption of mental health, in particular in LMICs.</p> <p><b>Two successful global mental health strategies:</b></p> <ol style="list-style-type: none"> <li>1- <b>Task shifting</b>—the use of trained lay health workers to deliver health care in non-specialist settings (can be used to address the urgent need to build a provider base in developing countries)</li> <li>2- <b>Use of digital health technology</b> to strengthen health systems (use of mobile phone for health interventions in LMICs) -&gt; to increase access and coverage in hard-to-reach areas calls for more research on their effectiveness in LMICs -</li> </ol> <p>-&gt; the <i>Lancet</i> Commission on global mental health recommended adoption of digital interventions alongside traditional treatments/ -&gt; Efforts to respond to mental health needs present researchers with an important opportunity to build on what we know and advance progress in achieving the mental health objectives of universal health coverage.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Lancet 01JUN2020	<b>Physical distancing, face masks, and eye protection to prevent person-to-person transmission of SARS-CoV-2 and COVID-19: a systematic review and meta-analysis</b>	Chu et al, Canada <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Aim: to investigate the effects of physical distance, face masks, and eye protection on virus transmission in health-care and community settings, through a systematic review and meta-analysis of 172 observational studies (25 697 patients in total).</p> <ul style="list-style-type: none"> <li>- Transmission of viruses was lower with physical distancing of 1 m or more, compared with a distance of less than 1. protection was increased as distance was lengthened.</li> <li>- Face mask use resulted in a large reduction in risk of infection, with stronger associations with N95 or similar respirators compared with disposable surgical masks or similar.</li> <li>- Eye protection was associated with less infection</li> </ul> <p>These findings support physical distancing of 1 m or more and provide quantitative estimates for models and contact tracing to inform policy.</p>
Eur. Respir. J. 01JUN2020	<b>Estimates of the ongoing need for social distancing and control measures post-"lockdown" from trajectories of COVID-19 cases and mortality</b>	Loneragan, M., Chalmers, J.D. UK <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Predictions on trajectories, doubling times and R0 of Covid-19 under social distancing and lockdown measures, based on new cases and mortality data in 89 countries up to May 21<sup>st</sup> 2020</p> <ul style="list-style-type: none"> <li>- Estimates of R0 before lockdown was between 2.0 and 3.7 (USA, Italy, Spain, France and UK). There was little evidence that the restrictions had reduced R far below 1 in many places (France showed the most rapid reductions - R0 0.76, based on cases and 0.77 based on mortality).</li> <li>- With an intermittent lockdown strategy, few countries could have even 1 week per month unrestricted without resurgence of the epidemic.</li> <li>- Restoring 20% of the activity that has been prevented by the lockdowns does not reconcile with preventing the resurgence of the disease in most countries.</li> </ul>
Nature Communications 01JUN2020	<b>Two linear epitopes on the SARS-CoV-2 spike protein that elicit neutralizing antibodies in COVID-19 patients</b>	Poh, Chek Meng et al; Singapore <a href="#">gotopaper</a>	Immuno	<p><b>Identification of immunogenic targets against the coronavirus spike glycoprotein</b></p> <ul style="list-style-type: none"> <li>• In this study, using pools of overlapping linear B-cell peptides, two immunodominant linear B-cell epitopes on the S glycoprotein of SARS-CoV-2 were identified. SP14P5 is located in close proximity to the receptor binding domain whereas S21P2 is in the region that encompasses the fusion peptide, which is highly conserved among coronaviruses suggesting a potential pan-SARS epitope at this location.</li> <li>• Detection for both S14P5 and S21P2 was consistently and significantly higher in COVID-19 patients</li> <li>• Interestingly, antibody depletion assays demonstrate that antibodies recognized these two linear epitopes and can neutralise SARS-CoV-2.</li> </ul> <p>Conclusion: SP14P5 et S21P2 epitopes can potentially be used in the design of more sensitive serological assays for epidemiological or vaccine efficiency assessments since that antibodies targeting these two linear epitopes account for a significant fraction of the anti-S-neutralising response.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Sensors 31MAY2020	<b>Development of a Portable, Ultra-Rapid and Ultra-Sensitive Cell-Based Biosensor for the Direct Detection of the SARS-CoV-2 S1 Spike Protein Antigen</b>	Mavrikou, Sophie et al. Greece <a href="#">gotopaper</a>	Diagnostic	<p>One of the key challenges of the recent COVID-19 pandemic is the ability <b>to accurately estimate the number of infected individuals</b>, particularly asymptomatic and/or early-stage patients. =&gt; <b>Proof-of-concept development of a biosensor able to detect the SARS-CoV-2 S1 spike protein</b>. The biosensor is based on <b>membrane-engineered mammalian cells bearing the human chimeric spike S1 antibody</b>.</p> <p><b>Results:</b> We demonstrate that the attachment of the protein to the membrane-bound antibodies resulted in a selective and considerable change in the cellular bioelectric properties measured by means of a Bioelectric Recognition Assay. <b>The novel biosensor provided results in an ultra-rapid manner (3 min), with a detection limit of 1 fg/mL and a semi-linear range of response between 10 fg and 1 µg/mL</b>. In addition, the observed high sensitivity of the biosensor could allow for screening the virus in easy-to-obtain patient samples such as saliva. <b>No cross-reactivity</b> was observed against the SARS-CoV-2 nucleocapsid protein. Furthermore, the biosensor was configured as a <b>ready-to-use platform</b>, including a portable read-out device operated via smartphone/tablet</p> <p><b>Conclusion:</b> The novel biosensor can be potentially applied for the mass screening without prior sample processing. The next step will be the actual clinical validation of the assay using patient samples and comparison to current serological and molecular tests</p>
Int. J. Infect. Dis. 31MAY2020	<b>Fast SARS-CoV-2 detection by RT-qPCR in preheated nasopharyngeal swab samples</b>	Alcoba-Florez, Julia et al. Spain <a href="#">gotopaper</a>	Diagnostic	<p>Performance of <b>three alternative, simple and affordable protocols to rapidly detect SARS-CoV-2</b>, bypassing the long and tedious RNA extraction step and reducing the time to viral detection: three methods based on direct nasopharyngeal swab viral transmission medium (VTM) heating before the RT-qPCR: a) direct without additives; b) in a formamide-EDTA (FAE) buffer, c) in a RNAsnap™ buffer.</p> <p>Although with a delay in cycle threshold compared to the gold-standard, we found consistent results in nasopharyngeal swab samples that were subject to a direct 70°C incubation for 10 min.</p> <p><b>CI°:</b> This study provides valuable options <b>to overcome any supply chain issue</b> and help <b>to increase the throughput of diagnostic tests</b>, thereby complementing standard diagnosis.</p>
Int. J. Infect. Dis. 31MAY2020	<b>Aberrant hyperactivation of cytotoxic T-cell as a potential determinant of COVID-19 severity</b>	Kang, Chang Kyung et al. Rep. of Korea <a href="#">gotopaper</a>	Immuno	<p>We hypothesized that <b>immune response may contribute to progression of coronavirus disease-19 (COVID-19)</b> at the second week of illness. Therefore, <b>we compared cell-mediated immune (CMI) responses between severe and mild COVID-19 cases</b>.</p> <p>We examined peripheral blood mononuclear cells of laboratory-confirmed COVID-19 patients from their first and third weeks of illness.</p> <p><b>CI°:</b> <b>Severe COVID-19 had higher degree of proliferation, activation, and cytotoxicity of T-cells</b> at the late phase of illness without cytotoxic T-cell contraction, which might contribute to the development of severe COVID-19.</p> <p><b>Limitations:</b> Small numbers of patients. Requires more time points to be examined to elucidate the exact temporal changes of such responses or when the persistent cytotoxic T-cell activity returns to normal in severe cases. The potential immunomodulatory effects of lopinavir/ritonavir could not be adjusted because the drug was prescribed to all severe patients in this study.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
J. Clin. Virol. 30MAY2020	<b>Persistent detection of SARS-CoV-2 RNA in patients and healthcare workers with COVID-19</b>	Gombar, Saurabh et al. USA <a href="#">gotopaper</a>	Diagnostic	<p>Current <b>guidelines for returning health care workers (HCW) to service after a positive SARS-CoV-2 RT-PCR test</b> and ceasing of transmission precautions for patients is based on two general strategies: <b>A test-based strategy</b> that requires negative respiratory RT-PCR tests obtained after the resolution of symptoms; <b>a symptom-based strategy</b> that recommends excluding HCW from the workforce until a fixed period of time has elapsed from symptom recovery.</p> <p><b>Objective:</b> to better understand the appropriate length of symptom-based return to work and contact precaution strategies.</p> <p>Observational analysis of 150 patients and HCW shows that the <b>average time to transition from RT-PCR positive to negative was 24 days</b> after symptom onset and 10 % remained positive even 33 days after symptom onset.</p> <p><b>CL<sup>o</sup>:</b> the fixed length of time before returning to work or ceasing contract precautions be revised <b>to over one-month</b>. Note: our analysis could be overestimating the length of infectious spreading by detecting non-infectious viral shedding. Large trials that rely on methods that detect the infective virus (ie viral culture) have not yet been reported in the literature.</p>
Brain, Behavior and Immunity 30MAY2020	<b>COVID-19 pandemic and mental health consequences: systematic review of the current evidence</b>	Vindegard and Eriksen Benros, Denmark <a href="#">gotopaper</a>	Psy	<p>COVID-19 patients displayed high levels of PTSS and increased levels of depression.</p> <ul style="list-style-type: none"> <li>•Patients with preexisting psychiatric disorders reported worsening of psychiatric symptoms.</li> <li>•Higher levels of psychiatric symptoms were found among health care workers.</li> <li>•A decrease in psychological well-being was observed in the general public.</li> <li>•However, well conducted large-scale studies are highly needed.</li> </ul>
Clin. Infect. Dis. 30MAY2020	<b>Maximum Daily Temperature, Precipitation, Ultra-Violet Light and Rates of Transmission of SARS-Cov-2 in the United States</b>	Shera T et al., USA <a href="#">gotopaper</a>	Public Health/Epidemio	<p>Aim: to investigate effects of temperature, precipitation, and UV Light on community transmission of SARS-CoV-2 in the USA.</p> <ul style="list-style-type: none"> <li>- A maximum temperature &gt;52° F on a given day was associated with a lower rate of new cases at 5 days.</li> <li>- Below 52° F, there was a significant inverse association between the maximum daily temperature and the rate of cases at 5 days.</li> <li>- In a theoretical state with a stable maximum daily temperature &gt;52° F, the rate of new cases is predicted to be of 23-fewer cases per-million per-day by 25 days of the epidemics.</li> <li>- A 1-unit higher UV index was associated with a lower rate at 5 days.</li> <li>- Precipitation was not associated with a greater rate of cases at 5 days.</li> </ul> <p>The incidence of disease is lower at warmer versus cooler temperatures, but this association is small, and transmission is likely to remain high at warmer temperatures.</p>
Cell Reports 30MAY2020	<b>Structural and biochemical characterization of nsp12-nsp7-nsp8 core polymerase complex from SARS-CoV-2</b>	Peng, Qi et al. China <a href="#">gotopaper</a>	Structural biology	<p>Cryo-EM structure of SARS-CoV-2 core polymerase complex (nsp12 catalytic subunit + nsp7-nsp8 cofactors):</p> <ul style="list-style-type: none"> <li>- Structure highly resembles SARS-CoV counterpart with conserved motifs for all viral RNA-dependent RNA polymerases, and suggests a mechanism for activation by cofactors.</li> <li>- <b>SARS-CoV-2 core complex has lower enzymatic activity than SARS-CoV.</b></li> <li>- <b>SARS-CoV-2 nsp7/8/12 subunits are less thermostable than the SARS-CoV counterpart.</b></li> </ul> <p>-&gt; Provides insights into RNA synthesis by coronavirus polymerase and <b>indicate adaptation of SARS-CoV-2 towards humans with relatively lower body temperatures than the natural bat hosts.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Clin. Infect. Dis. 30MAY2020	<b>Surgical mask partition reduces the risk of non-contact transmission in a golden Syrian hamster model for Coronavirus Disease 2019 (COVID-19)</b>	Chan, Jasper Fuk-Woo <i>et al.</i> China <a href="#">gotopaper</a>	Transmission - Animal model	<p>Golden Syrian hamster SARS-CoV-2 model to experimentally address effect of surgical mask on transmission :</p> <p>Surgical mask partition placed between cages of SARS-CoV-2-challenged index hamsters and naïve hamsters (closed system units separated by a polyvinyl chloride air porous partition + unidirectional airflow).</p> <p>- <b>Surgical mask partition for challenged hamsters significantly reduced transmission to 16.7% (2/12, P=0.019) of exposed naïve hamsters compared to exposed naïve hamsters without surgical mask partition (66.7%).</b></p> <p>- <b>Unlike severe COVID-19 manifestations of challenged hamsters, infected naïve hamsters had lower clinical scores, milder histopathological changes, and lower viral nucleocapsid antigen expression respiratory tract tissues.</b></p> <p>-&gt; <b>SARS-CoV-2 could be transmitted by respiratory droplets or airborne droplet nuclei in the hamster model. Such transmission could be reduced by surgical mask usage, especially when masks were worn by infected individuals.</b></p>
J Med Virol 29MAY2020	<b>Serum KL-6 concentrations as a novel biomarker of severe COVID19</b>	d'Alessandro, Miriana <i>et al.</i> Italy <a href="#">gotopaper</a>	Virology	<p>SARS-CoV-2 induced direct cytopathic effects against type I and II pneumocytes mediate lung damage. <b>Krebs von den Lungen-6 (KL-6) is mainly produced by damaged or regenerating alveolar type II pneumocytes.</b> This preliminary study analysed <b>serum concentrations of KL-6 in COVID19 patients to verify its potential as a prognostic biomarker of severity.</b></p> <p><b>CI°: NK cell analysis and assay of KL-6 in serum can help identify severe COVID19 patients.</b> Increased KL-6 serum concentrations were observed in patients with severe pulmonary involvement, revealing a prognostic value and supporting the potential usefulness of KL-6 measurement to evaluate COVID19 patients prognosis.</p> <p><b>Limitations:</b> these results are worthy of further validation in a larger cohort to define the cut-off value for identifying patients at high risk of severe respiratory failure.</p>
BMJ 29MAY2020	<b>Covid-19: the ethics of clinical research in quarantine</b>	Nicholas G Evans, USA <a href="#">gotopaper</a>	SHS/Sciences Po	<p>Quarantine = an opportunity to gain scientific knowledge of covid-19. Quarantine provides a model community in which to study both the social and epidemiological characteristics of a disease outbreak.</p> <p>As a closed system, quarantine offers the possibility for highly controlled research into the development and transmission of covid-19</p> <p>But this opportunity is also an ethical risk. This knowledge would be obtained with human rights violation (such as liberty). Moreover, individuals under quarantine seem to me more vulnerable to researchers. Confinement could affect their ability to choose or not to participate into research.</p> <p>Finally, research in this context could be a major opportunity to clarify when quarantine is, or is not, effective and proportionate (with high quality data on COVID-19).</p>
Journal of Allergy and Clinical Immunology 29MAY2020	<b>Successful use of methylprednisolone for treating severe COVID-19</b>	Liu, Jing <i>et al.</i> China <a href="#">gotopaper</a>	Therapeutic	<p><b>Case series of 101 consecutive hospitalized patients with confirmed COVID-19 infection, among which 26 were classified as severe or critical (25.74%), with at least 10 patients had a PaO<sub>2</sub>/FiO<sub>2</sub> ratio of less than 150 mmHg, treated with methylprednisolone.</b></p> <p>-&gt; <b>Timely and appropriate application</b> of glucocorticoid in severe and critical COVID-19 patients may <b>improve outcomes</b> and lung function and could avoid the need for invasive mechanical ventilation, compared with outcomes in reported studies.</p> <p>-&gt; Single-dose pulse methylprednisolone (40-500mg methylprednisolone) had <b>no apparent negative impact on SARS-CoV-2 removal and production of specific IgG.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature Communications 29MAY2020	<b>Detection of air and surface contamination by SARS-CoV-2 in hospital rooms of infected patients</b>	Chia, PY et al., Singapore <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Screening for SARS-CoV-2 RNA on surface and air samples from 3 airborne infection isolation rooms (AIIRs) in ICU and 27 AIIRs in the hospital general ward hosting Covid-19 patients. From 245 surface samples collected:</p> <ul style="list-style-type: none"> <li>- 56.7% of rooms have at least one surface contaminated</li> <li>- High touch surface contamination is shown in 10 (66.7%) out of 15 patient environments in the first week of illness, and 3 (20%) beyond the first week of illness</li> <li>- Air sampling performed in 3 of 27 general ward AIIRs ward detected SARS-CoV-2 PCR-positive particles of sizes &gt;4 microm and 1-4 microm in two rooms, despite these rooms having 12 air changes per hour.</li> </ul> <p>This warrants further study of the particle size distribution and airborne transmission potential of SARS-CoV-2.</p>
Science 29MAY2020	<b>Introductions and early spread of SARS-CoV-2 in the New York City area</b>	Gonzalez-Reiche et al, USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Aim: to identify the early transmission events underlying the rapid spread of the virus in the NYC metropolitan area, by sequencing SARS-CoV-2 from patients. Phylogenetic analysis of 84 distinct SARS-CoV-2 genomes indicated:</p> <ul style="list-style-type: none"> <li>- multiple, independent but isolated introductions mainly from Europe and other parts of the United States,</li> <li>- evidence for community transmission of SARS-CoV-2 as suggested by clusters of related viruses found in patients living in different neighborhoods of the city.</li> </ul>
Science Advances 29MAY2020	<b>Emergence of SARS-CoV-2 through recombination and strong purifying selection</b>	Li, Xiaojun <i>et al.</i> USA - China <a href="#">gotopaper</a>	Phylogenetic	<p>Localised genomic analysis of patterns of evolutionary recombination between CoVs from distinct host species that likely originated SARS-CoV-2, reveal:</p> <ul style="list-style-type: none"> <li>- <b>Strong purifying selection around the receptor binding motif (RBM) of the spike among bat, pangolin, and human coronaviruses.</b></li> <li>- <b>SARS-CoV-2's entire RBM was introduced through recombination with coronaviruses from pangolins</b>, possibly a critical step in the evolution of SARS-CoV-2's ability to infect humans.</li> </ul> <p><b>Note:</b> all 3 human CoVs (SARS, MERS and SARS-2) are the result of recombination among CoVs involving the S gene, likely a precondition to zoonosis that enabled efficient binding to human receptors.</p> <p><b>-&gt; Similar evolutionary selection in different host species, together with frequent recombination among coronaviruses, suggest a common evolutionary mechanism that could lead to new emerging human coronaviruses.</b></p>
Lancet 29MAY2020	<b>Mortality and pulmonary complications in patients undergoing surgery with perioperative SARS-CoV-2 infection: an international cohort study</b>	COVIDSurg Collaborative UK <a href="#">gotopaper</a>	Clinic	<p>International – multicentre – 235 hospitals – 24 countries Surgery + SARS-CoV-2 positive within 7 before or 30 days after → <b>1128 patients</b> Pulmonary complication: ARDS or pneumonia or unexpected postoperative ventilation 74% had emergency surgery and 24,8% elective surgery <b>30-day mortality= 23,8%</b> <b>Pulmonary complications= 51,2% with 38% of mortality</b> <u>Association with mortality (adjusted analysis):</u></p> <ul style="list-style-type: none"> <li>- Male sex OR: 1,75 [1,28 – 2,40]</li> <li>- &gt; or = 70 years OR: 2,30 [1,65 – 3,22]</li> <li>- ASAS grade 3-5 OR: 2,35 [1,57 – 3,53]</li> <li>- Malignant diagnosis OR: 1,55 [1,01 – 2,39]</li> <li>- Emergency surgery OR: 1,67 [1,06 – 2,63]</li> <li>- Major surgery OR: 1,52 [1,01 – 2,31]</li> </ul> <p>→ pulmonary complication in more than half of patients with perioperative SARS-CoV-2 infection → <b>postponing non urgent procedure and promoting non operative treatment</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Journal of clinical investigation 28MAY2020	<b>Impaired immune cell cytotoxicity in severe COVID-19 is IL-6 dependent</b>	Mazzoni, Alessio et al, Italy <a href="#">gotopaper</a>	Immuno	<p><b>Characterization of the immune response in SARS-CoV-2 infected patients hospitalized at the Careggi University Hospital, Florence, Italy</b></p> <ul style="list-style-type: none"> <li>A flow cytometric characterization of immune cells subsets from 30 COVID-19 patients correlated with clinical outcomes confirms a decreased number of circulating T, B and NK cells and that T CD4+, T CD8+ but also NK cells displayed reduced anti-viral cytokine production capability.</li> <li>The study shows a skewing of CD8+ T cells towards a terminally differentiated/senescent phenotype via a TNF-mediated T cell apoptosis. This may contribute to an uncontrolled inflammatory response.</li> <li>In a group of patients that required intensive care, serum IL-6 levels are inversely correlated with the frequency of granzyme-expressing NK cells. This underlines that the exposure to high levels of IL-6 inhibits NK cell cytotoxicity and down-regulates the expression of perforin and granzyme.</li> <li>In all patients treated with tocilizumab an increased expression of both perforin and granzyme in NK cells was observed as well as a decrease of CRP, which is considered as a marker of IL-6 mediated inflammation. This study points out that tocilizumab might restore the cytotoxic potential of NK cells.</li> </ul> <p>Conclusion: This current work endorses that targeting IL-6 cytokine might restore anti-viral mechanisms and the use an anti-IL-6 receptor monoclonal antibody in COVID-19 patients remains a potential therapeutic option.</p>
Natl Sci Rev 28MAY2020	<b>Preliminary evidence from a multicenter prospective observational study of the safety and efficacy of chloroquine for the treatment of COVID-19</b>	Huang, Mingxing et al. China <a href="#">gotopaper</a>	Therapeutic	<p><b>Multicenter prospective observational study</b> to assess the <b>efficacy and safety of chloroquine</b> with different doses in COVID-19.</p> <p>A total of 197 patients completed chloroquine treatment, and 176 patients treated with non-chloroquine therapy were included as historical controls. Across the two treatment groups, the majority patients were classified as <b>moderate cases</b> (93.4% in chloroquine; 89.2% in nonchloroquine).</p> <p>The primary endpoint is the <b>time to undetectable viral RNA</b>. Secondary outcomes include the proportion of patients with undetectable viral RNA by day 10 and 14, hospitalization time, duration of fever, and adverse events.</p> <ul style="list-style-type: none"> <li>Patients in the chloroquine group experienced <b>significantly faster and higher rate of viral suppression</b> comparing to the nonchloroquine group in both the full analysis and the post hoc stratified analysis. Even when the dose reduced to half, the benefit of chloroquine still remained.</li> <li><b>Duration of fever is shorter</b> in chloroquine (geometric mean ratio 0.6; 95% CI 0.5 to 0.8).</li> <li><b>No serious adverse events</b> were observed in the chloroquine group. Patients treated with half dose experienced lower rate of adverse events than with full dose.</li> <li><b>No beneficial effect</b> of chloroquine in the <b>length of hospital stay</b> and the <b>duration of oxygen support</b>.</li> <li>Unprecedentedly, 3 cases of so called <b>“re-positive” patients</b> observed in the chloroquine group.</li> </ul> <p><b>Limitations:</b> observational study ; dramatic improvement in the primary outcome in chloroquine could be due to the later treatment initiation since symptom onset ; relatively young study population with few patients with severe symptoms that requires ventilation.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Lancet HIV 28MAY2020	<b>Description of COVID-19 in HIV-infected individuals: a single-centre, prospective cohort</b>	Vizcarra P et al Spain <a href="#">gotopaper</a>	Clinic	<p>Observational &amp; prospective study: 51 HIV-infected COVID-19+ &amp; 1288 HIV-infected without 1 center in Madrid Mean age of COVID-19= 53,3y (SD:9,5) – 84% of men <b>No difference for age and CD4 cell counts</b> 63% with COVID-19 had at least one comorbidity Clinical presentations similar than in general population 12% critically ill and 4% died <u>Covid-19 vs without COVID-19:</u>  <ul style="list-style-type: none"> <li>- 73% vs 32% received tenofovir before COVID-19 diagnosis (<math>p=0,0036</math>)</li> <li>- 22% vs 14% had previous protease inhibitor (<math>p=0,578</math>)</li> </ul> </p> <p>RT-PCR remained positive after a median of 40 days in 6 patients</p> <p>→ HIV-infected patients should receive the same treatment to the general population</p>
Lancet 28MAY2020	<b>Clinical impact of COVID-19 on patients with cancer (CCC19): a cohort study</b>	Kuderer NM et al USA <a href="#">gotopaper</a>	Clinic	<p>Cohort study from the USA, Canada &amp; Spain from the COVID-19 &amp; cancer consortium (CCC19) Patients with active or previous malignancy &amp; SARS-CoV-2 infection: <b>928 patients included</b> <u>Malignancy:</u> breast cancer (21%) – prostate (16%) – gastrointestinal (12%) – thoracic (10%) ... <u>Status:</u> active cancer (43%) – remission (45%) – anticancer treatment (39%) <b>13% had died</b> Median age: 66 y IQR [57 – 76] – 50% were male <u>Independent factor associated with increased 30-day mortality</u>  <ul style="list-style-type: none"> <li>- Increased age per 10 years OR: 1,84 [1,53 – 2,21]</li> <li>- Male OR: 1,63 [1,07 – 2,28]</li> <li>- Former smoker vs never OR: 1,60 [1,03 – 2,47]</li> <li>- Two comorbidities vs none OR: 4,50 [1,33 – 15,28]</li> <li>- Active cancer OR: 5,2 [2,77 – 9,77]</li> <li>- Receipt azithro+hydroxy vs neither OR: 2,93 [1,79 – 4,79] → <i>confounding by indication not excluded</i></li> </ul> </p> <p>Obesity, cancer type, ethnicity, type of anticancer therapy, recent surgery → not associated</p> <p>Limits: regional variations in the primary and secondary outcomes</p> <p>→ High mortality among patients with cancer + COVID-19 → <b>longer follow-up is needed</b></p>
The Lancet ID 28MAY2020	<b>Implication of SARS-CoV-2 evolution in the sensitivity of RT-qPCR diagnostic assays</b>	Sampaio Osorio and Correia-Neves, Portugal <a href="#">gotopaper</a>	Diagnostic	<p>Reports suggest the virus might be evolving → redesign of the oligonucleotide sequences in use in RT-qPCR assays to circumvent potential primer–sample mismatches ? Analysis of all high-coverage SARS-CoV-2 genome sequences (1825 in total) deposited in the Global Initiative on Sharing All Influenza Data (GISAID) database nucleotide diversity (<math>\pi</math>) WAS CALCULATED in the binding region of each oligonucleotide.</p> <p>-&gt; 79% (26 of 33) of the primer binding sites used in the RT-qPCR assays were mutated in at least one genome</p> <p>-&gt; at least one of the previously designed primers is now likely to be ineffective at detecting up to 14% of the virus variants in circulation</p> <p>-&gt; Oligonucleotide optimisation will be facilitated by global sharing of SARS-CoV-2 genomes and the frequently updated reports on sequence analysis that are available on the GISAID website.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
BMJ 28MAY2020	Kawasaki-like multisystem inflammatory syndrome in children during the covid-19 pandemic in Paris, France: prospective observational study	Toubiana J et al France <a href="#">gotopaper</a>	Clinic	<p><b>21 children in a hospital in Paris:</b> median age: 7,9y [3,7 – 16,6] – 43% male            9 had a recent history of a viral like symptoms – 10 reported a recent contact            Children of African ancestry were overrepresented (57%)</p> <p><u>Symptoms</u></p> <ul style="list-style-type: none"> <li>- 57% presented with Kawasaki disease shock syndrome</li> <li>- 76% presented with myocarditis</li> <li>- 100% gastrointestinal symptoms during the early stage</li> </ul> <p><u>Biology:</u></p> <ul style="list-style-type: none"> <li>- High level of inflammatory markers (PCT, CRP, IL-6)</li> <li>- 81% had lymphopenia</li> <li>- 95% had elevated D-dimer (&gt;500 µg/L)</li> <li>- 90% recent SARS-CoV-2 infection (PCR or IgG antibody)</li> </ul> <p><u>Treatment:</u></p> <ul style="list-style-type: none"> <li>- All received intravenous Ig</li> <li>- 48% received also corticosteroid</li> </ul> <p><u>Outcomes:</u></p> <ul style="list-style-type: none"> <li>- 81% required intensive care</li> <li>- 24% had moderate coronary artery dilations</li> <li>- <b>No death</b> – all discharged home after 8 days of hospital stay</li> </ul> <p>→ high proportion of children with KD shock syndrome – <b>differ from classic KD</b>            → temporal association between the covid-19 pandemic &amp; results of test for SARS-CoV-2 in children with Kawasaki-like disease suggest a causal link → <b>further study is needed</b></p>
BMJ 27MAY2020	Characteristics and outcomes of pregnant women admitted to hospital with confirmed SARS-CoV-2 infection in UK: national population based cohort study	Knight M et al UK <a href="#">gotopaper</a>	Clinic	<p>National cohort of pregnant women hospitalized with SARS-CoV-2 → <b>427 women</b>            Estimated incidence: 4,9 per 1000 maternities [4,5 – 5,4]</p> <p><u>Comorbidities:</u></p> <ul style="list-style-type: none"> <li>- 69% were obese or overweight</li> <li>- 34% had co-existence comorbidities: asthma, hypertension, diabetes, cardiac disease</li> </ul> <p><u>Symptoms</u></p> <ul style="list-style-type: none"> <li>- Have symptom at median of 34 completely weeks'</li> <li>- Most have symptom on the third trimester: fever – cough – breathlessness ...</li> </ul> <p><u>Outcomes</u></p> <ul style="list-style-type: none"> <li>- 62% women admitted gave birth or had a pregnancy loss</li> <li>- 15% women were given corticosteroids for fetal lung maturation</li> <li>- 196 women gave birth at term and 66 preterm → 265 infants</li> <li>- 41 needed respiratory support</li> <li>- 5 women died</li> </ul> <p><u>Neonatal</u></p> <ul style="list-style-type: none"> <li>- 5 babies died: 3 stillborn &amp; 2 neonatal period</li> <li>- 12 were positive for SARS-CoV-2 which 6 within the 12 hours → no death</li> </ul> <p>→ <b>Most women had good outcome</b>            → <b>transmission to infant was uncommon</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Gene Reports 2020	<b>COVID-19 target: A specific target for novel coronavirus detection</b>	Kakhki, Reza Kamali et al. Iran <a href="#">gotopaper</a>	Diagnostic	<p><b>The diagnosis and differentiation of this virus from other types of coronavirus is essential to control of the disease</b></p> <p>The <b>analysis of genomics data</b> plays a vital role in introducing a stronger target and consequently provides better results in laboratory examinations. The modified comparative genomics approach helps us to find novel specific targets by comparing two or more sequences on the nucleotide collection database.</p> <p>Unlike previous reported genes (RdRP, E and N genes), <b>ORF8 is entirely specific to the novel coronavirus (COVID-19)</b> and has no cross-reactivity with other kinds of coronavirus <b>CI*</b>: ORF8 gene can be used as an additional confirmatory assay.</p> <p><b>Limitations:</b> this study was conducted bioinformatically, and laboratory examinations are needed to confirm ORF8 gene as a potential target using RT-PCR, Real time PCR, or Line probe assay</p>
Critical care medicine 27MAY2020	<b>Routine Venous Thromboembolism Prophylaxis May Be Inadequate in the Hypercoagulable State of Severe Coronavirus Disease 2019</b>	Maatman, Thomas K. et al. USA <a href="#">gotopaper</a>	Therapeutic/ Clinic	<p><b>Observational multicenter study</b>, enrolled 240 consecutive patients among whom <b>109 critically ill COVID-19</b> patients admitted to the ICU were included in the analysis. All patients received <b>routine subcutaneous chemical venous thromboembolism prophylaxis</b>.</p> <p>Primary outcome: <b>frequency of venous thromboembolism (VTE)</b> and the degree of inflammatory and coagulation marker elevation associated with venous thromboembolism development.</p> <ul style="list-style-type: none"> <li>- <b>VTE was diagnosed in 31 patients (28%)</b> 8 ± 7 days after hospital admission, including two patients diagnosed with venous thromboembolism at presentation to the hospital.</li> <li>- Elevated admission D-dimer and peak D-dimer were associated with VTE development (p &lt; 0.05).</li> <li>- D-dimer greater than 2,600 ng/mL predicted VTE with an area under the receiver operating characteristic curve of 0.760 (95% CI, 0.661-0.858; p &lt; 0.0001), sensitivity of 89.7%, and specificity of 59.5%.</li> <li>- Twelve patients (11%) had thromboelastography performed and 58% of these patients had a hypercoagulable study. The calculated coagulation index was hypercoagulable in 50% of patients with thromboelastography</li> <li>-&gt; SARS-CoV-2 infection results in <b>systemic hypercoagulability resulting in VTE</b>. Although current data on outcomes in patients receiving therapeutic anticoagulation in COVID-19 are lacking, it is apparent that <b>routine chemical VTE prophylaxis may be inadequate in preventing thrombotic complications in severe COVID-19</b>.</li> </ul> <p><b>Limitations:</b> Although a relatively large series, it is an observational study ; retrospective study ; lack of comparator group of ICU patients without COVID-19 in which to compare VTE frequency ; degree of viremia not evaluated as a risk factor for VTE.</p>
NEJM 27MAY2020	<b>Remdesivir for 5 or 10 Days in Patients with Severe Covid-19</b>	Jason D. Goldman et al. <a href="#">gotopaper</a>	Therapeutic	<p><b>Open-label, randomized, multicenter trial</b> evaluating the <b>efficacy and safety</b> of treatment with <b>remdesivir for 5 or 10 days</b> in <b>397 patients</b> with <b>severe Covid-19</b> disease. The primary end point was <b>clinical status on day 14</b>, assessed on a 7-point ordinal scale.</p> <p>=&gt; <b>No significant difference in efficacy between 5-day and 10-day courses of remdesivir</b>.</p> <p>After adjustment for baseline imbalances in disease severity, outcomes were similar as measured by a number of end points: clinical status at day 14, time to clinical improvement, recovery, and death from any cause. Similar percentages of patients experiencing adverse events.</p> <p>-&gt; Results cannot be extrapolated to critically ill patients receiving mechanical ventilation; further evaluation of this subgroup and of other high-risk groups is needed to determine the shortest effective duration of therapy.</p> <p><b>Limits:</b> lack of a randomized placebo control group (magnitude of benefit not determined); open-label design.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
SCIENCE 27MAY2020	Reducing transmission of SARS-CoV-2	Prather et al., USA-China <a href="#">gotopaper</a>	Public Health/Epidemiology	<p><b>PERSPECTIVE</b></p> <ul style="list-style-type: none"> <li>- Traditional respiratory disease control measures: designed to reduce transmission by droplets produced in the sneezes and coughs of infected individuals. However, a large proportion of the spread: through <u>airborne transmission of aerosols</u> produced by asymptomatic individuals during breathing and speaking. Aerosols can <b>accumulate, remain infectious in indoor air for hours</b>, and be easily inhaled deep into the lungs. Contact (direct or indirect) is also a major source of contamination.</li> <li>- Respiratory droplet size has been shown to affect the severity of disease</li> <li>- “Silent shedders” (asymptomatic / pre-symptomatic) could be critical drivers. In China, undiagnosed cases, presumably asymptomatic, may be responsible for up to 79% of infections.</li> <li>- Many countries have not yet acknowledged airborne transmission as a possible pathway</li> </ul> <p>Recommendations for social distancing of 6 ft are based on studies of respiratory droplets carried out in the 1930s, but on large droplets (no technology for submicron aerosols). -&gt;Intense coughs and sneezes that propel larger droplets more than 20 ft can also create thousands of aerosols that can travel even further (<u>like a cigarette smoke</u>)-&gt; <b>6 ft WHO recommendation is likely not enough.</b></p> <p>-Viruses can attach to other particles such as dust and pollution, which can modify the aerodynamic characteristics and increase dispersion</p> <p><b>Aerosol transmission of viruses must be acknowledged as a key factor leading to the spread of infectious respiratory diseases. Evidence suggests that SARS-CoV-2 is silently spreading in aerosols exhaled by highly contagious infected individuals with no symptoms.</b></p> <p><b>-&gt; It is essential that control measures be introduced to reduce aerosol transmission (face masks +++)</b></p>
Nature reviews Immunology 26MAY2020	Dysregulation of type I interferon responses in COVID-19	Dhiraj Acharya et al, USA <a href="#">gotopaper</a>	Immunology	<p>How imbalanced interferon responses may contribute to the pathology of COVID-19:</p> <ul style="list-style-type: none"> <li>• The lung injury in patients with severe COVID-19 underlines a possible failure to activate immunosuppressive mechanisms in a timely manner. One suggestion is that the deficient or dysregulated IFN responses elicited by SARS-CoV-2 infection may influence the generation of regulatory T cells during the recovery phase of COVID-19.</li> <li>• The use of IFNs as a treatment for COVID-19 remains controversial, particularly regarding the timing of administration in mice model and in human</li> <li>• Since ACE2 has been identified as an ISG in human airway epithelial cells<sup>10</sup>. This finding raises the question of whether prophylactic or therapeutic IFN administration enhance the entry and replication of SARS-CoV-2 in target cells during disease progression</li> </ul> <p><b>Conclusion:</b> a deeper understanding of the spatiotemporal kinetics of IFN responses during clinical SARS-CoV-2 infections is warranted to inform IFN-related therapeutics and vaccine design.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Cell Death Discovery 26MAY2020	<b>SARS-CoV-2 infection serology: a useful tool to overcome lockdown?</b>	Nuccetelli, M. et al, Italy <a href="#">gotopaper</a>	Diagnostic	<p><b>Aim:</b> to compare and to evaluate different serological assays analytical performances (two different immunochromatographic cards, an immunofluorescence chromatographic card, and a chemiluminescence-automated immunoassay) on 43 positive samples with RT-qPCR-confirmed SARS-CoV-2 infection and 40 negative control subjects.</p> <p>→ excellent IgG/IgM specificities for all the immunochromatographic card tests (100% IgG and 100% IgM) and for the chemiluminescence-automated assay (100% IgG and 94% IgM);</p> <p>→ IgG/IgM sensitivities are moderately lower for all methods (94% and 84% for IgG and IgM, respectively), probably due to the assay viral antigen's nature and/or to the detection time of nasopharyngeal swab RT-qPCR, with respect to symptoms onset.</p>
Clinical Infectious Diseases 25MAY2020	<b>Characterization of an asymptomatic cohort of SARS-COV-2 infected individuals outside of Wuhan, China</b>	Wang Y et al China <a href="#">gotopaper</a>	Clinic	<p>Epidemiologic and clinical characteristics of asymptomatic SARS-COV-2 infections 279 hospitalized SARS-CoV-2+ contacts of COVID-19 patients → <b>63 asymptomatic</b> included Mean time to diagnosis after contact: 16 days Mean age: 39,3 - 87,3% had no comorbidities <u>Laboratory findings:</u> quasi normal for all</p> <p><b>2 groups:</b> abnormal chest CT findings (29) &amp; normal chest CT findings (34) - Patient with abnormal findings were older (<math>p &lt; 0,05</math>) - Time from exposure to illness shorter in patient with abnormal CT (<math>p &gt; 0,05</math>)</p> <p><u>Outcomes:</u> - 9 transmitted the virus to others with and without abnormal chest CT - No one died → asymptomatic infections play a large role in transmission → impact on treatment of symptomatic cases on transmission?</p>
The Lancet 25MAY2020	<b>Assistance Publique–Hôpitaux de Paris' response to the COVID-19 pandemic</b>	The COVID19- APHP Group, France <a href="#">gotopaper</a>	Public Health/Epidemio	<p><b>Key points of an effective AP-HP response to the Covid-19 crisis:</b></p> <ul style="list-style-type: none"> <li>- Establishment of a medical organisation led by a central crisis medical director and supported by medical directors in each hospital</li> <li>- Allocation of human resources to recruit and train specialised staff from a single platform</li> <li>- Centralised logistics to adjust to shortages the equipment and consumables daily supply</li> <li>- Recruitment of students of various medical branches to support as paramedics, research assistants, operators of a telemedicine platform.</li> <li>- Regional centralised ICU bed-allocation system</li> <li>- Regularly updated practical guidelines for all hospitals</li> <li>- Development of a common research strategy prioritising patient cohorts, biobanking, clinical trial</li> <li>- Large scale initiatives: equipment 3D printing, Covidom telemedicine platfor</li> <li>- Setup of a region-wide patient-tracing programme</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature Climate Change 23MAY2020	<b>Temporary reduction in daily global CO2 emissions during the COVID-19 forced confinement</b>	Le Quéré et al. UK, <a href="#">gotopaper</a>	SHS/SciPo	<p>&gt; Emissions of carbon dioxide rise by about 1% per year over the previous decade</p> <p>&gt; COVID 19 imposed confinement leads to drastic changes in energy use, with expected impacts on CO2 emissions. However CO2 emissions are reported as annual values and there is no available real time data</p> <p>&gt; An alternative approach using a combination of energy, activity and policy data was used to estimate changes in CO2 daily emissions during the confinement is proposed (69 countries, 50 US states and 30 Chinese provinces; 85% of global population; 87% of global emissions)</p> <p><b>Results</b></p> <p>&gt; Estimated decrease in daily fossil CO2 emissions from the severe and forced confinement of world populations: -17% (-11 to -25%)</p> <p>&gt; Annual associated decrease: -4.2 to -7.5% (rates needed over the next decades to limit climate change to 1.5 °C warming)</p> <p>&gt; However observed changes are likely to be temporary as they do not reflect structural changes in the economic, transport or energy systems.</p>
Biosensors and Bioelectronics 23MAY2020	<b>Ultra-sensitive and high-throughput CRISPR-Powered COVID-19 diagnosis</b>	Huang, Zhen and al. USA <a href="#">gotopaper</a>	Diagnostic	<p>A rapid, sensitive SARS-CoV-2 diagnostic assay capable of high-throughput operation that can preferably utilize existing equipment to facilitate broad, large-scale screening efforts.</p> <p>The developed assay utilizes a custom CRISPR Cas12a/gRNA complex and a fluorescent probe to amplify target amplicons produced by standard RT-PCR or isothermal recombinase polymerase amplification (RPA), to allow sensitive detection at sites not equipped with real-time PCR systems required for qPCR diagnostics.</p> <p>The results obtained on nasal swab samples of individuals with suspected COVID-19 cases were comparable to paired results from a CDC-approved qPCR assay performed in a state testing lab, and superior to those produced by same assay in a clinical lab, where the qPCR assay exhibited multiple invalid or inconclusive results. It also demonstrated greater analytical sensitivity and more robust diagnostic performance than other recently reported CRISPR-based assays.</p> <p>CI*: a CRISPR-based fluorescent application has potential to improve current COVID-19 screening efforts.</p>
BMJ 22MAY2020	<b>Features of 20 133 UK patients in hospital with covid-19 using the ISARIC WHO Clinical Characterisation Protocol: prospective observational cohort study</b>	Docherty, AB. Et al. UK <a href="#">gotopaper</a>	Clinic	<p>Aim: To characterise the clinical features of 20 133 hospital patients with Covid-19 enrolled in the ISARIC WHO CCP-UK prospective cohort study, and to explore risk factors associated with admission to critical care and mortality in hospital. Patient follow-up time was of two weeks minimum.</p> <p>→ Median age: 73 years (interquartile range (IR) 58-82, range 0-104). 60% of patients were men, 40% women. Median duration of symptoms before admission was 4 days (IR 1-8).</p> <p>→ Comorbidities: chronic cardiac disease (31%), uncomplicated diabetes (21%), non-asthmatic chronic pulmonary disease (18%), chronic kidney disease (16%); 23% had no reported major comorbidity.</p> <p>→ 41% of patients were discharged alive, 26% died, 34% continued receiving care as of reporting date. 17% (3001/18 183) required admission to high dependency or ICU; of these, 28% were discharged alive, 32% died, and 41% continued to receive care. Of those receiving mechanical ventilation (1658), 17% were discharged alive, 37% died, and 46% remained in hospital. Increasing age, male sex, and chronic comorbidities were associated with higher mortality in hospital.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Circulation 22MAY2020	<b>Cardiovascular Toxicities Associated with Hydroxychloroquine and Azithromycin: An Analysis of the World Health Organization Pharmacovigilance Database</b>	Nguyen, Lee S. et al. France/USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Observational, retrospective study</b>, that used VigiBase®, the <b>WHO pharmacovigilance database</b> encompassing over 21 million reports from over 130 countries, to compare CV-ADR reporting in patients who received <b>hydroxychloroquine, azithromycin, or their combination</b> with <b>cardiovascular adverse-drug-reactions</b> (CV-ADRs) reported with all other drugs in the full database.</p> <p>Extraction of 76,822 ADR cases associated with hydroxychloroquine alone, 89,692 with azithromycin alone, and 607 with the combination of both drugs. The cases were retrieved from 21,275,867 total ADR reports in VigiBase®. The lower end of the IC's 95% credibility interval is IC025. It is considered significant when above 0.</p> <ul style="list-style-type: none"> <li>- Significant greater reporting of <b>prolonged-QT (LQT)</b> and/or <b>ventricular tachycardia</b> including <b>Torsades-de-Pointes</b> (TdP/VT) for each drug individually in suspected cases (IC025=1.67 for azithromycin and IC025=1.04 for hydroxychloroquine).</li> <li>- Hydroxychloroquine was also associated with <b>conduction disorders</b> (atrioventricular and bundle branch blocks) (IC025=1.04) and heart failure (HF, IC025=0.06).</li> <li>- Azithromycin monotherapy was associated with a greater reporting of LQT and/or TdP/VT than hydroxychloroquine monotherapy. The <b>combination of azithromycin and hydroxychloroquine</b> was associated with a <b>greater reporting of LQT and/or TdP/VT</b> than either drug in monotherapy.</li> <li>- The proportion that resulted in <b>death for TdP/VT</b> cases was 8.4% with hydroxychloroquine and 20.2% with azithromycin versus 0% and 5.4% for LQT without TdP/VT with hydroxychloroquine and azithromycin, respectively (p&lt;0.001 for both).</li> <li>- Corresponding death rate was 20.7% for HF associated with hydroxychloroquine. Dose of hydroxychloroquine was higher in HF compared to LQT and/or TdP/VT cases.</li> </ul> <p><b>Main limitation:</b> without data on numbers exposed in Vigibase, this work cannot assess the incidence or risk for QT prolongation with these drugs.</p>
Annals of Translational Medicine MAY2020	<b>Preliminary study to identify severe from moderate cases of COVID-19 using combined hematology parameters</b>	Wang, Changzheng et al. China <a href="#">gotopaper</a>	Diagnostic	<p>The aim of this study was to investigate the <b>characteristics and rules of hematology changes in patients with COVID-19</b>, and to explore the possibility differentiating moderate and severe patients using conventional hematology parameters or combined parameters</p> <p>As the disease progressed, white blood cell count (WBC), neutrophil count, neutrophil-to-lymphocyte ratio (NLR), platelet-to-lymphocyte ratio (PLR), red blood cell distribution width-coefficient of variation (RDW-CV), and red cell volume distribution width-standard deviation (RDW-SD) parameters in the severe group were significantly higher than those in the moderate group.</p> <p><b>CI°: the combined NLR and RDW-SD parameter is the best hematology index.</b> It may help clinicians to predict the severity of COVID-19 patients and can be used as a useful indicator to help prevent and control the epidemic</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Gastroenterology 22MAY2020	<b>Famotidine Use is Associated with Improved Clinical Outcomes in Hospitalized COVID-19 Patients: A Propensity Score Matched Retrospective Cohort Study</b>	Freedberg, Daniel E. et al. USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Retrospective cohort study</b> of 1,620 hospitalized patients tested positive for SARS-CoV-2 within 72 hours following admission including <b>84 patients</b> who received <b>famotidine</b> within 24 hours of hospital admission.</p> <p><b>Primary outcome</b> was a composite of death or endotracheal intubation.</p> <p>Although Famotidine has not previously been studied in patients for antiviral effects, an untargeted computer modelling analysis identified famotidine as one of the highest-ranked matches for drugs <b>predicted to bind 3CLpro</b>, a SARS-CoV-2 protease which generates non-structure proteins critical to viral replication.</p> <p>340 (21%) patients met the composite study outcome. Famotidine use was <b>significantly associated</b> with a <b>reduced risk of clinical deterioration leading to intubation or death</b>. A randomized controlled trial is currently underway to determine whether famotidine can improve clinical outcomes in hospitalized COVID-19 patients.</p> <p><b>Limitations:</b> observational; no samples were gathered, and mechanism cannot be directly assessed; single center study.</p>
Clinical infectious diseases 22MAY2020	<b>Association of renin-angiotensin-aldosterone system inhibitors with COVID-19-related outcomes in Korea: a nationwide population-based cohort study</b>	Jung, Sun-Young et al. Korea <a href="#">gotopaper</a>	Therapeutic/ Clinic	<p><b>Nationwide population-based cohort study</b> in Korea comparing the <b>clinical outcomes</b> of confirmed COVID-19 cases between <b>RAAS inhibitor users and nonusers</b>.</p> <p>The study revealed a <b>significantly higher mortality rate</b> among patients with COVID-19 who were <b>using RAAS inhibitors</b>, relative to patients who were not receiving RAAS inhibitors. However, <b>RAAS inhibitor users were older</b>, had <b>more comorbidities</b>, and were <b>more likely to receive in-hospital treatments</b>. The <b>elevated risk of mortality</b> among RAAS inhibitor users <b>disappeared after adjusting for these confounding factors</b>.</p> <p>This study in an <b>Asian population</b> is clinically relevant, given that the East Asian populations <b>have higher ACE2 expression in tissues</b> than other populations under the similar conditions.</p> <p><b>Limitations:</b> accuracy of diagnostic codes may be limited; retrospective observational design.</p>
Clinical infectious diseases 22MAY2020	<b>Thymosin alpha 1 (Talpha1) reduces the mortality of severe COVID-19 by restoration of lymphocytopenia and reversion of exhausted T cells</b>	Liu, Yueping et al. China <a href="#">gotopaper</a>	Therapeutic	<p><b>Retrospective cohort study</b> to evaluate the clinical outcomes of <b>severe or critical</b> COVID 19 hospitalized patients receiving <b>Thymosin alpha 1 (Tα1)</b> supplement. A total of <b>76 patients</b> were enrolled (36 in the treatment group and 40 in the non treatment group)</p> <p>Compared with untreated group, <b>Tα1 treatment significantly reduces mortality</b> of severe COVID-19 patients (11% vs. 30%, p=0.044). <b>Tα1 timely enhances blood T cell numbers</b> in COVID-19 patients with <b>severe lymphocytopenia</b>. Under such conditions, Tα1 also successfully <b>restores CD8+ and CD4+ T cell numbers in aged patients</b>. Meanwhile, Tα1 <b>reduces PD-1 and Tim-3 expression on CD8+ T cells</b> from severe COVID-19 patients in comparison with untreated cases.</p> <p><b>Limitations:</b> issue with the normalization of TREC levels among individuals; retrospective study and small sample size; mortality as primary clinical outcome and not clinical improvement.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
NEJM 22MAY2020	Remdesivir for the Treatment of Covid-19 — Preliminary Report	Beigel, John H. et al., USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Double-blind, randomized, placebo-controlled trial of intravenous remdesivir in 1063 adults hospitalized with Covid-19 with evidence of lower respiratory tract involvement in 60 trial sites in 10 countries.</b></p> <p>The <b>primary outcome</b> was the <b>time to recovery</b>, defined by either discharge from the hospital or hospitalization for infection-control purposes only.</p> <p><b>Early unblinding</b> of the results recommended by the DSMB based on findings from an analysis that showed shortened time to recovery in the remdesivir group.</p> <p>Preliminary results from the 1059 patients (538 assigned to remdesivir and 521 to placebo) suggest that a <b>10-day course of remdesivir was superior to placebo in the treatment of hospitalized patients with Covid-19.</b></p> <ul style="list-style-type: none"> <li>- This benefit was seen in the <b>number of days to recovery</b> (median, 11 days, as compared with 15; rate ratio for recovery, 1.32 [95% CI, 1.12 to 1.55]) and in <b>recovery according to the ordinal scale score at day 15</b> (odds ratio, 1.50; 95% CI, 1.18 to 1.91).</li> <li>- Mortality was numerically lower in the remdesivir group than in the placebo group, but the difference was not significant (hazard ratio for death, 0.70; 95% CI, 0.47 to 1.04; 1059 patients).</li> <li>- Incidence of adverse events was not found to be significantly different between the remdesivir group and the placebo group.</li> </ul> <p>Awaiting final visits, data entry, monitoring, and data lock for the last of the 1063 patients enrolled, to provide an update of the results.</p>
The Lancet 22MAY2020	Hydroxychloroquine or chloroquine with or without a macrolide for treatment of COVID-19: a multinational registry analysis	Mandeep R Mehra et al., USA <a href="#">gotopaper</a>	Therapeutic	<b>RETRACTED</b>
mBio 22MAY2020	Antiviral Efficacies of FDA-Approved Drugs against SARS-CoV-2 Infection in Ferrets	Park, Su-Jin et al. Korea <a href="#">gotopaper</a>	Therapeutic	<p>FDA-approved drugs <b>lopinavir-ritonavir</b>, <b>hydroxychloroquine sulfate</b>, and <b>emtricitabine-tenofovir</b> were <b>tested against SARS-CoV-2</b> infection in a highly susceptible <b>ferret infection model</b>.</p> <p>While <b>most of the drug treatments marginally reduced clinical symptoms</b>, they <b>did not reduce virus titers</b>, with the <b>exception of emtricitabine-tenofovir</b> treatment, which led to diminished virus titers in nasal washes at 8 dpi.</p> <p>Further, the <b>azathioprine-treated immunosuppressed ferrets</b> showed delayed virus clearance and low SN titers, resulting in a prolonged infection.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
BMJ 22MAY2020	<b>Factors associated with hospital admission and critical illness among 5279 people with coronavirus disease 2019 in New York City: prospective cohort study</b>	Petrilli CM, et al., USA <a href="#">gotopaper</a>	Clinic	<p><b>Aim:</b> To describe outcomes, and clinical and laboratory characteristic associated with severity of illness of 5279 patients admitted with Covid-19 in NYC/Long Island (US).</p> <p>→ 48.2% of people tested (5566/11544) were positive for SARS-CoV2; of the 5279 patients included, 51.9% were admitted to hospital. Of these, 69.5% were discharged without hospice care, 24.3% were discharged to hospice care or died. Of 647 (23.6% of hospitalised) patients requiring mechanical ventilation, 60.4% died and 26.2% were extubated or discharged.</p> <p>→ Risk for hospital admission was associated with age (odd ratio &gt;2 for age groups &gt;44 years, 37.9 (95% CI[26.1 to 56.0]) for &gt;75 years), heart failure (4.4, 2.6 to 8.0), male sex (2.8, 2.4 to 3.2), chronic kidney disease (2.6, 1.9 to 3.6), increase in body mass index (BMI) (eg, for BMI &gt;40: 2.5, 1.8 to 3.4).</p> <p>→ Risk for critical illness besides age was associated with heart failure (1.9, 1.4 to 2.5), BMI &gt;40 (1.5, 1.0 to 2.2), and male sex (1.5, 1.3 to 1.8). Admission oxygen saturation of &lt;88% (3.7, 2.8 to 4.8), troponin level &gt;1 (4.8, 2.1 to 10.9), C reactive protein level &gt;200 (5.1, 2.8 to 9.2), and D-dimer level &gt;2500 (3.9, 2.6 to 6.0) were more strongly associated with critical illness than age or comorbidities.</p> <p>→ Risk of critical illness decreased significantly over the study period, and similar associations were found for mortality alone, potentially <b>suggesting improvement in care.</b></p>
The Lancet 22MAY2020	<b>Safety, tolerability, and immunogenicity of a recombinant adenovirus type-5 vectored COVID-19 vaccine: a dose-escalation, open-label, non-randomised, first-in-human trial</b>	Zhu FC et al. China, <a href="#">gotopaper</a>	Vaccine	<p>Phase 1 vaccine trial using the recombinant non-replicating adenovirus 5 type vectored COVID 19 vaccine expressing the S protein (open label; non randomised; dose escalation-5x10<sup>10</sup>/1x10<sup>11</sup>/5x10<sup>11</sup> viral particles). NCT04313127 108 patients recruited (18-60 years; 51% male-49% female)</p> <p><b>Endpoint for safety:</b> 7D post vaccination/recording of AE until 28D post-vaccination</p> <p><b>Humoral immunogenicity endpoints:</b> The specific ELISA antibody titres to RBD and S protein, and the neutralising antibody amounts against live SARS-CoV-2</p> <p><b>Positive antibody response (seroconversion):</b> at least a four-fold increase in post-vaccination titre from baseline</p> <p><b>SAE and Safety</b> &gt; 83%/83%/75% (related to increasing dose, non-significant difference) of patients had at least an AE of moderate/mild severity within the first 7 days (pain, fever, fatigue, headache or muscle pain) &gt; no serious AE were recorded at 28d post vaccination</p> <p><b>Protection</b> Elisa antibodies and neutralizing antibodies increased at D14 and peaked at D28 post-vaccination. Specific T cells response peaked at D14 post-vaccination</p> <p><b>Comments</b> &gt; Older age participants could have a negative effect on the vaccine-elicited responses to SARS-CoV-2 &gt; ADE not evaluated (because of low number of participants) &gt; Pre-existing Ad5 immunity might also have a negative effect on the persistence of the vaccine-elicited immune responses. Issues to evaluate in Phase 2</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Neurology 22MAY2020	<b>Clinical characteristics and outcomes of inpatients with neurologic disease and COVID-19 in Brescia, Lombardy, Italy</b>	Benussi A et al Italy <a href="#">gotopaper</a>	Clinic	<p>Outcomes of patients admitted with neurological disorders with and without COVID-19 173 patients included: <b>56 COVID-19 pos &amp; 117 COVID-19 neg</b> No difference for comorbidities Patients with COVID-19: -Older: 77,0 versus 70,1 years (<math>p=0,006</math>) -More cerebrovascular disorders: 76,8% versus 58,1% (<math>p=0,035</math>) -Higher qSOFA: 0,9 versus 0,5 (<math>p=0,006</math>) -Higher incidence of delirium: 26,8% versus 7,7% (<math>p=0,003</math>) -Higher in-hospital mortality: 75,5% versus 4,3% (<math>p&lt;0,001</math>) -Wider use of high flow oxygenation: 76,8% versus 9,4% (<math>p&lt;0,001</math>) -Prolonged length of stay Potential risk factor of poor prognosis: high qSOFA score – thrombocytopenia – increase lactate deshydrogenase level</p>
Clinical infectious diseases 22MAY2020	<b>Predicting infectious SARS-CoV-2 from diagnostic samples</b>	Bullard, Jared and al. Canada <a href="#">gotopaper</a>	Diagnostic	<p>RT-PCR detects RNA, not infectious virus, thus its ability to determine duration of infectivity of patients is limited.</p> <p>Objective: to determine the relationship between E gene SARS-CoV-2 RT-PCR cycle threshold (Ct) values from respiratory samples, symptom onset to test (STT) and infectivity in cell culture.</p> <p>=&gt; SARS-CoV-2 Vero cell infectivity was only observed for RT-PCR Ct &lt; 24 and STT &lt; 8 days. Infectivity of patients with Ct &gt;24 and duration of symptoms &gt;8 days may be low.</p> <p>CI*: this information can inform public health policy and guide clinical, infection control and occupational health decisions. However, further studies of larger size are needed.</p>
Analytical chemistry 22MAY2020	<b>A novel one-step single-tube nested quantitative Real-Time PCR assay for highly sensitive detection of SARS-CoV-2</b>	Wang, Ji and al. China <a href="#">gotopaper</a>	Diagnostic	<p>qRT-PCR results could be false-negative due to the inadequate sensitivity of qRT-PCR.</p> <p>In this study, we have developed and evaluated a novel one-step single-tube nested quantitative Real-Time PCR (OSN-qRT-PCR) assay for highly sensitive detection of SARS-CoV-2 targeting the ORF1ab and N genes.</p> <p>The sensitivity of the OSN-qRT-PCR assay was 1 copy/reaction and 10-fold higher than that of commercial qRT-PCR kit (10 copies/reaction).</p> <p>CI* : Compared to the qRT-PCR kit, OSN-qRT-PCR assay revealed higher sensitivity and specificity hence better suited to clinical applications for the detection of SARS-CoV-2 in patients with low viral load.</p>
New England Journal of Medicine 21MAY2020	<b>Pulmonary Vascular Endothelialitis, Thrombosis, and Angiogenesis in Covid-19</b>	Ackermann, Maximilian, et al. Germany-Belgium-UK-USA <a href="#">gotopaper</a>	Clinic	<p>Lungs (autopsy) from patients: 7 who died from Covid-19 vs 7 who died from acute respiratory distress syndrome secondary to influenza A (H1N1) infection vs 10 age-matched, uninfected control lungs.</p> <p>Common to both Covid-19 and influenza-associated respiratory failure :</p> <ul style="list-style-type: none"> <li>- diffuse alveolar damage with perivascular T-cell infiltration</li> </ul> <p><b>In Covid-19 patients :</b></p> <ul style="list-style-type: none"> <li>- <b>severe endothelial injury</b> (associated to intracellular virus and disrupted cell membranes)</li> <li>- <b>widespread thrombosis with microangiopathy</b></li> <li>- <b>capillary microthrombi 9x more prevalent</b> compared to influenza patients</li> <li>- <b>new vessel growth 2.7x</b> higher than in lungs from influenza patients</li> </ul> <p>-&gt; <b>Vascular angiogenesis distinguished pulmonary pathobiology of Covid-19 from that of equally severe influenza virus infection</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 21MAY2020	Postmortem Examination of Patients With COVID-19	Schaller, Tina, <i>et al.</i> Germany <a href="#">gotopaper</a>	Clinic	<p>Serial postmortem examinations, 10 patients with proven SARS-CoV-2 infection who died at University Medical Center Augsburg. Median age: 79; range: 64-90 yrs, 7 male; all SARS-CoV-2 +ve by nasopharyngeal swab at hospital admission. Admission median till death: 7.5 days (range, 1-26 days), median of 4 known preexisting comorbidities.</p> <ul style="list-style-type: none"> <li>- <b>At autopsy, SARS-CoV-2 detectable in respiratory tracts of all patients, and PCR +ve in pleural effusion but not in all CSF samples.</b></li> <li>- <b>Predominant histopathologic findings: acute and organizing diffuse alveolar damage and SARS-CoV-2 persistence in the respiratory tract, constituting the leading cause of death in patients with and without invasive ventilation.</b></li> <li>- Periportal liver lymphocyte infiltration considered unspecific inflammation.</li> <li>- Whether myoepicardial alterations represented systemic inflammation or early myocarditis is unclear (criteria for true myocarditis not met).</li> <li>- <b>Central nervous system involvement by COVID-19 could not be detected</b></li> </ul> <p>Limitations: small number of cases from a single center and missing proof of direct viral organ infection</p>
Nature 21MAY2020	Structure of replicating SARS-CoV-2 polymerase	Hillen, Hauke <i>S. et al.</i> Germany <a href="#">gotopaper</a>	Structural biology	<p>Cryo-EM structure of SARS-CoV-2 RdRp in active form, mimicking the replicating enzyme:</p> <ul style="list-style-type: none"> <li>- Active site cleft of nsp12 binds first turn of RNA and mediates RdRp activity with conserved residues.</li> <li>- Two copies of nsp8 bind to opposite sides of the cleft and position the second turn of RNA.</li> <li>- Long helical extensions in nsp8 protrude along exiting RNA, forming +vely charged 'sliding poles'.</li> <li>-&gt; sliding poles can account for known processivity of RdRp required for replicating the long coronavirus genome.</li> <li>- Previous study suggested remdesivir functions as 'immediate' RNA chain terminator, while <b>this study showed that several more nucleotides can be added to RNA following remdesivir incorporation, leading to 'delayed' termination</b> -&gt; mechanism that <b>could explain how remdesivir escapes removal from the RNA 3'-end by the viral exonuclease nsp14.</b></li> </ul>
Cell 21MAY2020	Pathogenesis of SARS-CoV-2 in transgenic mice expressing human angiotensin-converting enzyme 2	Jiang, Ren-Di. <i>et al.</i> China-USA <a href="#">gotopaper</a>	Animal model	<p>Developed a SARS-CoV-2 <b>hACE2 transgenic mouse</b> (HFH4-hACE2 in C3B6 mice) infection model, generating:</p> <ul style="list-style-type: none"> <li>- <b>Typical interstitial pneumonia and pathology, similar COVID-19 patients.</b></li> <li>- Viral quantification: lungs are the major site of infection (<b>viral RNA also found in eye, heart, and brain in some mice</b>).</li> <li>- Full-genome sequences of virus identical to SARS-CoV-2 isolated from the infected lung and brain tissues.</li> <li>- <b>Pre-exposure to SARS-CoV-2 could protect mice from severe pneumonia.</b></li> </ul> <p>-&gt; <b>The hACE2 mouse would be a valuable tool</b> for testing potential vaccines and therapeutics.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 20MAY2020	Viral and host factors related to the clinical outcome of COVID-19	Zhang, Xiaonan, <i>et al.</i> China <a href="#">gotopaper</a>	Clinic	<p>326 confirmed COVID-19 cases in Shanghai. (SARS-CoV-2 genomic sequences assembled from 112 samples + sequences in the Global Initiative on Sharing All Influenza Data (GISAID)).</p> <p>Observations:</p> <ul style="list-style-type: none"> <li>- <b>stable evolution</b> and 2 major lineages suggested with differential exposure history during the early phase of outbreak in Wuhan (similar virulence and clinical outcomes for both lineages).</li> <li>- <b>Lymphocytopenia predictive of disease progression</b> (especially reduced CD4+ and CD8+ T cell counts upon admission).</li> <li>- <b>High IL-6 and IL-8 levels during treatment</b> in patients with severe/critical disease, which correlated with decreased lymphocyte count.</li> </ul> <p>-&gt; <b>Determinants of disease severity seemed to stem mostly from host factors (age, lymphocytopenia and associated cytokine storm), whereas viral genetic variation did not significantly affect outcomes.</b></p>
Eur. Respir. J. MAY2020	A Fully Automatic Deep Learning System for COVID-19 Diagnostic and Prognostic Analysis	Wang et al., China <a href="#">gotopaper</a>	Diagnostic	<p><b>Retrospective collection of 5372 patients with computed tomography images</b> from 7 cities or provinces.</p> <p><u>Steps:</u></p> <p>1st-&gt; 4106 patients with computed tomography images were used to pre-train the DL system, making it learn lung features</p> <p>2<sup>nd</sup>-&gt; 1266 patients from 6 cities or provinces were enrolled to train and externally validate the performance of the deep learning system</p> <p>-&gt; <b>Deep learning system achieved good performance in identifying COVID-19 from other pneumonia (AUC=0.87 and 0.88) and viral pneumonia (AUC=0.86).</b></p> <p>-&gt; <b>Succeeded to stratify patients into high-risk and low-risk groups whose hospital-stay time have significant difference (p=0.013 and 0.014)</b></p> <p>-&gt; <b>Without human-assistance, the deep learning system automatically focused on abnormal areas that showed consistent characteristics with reported radiological findings</b></p>
Science 20MAY2020	SARS-CoV-2 infection protects against rechallenge in rhesus macaques	Chandrashekar A et al. USA <a href="#">gotopaper</a>	Vaccine	<p>Infection of macaques with SARS-CoV-2 results in protective immunity against re-exposure?</p> <p><b>Methods:</b> Infection of 9 rhesus macaques (6-12 years) with SARS-CoV-2 showing high viral loads in the upper and lower respiratory tract, humoral and cellular immune responses and pathologic evidence of viral pneumonia, re-challenged after virus clearance (D35 post initial infection), + 3 naïve animals as positive controls in the rechallenge experiment</p> <p><b>Results</b></p> <ul style="list-style-type: none"> <li>&gt; After primary infection all 9 macaques developed binding antibody responses to S protein and NAb responses and cellular immune responses.</li> <li>&gt; 5 log10 reductions in median viral loads in bronchoalveolar lavage and nasal mucosa compared with primary infection and infected naïve animals</li> <li>&gt; All animals developed anamnestic antibody responses following re-challenge-&gt; protection mediated by immunologic control</li> </ul>
Cell 20MAY2020	Targets of T Cell Responses to SARS-CoV-2 Coronavirus in Humans with COVID-19 Disease and Unexposed Individuals	Grifoni et al., USA <a href="#">gotopaper</a>	Immuno	<p><b>Measuring immunity to SARS-CoV-2 is key for understanding COVID-19 and vaccine development</b></p> <ul style="list-style-type: none"> <li>• Epitope pools detect CD4<sup>+</sup> and CD8<sup>+</sup> T cells in 100% and 70% of convalescent COVID patients</li> <li>• T cell responses are focused not only on spike but also on M, N, and other ORFs</li> <li>• T cell reactivity to SARS-CoV-2 epitopes is also detected in non-exposed individuals</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet Infectious Diseases 20MAY2020	<b>Individual quarantine versus active monitoring of contacts for the mitigation of COVID-19: a modelling study</b>	Peak, C. et al, USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p><b>Aim:</b> to estimate the comparative efficacy of individual quarantine and active monitoring of contacts to control SARS-CoV-2, fitting a model to the incubation period distribution (mean 5.2 days) and to two estimates of the serial interval distribution (4.8 and 7.5 days).</p> <p>Two feasibility settings:</p> <ul style="list-style-type: none"> <li>- high-feasibility (90% of contacts traced, half-day average delay in tracing and symptom recognition, 90% effective isolation)</li> <li>- low-feasibility (50% of contacts traced, 2-day average delay, 50% effective isolation).</li> </ul> <p>→ Mean time of infectiousness onset before symptom onset : 0.77 days (shorter serial interval) and 0.51 days (longer serial interval).</p> <p>→ Individual quarantine in high-feasibility settings (&gt;75% of infected contacts individually quarantined), contains an outbreak of SARS-CoV-2 with a short serial interval 84% of the time. In low-feasibility setting: the outbreak continues to grow and so does the burden of the number of contacts traced for active monitoring or quarantine, particularly asymptomatic contacts.</p> <p><b>Conclusion:</b> When resources are prioritised for scalable interventions such as physical distancing, active monitoring or individual quarantine of high-risk contacts, this can contribute synergistically to mitigation efforts.</p>
Science 20MAY2020	<b>DNA vaccine protection against SARS-CoV-2 in rhesus macaques</b>	Yu et al., USA <a href="#">gotopaper</a>	Vaccine	<p><b>Development of a series of DNA vaccine candidates expressing different forms of the SARS-CoV-2 Spike (S) protein and evaluated them in 35 rhesus macaques.</b></p> <p>-&gt; humoral and cellular immune response with neutralizing antibody (titers = those found in convalescent humans and macaques).</p> <p>-&gt; challenged with SARS-CoV-2: the vaccine encoding the full-length S protein resulted in &gt;3.1 and &gt;3.7 log10 reductions in median viral loads in bronchoalveolar lavage and nasal mucosa.</p> <p>-&gt; Vaccine-elicited neutralizing antibody titers correlated with protective efficacy: <b>suggests an immune correlate of protection</b></p>
JAMA 20MAY2020	<b>Nasal Gene Expression of Angiotensin-Converting Enzyme 2 in Children and Adults</b>	Bunyavanich, Supinda, <i>et al.</i> USA <a href="#">gotopaper</a>	Clinic	<p>Nasal epithelium from 305 individuals aged 4-60, with or without asthma (Mount Sinai Health System, New York, during 2015-2018) collected using a cytology brush.</p> <p><b>ACE2 gene expression in nasal epithelium:</b></p> <ul style="list-style-type: none"> <li>- lowest in aged &lt;10 yrs, and significantly higher in older children (10-17 yrs), young adults (18-24 yrs), and adults (≥25 yrs).</li> <li>- ACE2 gene expression and age was independent of sex and asthma.</li> </ul> <p>-&gt; <b>Lower ACE2 expression in children nasal epithelium relative to adults may help explain why COVID-19 is less prevalent in children.</b></p> <p>Limitation: study did not include individuals older than 60 years.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
NEJM 19MAY2020	Respecting disability rights – towards improved crisis standards of care	Mello M. et al, US <a href="#">gotopaper</a>	HSS/Politic	<p>Policymakers and hospitals can take key steps to honor commitments to antidiscrimination principles while appropriately stewarding scarce resources during a public health emergency.</p> <p><b>1- Do not use categorical exclusions.</b> Patients must not be categorically excluded from access to treatment because of a disability or diagnosis. Consider not whether someone has a disability but patient's prospects of benefiting from treatment.</p> <p><b>2- Do not use perceived quality of life</b> = biases in how the public/physicians evaluate the quality of life of persons with disabilities.</p> <p><b>3- Use hospital survival and near-term prognosis but not long-term life expectancy.</b> Predictions of long-term life expectancy are much more uncertain + prone to bias than predictions of short-term survival + affected by social circumstances (poverty). Ignoring near-term prognosis can produce outcomes inconsistent with responsible stewardship of scarce resources.</p> <p><b>4- Designate triage officers as the decision-makers and train them to respect disability rights.</b> <b>Include disability rights advocates in policy development and dissemination</b> = it will also help in avoiding inflammatory language and ensure public understanding for operationalization.</p>
Clinical Infectious Diseases 19MAY2020	Early Short Course Corticosteroids in Hospitalized Patients with COVID-19	Fadel, Raef et al. USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Multi-center quasi-experimental study</b> of 213 adult patients with confirmed moderate to severe COVID, 81 (38%) and 132 (62%) in SOC and <b>early short course corticosteroid</b> (methylprednisolone) groups, respectively. Outcomes were evaluated with a primary composite endpoint of escalation of care from ward to ICU, new requirement for mechanical ventilation, and mortality.</p> <p>- The composite endpoint occurred at a significantly lower rate in the early corticosteroid group (34.9% vs. 54.3%, <math>p=0.005</math>). Treatment effect observed within each individual component of the composite endpoint.</p> <p>- Significant reduction in median hospital length of stay was also observed in the early corticosteroid group (8 vs. 5 days, <math>p &lt; 0.001</math>).</p> <p>- Multivariate regression analysis demonstrated an independent reduction in the composite endpoint at 14-days controlling for other factors (aOR: 0.41; 95% CI [0.22 – 0.77]).</p> <p><b>Conclusion: An early short course of methylprednisolone in patients with moderate to severe COVID-19 reduced escalation of care and improved clinical outcomes.</b></p> <p><b>Limitations:</b> pragmatic quasi-experimental design was used and there are some differences in the baseline characteristics of the comparator groups ; limited follow up period of 14 days.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
J Infect Dis 19MAY2020	<b>Influence of storage conditions on SARS-CoV-2 nucleic acid detection in throat swabs</b>	Li, Lin and al. China <a href="#">gotopaper</a>	Virology	<p>For the detection of SARS-CoV-2 infection, samples often need to be shipped or inactivated before SARS-CoV-2 testing. In this study, we checked the influence of sample storage conditions on SARS-CoV-2 nucleic acid testing results, including sample inactivation time, storage temperature and the time.</p> <p>All of these conditions caused an increase in the Ct values of the nucleic acid tests and led to the misclassification of at least 10.2% of positive cases as negative or suspected.</p> <p>CL<sup>o</sup>: results highlight the importance of immediate testing of samples for SARS-CoV-2 nucleic acid and detection.</p>
Lancet 19MAY2020	<b>Epidemiology, clinical course, and outcomes of critically ill adults with COVID-19 in New York City: a prospective cohort study</b>	Cummings MJ et al USA <a href="#">gotopaper</a>	Clinic	<p>2 hospitals in NY – critically ill patient with COVID-19 – At least 28 days of observation</p> <p>1150 adults admitted COVID-19 which <b>257 (22%) were critically ill</b> (included) Median age: 62 years – 67% male – 82% at least one comorbidity (HTA, diabetes) 46% had obesity</p> <p><u>Treatment</u></p> <ul style="list-style-type: none"> <li>- 72% received hydroxychloroquine and 9% remdesivir</li> <li>- 26% received corticosteroid</li> <li>- 66% received vasopressor</li> <li>- 31% received RRT</li> </ul> <p><u>Outcomes:</u></p> <ul style="list-style-type: none"> <li>- 79% received IMV median of 18 days</li> <li>- <b>39% died (median of 9 days in the hospital)</b> and 37% remained hospitalized</li> </ul> <p><u>Association with in hospital death</u> (significantly)</p> <ul style="list-style-type: none"> <li>- Older age aHR: 1,31 [1,09 – 1,57] per 10 years increase</li> <li>- Chronic cardiac disease aHR: 1,76 [1,08 – 2,86]</li> <li>- Chronic pulmonary disease aHR: 2,94 [1,48 – 5,84]</li> <li>- Concentration of IL-6 aHR: 1,11 [1,02 – 1,20] per decile increase</li> <li>- Concentration of D-dimer aHR: 1,10 [1,01 – 1,19] per decile increase</li> </ul> <p>→ <b>high frequency of IMV &amp; in hospital mortality</b></p>
Nature Medicine 19MAY2020	<b>Artificial intelligence-enabled rapid diagnosis of patients with COVID-19</b>	Mei et al., USA <a href="#">gotopaper</a>	Diagnostic	<p>Use of artificial intelligence (AI) algorithms to integrate chest CT findings with clinical symptoms, exposure history and laboratory testing to rapidly diagnose patients who are positive for COVID-19.</p> <p>-&gt; In a test set of 279 patients, the AI system achieved an area under the curve of 0.92 and had equal sensitivity as compared to a senior thoracic radiologist.</p> <p>-&gt; The AI system also improved the detection of patients who were positive for COVID-19 via RT-PCR who presented with normal CT scans, correctly identifying 17 of 25 (68%) patients, whereas radiologists classified all of these patients as COVID-19 negative</p> <p><b>When CT scans and associated clinical history are available, the proposed AI system can help to rapidly diagnose COVID-19 patients.</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Circulation 19MAY2020	<b>Deep Vein Thrombosis in Hospitalized Patients with Coronavirus Disease 2019 (COVID-19) in Wuhan, China: Prevalence, Risk Factors, and Outcome</b>	Zang L et al, China <a href="#">gotopaper</a>	Clinic	<p>Investigation of deep vein thrombosis (DVT) in hospitalized patient with COVID-19 143 patients from Jan 29 and Feb 29 <u>Demographic</u>: 51,7% man – median age= 63 46,1% (66) lower extremity DVT= 23 proximal DVT and 43 distal DVT <u>DVT vs no DVT</u>:</p> <ul style="list-style-type: none"> <li>- Older</li> <li>- Lower oxygenation index</li> <li>- Higher rate of cardiac injury</li> <li>- Increase death (23 vs 9, p=0,001)</li> </ul> <p><u>Multivariate analysis, DVT associated with</u></p> <ul style="list-style-type: none"> <li>- CURB-65 score3-5, OR:6,12</li> <li>- Padua prediction <math>\geq 4</math>, OR: 4,01</li> <li>- D-dimer&gt;1<math>\mu</math>g/ml OR :5,81</li> </ul> <p><u>Predicting DVT</u> <b>CURB-65 score3-5, Padua prediction score <math>\geq 4</math> + D-dimer&gt;1 had Se 88% and Spe 61,4%</b></p> <p>→ prevalence of DVT is high → importance of prophylaxis for venous thromboembolism (Padua prediction score <math>\geq 4</math>)</p>
Science 18MAY2020	<b>Susceptible supply limits the role of climate in the early SARS-CoV-2 pandemic</b>	Baker, Rachel E. et al. USA <a href="#">gotopaper</a>	Climate	<p>Climate-dependent model to simulate SARS-CoV-2 pandemic, probing different scenarios based on known coronavirus biology.</p> <p>Results suggest:</p> <ul style="list-style-type: none"> <li>• <b>While climate may play a role in details of the size and timescales of an endemic outbreak, population immunity is a much more fundamental driver of pandemic invasion dynamics.</b></li> <li>• <b>Both tropical and temperate locations should prepare for severe outbreaks and summertime temperatures will not effectively limit spread of infection.</b></li> <li>• <b>Endemic cycles will likely be tied to climate factors and seasonal peaks may vary with latitude.</b></li> </ul>
Cell 18MAY2020	<b>Potent neutralizing antibodies against SARS-CoV-2 identified by high-throughput single-cell sequencing of convalescent patients' B cells</b>	Yunlong Cao et al., Chine, <a href="#">gotopaper</a>	Immunology	<p>Rapid and efficient identification of SARS-CoV-2 neutralizing antibodies achieved by high-throughput single-cell RNA and VDJ sequencing of antigen-binding B cells from 60 convalescent COVID-19 patients reveal over 8,500 antigen-binding B cell clonotypes expressing IgG1 antibodies.</p> <p>=&gt; among of which,14 potent neutralizing mAbs were identified</p> <p>=&gt; one of them, BD-368-2, exhibited an IC50 of 1.2 ng/mL and 15 ng/mL against pseudotyped and authentic SARS-CoV-2.</p> <p>=&gt; in vivo experiments confirmed that BD-368-2 provide strong therapeutic efficacy and prophylactic protection against SARS-CoV-2, using the hACE2 transgenic mice model</p> <p>Conclusion: The potent neutralizing antibodies we identified may provide an effective therapeutic and prophylactic solution</p> <p>Limitation: deeper sequencing of the scRNA libraries is needed to further evaluate the effectiveness of removing exhausted memory B cells to improve the identification of neutralizing mAbs.</p>
Ann. Intern. Med. 18MAY2020	<b>Tocilizumab for Hemophagocytic Syndrome in a Kidney Transplant Recipient With COVID-19</b>	Faguer, Stanislas et al. France, <a href="#">gotopaper</a>	Therapeutic	<p><b>Case Report</b> describing an <b>immunocompromised patient with COVID-19 and a related hemophagocytic syndrome</b> who was treated with <b>tocilizumab</b>.</p> <p>The cytokine storm and multiorgan failure <b>rapidly reversed</b>, and the patient made a <b>speedy recovery</b>. On hospital day 30, the patient was breathing spontaneously with protective tracheotomy, and rehabilitation is ongoing.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet Diabetes & Endocrinology 18MAY2020	Prevalence of obesity among adult inpatients with COVID-19 in France	Caussy, Cyrielle et al. France <a href="#">gotopaper</a>	Population studies	<p><b>Lyon University Hospital admissions</b> with BMI values: 340 patients with confirmed severe COVID-19 (68% non-critical and 32% critical COVID-19 patients) vs 1210 retrospective non-COVID-19 ICU patients admitted each year between 2007 - 2019.</p> <p>Results :</p> <ul style="list-style-type: none"> <li>- 25% of severe COVID-19 patients had obesity, vs 15-3% in French adult population in 2014.</li> <li>-&gt; <b>Obesity prevalence 1.35 times higher in severe COVID-19 patients vs general French population.</b></li> <li>-&gt; <b>In ICU, obesity prevalence 1.89 times higher vs general population.</b></li> <li>-&gt; <b>Obesity prevalence higher in critical vs non-critical COVID-19 patients.</b></li> </ul> <p>In agreement with preliminary data from 124 patients with critical COVID-19 and 306 ICU patients without COVID-19 from <b>Lille University Hospital (Simonnet A <i>et al.</i> 2020)</b> :</p> <ul style="list-style-type: none"> <li>-&gt; <b>obesity prevalence 2.88 times higher in critical COVID-19 vs French general population</b></li> <li>-&gt; <b>Obesity prevalence higher in critical COVID-19 vs Lille ICU patients without COVID-19 (n=306).</b></li> </ul> <p><b>=&gt; Report significant association in obesity prevalence and severe COVID-19, including critical COVID-19, and suggests obesity =&gt; risk factor of pejorative evolution of COVID-19, increasing risk of ICU admission.</b></p>
Circulation 17MAY2020	Acute heart failure in multisystem inflammatory syndrome in children (MIS-C) in the context of global SARS-CoV-2 pandemic	Belhadjer S et al, France <a href="#">gotopaper</a>	Clinic	<p>A series of children admitted to PICU for cardiogenic shock + left ventricular dysfunction + severe inflammatory state (14 centers)</p> <p>35 children – median age: 10 y [2 – 16]</p> <p>Comorbidities: 28% of the children which 17% were overweight</p> <p><u>Symptoms:</u> Fever and asthenia (100%) / Gastrointestinal symptoms (83%) / Respiratory distress (65%) – rhinorrhea (43%)</p> <p><u>Left ventricular ejection at baseline</u> &lt; 30 for 28% / 30 to 50 for 72%</p> <p><u>Laboratory:</u> cytokine storm → high IL-6 and D-dimer (= macrophage activation) + elevation of CRP and PCT 88% were positive for SARS-CoV-2 (nasopharyngeal swabs or serology)</p> <p><u>Treatment:</u> 94% Respiratory support: invasive (62%) – noninvasive (32%) / 28% ECMO / 80% inotropic support / 100% IV globuline / 1/3 received steroid therapy</p> <p><u>At discharge:</u> 25/35 had left ventricular function restored - no death</p> <p><b>→ SARS-COV-2 + severe inflammatory state in children → acute cardiac decompensation</b></p>
The Journal of Antimicrobial Chemotherapy 17MAY2020	COVID-19 infection also occurs in patients taking hydroxychloroquine	Lahouati, M et al. France <a href="#">gotopaper</a>	Therapeutic	<p>Report on <b>two severe cases of COVID-19 in patients already using hydroxychloroquine for a long time</b> to treat inflammatory disease.</p> <p><b>High plasma levels of hydroxychloroquine</b> collected on admission in those cases confirm chronic exposure and adherence to hydroxychloroquine. These values are <b>close to or higher than the EC50</b> described by Yao et al. not taking into account lung diffusion. Those potentially immunosuppressed patients do not represent the general population exposed to COVID-19.</p> <p>It cannot be excluded that chloroquine and hydroxychloroquine negatively impact the early inflammatory response to the virus and the risk of acquisition of infection owing to their anti-inflammatory activity.</p> <p>These <b>observational data are not in favour of a universal protective effect of hydroxychloroquine</b>, and clinicians should use it carefully, awaiting the results of clinical trials, particularly in the context of prevention.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Water research 16MAY2020	<b>SARS-CoV-2 RNA in wastewater anticipated COVID-19 occurrence in a low prevalence area</b>	Randazzo et al, Spain <a href="#">gotopaper</a>	Public Health/Epidemiology	<p><b>Faecal shedding of SARS-CoV-2 RNA from COVID-19 patients has extensively been reported.</b></p> <p>We investigated the occurrence of SARS-CoV-2 RNA in six wastewater treatment plants serving the major municipalities within the Region of Murcia, Spain (low COVID-19 prevalence).</p> <p>-&gt; The estimated quantification of SARS-CoV-2 RNA titers in untreated wastewater waters of <math>5.4 \pm 0.2 \log_{10}</math> genomic copies/L on average.</p> <p>-&gt; Two secondary water samples resulted positive (2/18) and all tertiary water samples tested as negative (0/12). SARS-CoV-2 RNA shedding in stools was present even before the first cases were reported by the authorities.</p> <p><b>This strategy could be implemented in environmental surveillance as an early indicator of the infection within a specific population.</b></p>
Developmental Cell 16MAY2020	<b>Cigarette smoke exposure and inflammatory signaling increase the expression of the SARS-CoV-2 receptor ACE2 in the respiratory tract</b>	Smith, Joan C. <i>et al.</i> USA <a href="#">gotopaper</a>	Fundamental research	<p>Cigarette smoke causes dose-dependent upregulation of ACE2 receptor in rodent and human lungs.</p> <p>Single-cell sequencing data:</p> <ul style="list-style-type: none"> <li>- <b>ACE2 expressed in a subset of secretory cells in respiratory tract.</b></li> <li>- <b>Chronic smoke exposure triggers expansion of this cell population and increased ACE2 expression.</b></li> <li>- <b>Quitting smoking has convers effect</b> (decreases this cell population and ACE2 levels).</li> <li>- ACE2 expression responsive to inflammatory signalling and upregulated by viral infections / interferon treatment.</li> </ul> <p>-&gt; May partially explain why smokers are particularly susceptible to severe SARS-CoV-2 infections</p> <p>-&gt; <b>identifies ACE2 as interferon-stimulated gene in lung cells -&gt; possible positive-feedback loops increasing ACE2 levels and facilitating viral dissemination.</b></p>
J. Clin. Virol. 16MAY 2020	<b>A combined oropharyngeal/nares swab is a suitable alternative to nasopharyngeal swabs for the detection of SARS-CoV-2</b>	LeBlanc, Jason J. and al., Canada <a href="#">gotopaper</a>	Diagnostic	<p>Given the global shortage of nasopharyngeal (NP) swabs typically used for respiratory virus detection, alternative collection methods were evaluated during the COVID-19 pandemic.</p> <p>This study showed that a combined oropharyngeal/nares swab is a suitable alternative to NP swabs for the detection of SARS-CoV-2, with sensitivities of 91.7% and 94.4%, respectively.</p>
The Lancet Infectious Diseases 15MAY2020	<b>Risk factors for SARS-CoV-2 among patients in the Oxford Royal College of General Practitioners Research and Surveillance Centre primary care network: a cross-sectional study</b>	Lusignan, Simon de et al. UK-South Africa <a href="#">gotopaper</a>	Clinic	<p>Routinely collected, pseudonymised data for patients in the RCGP Research and Surveillance Centre primary care network +ve for SARS-CoV-2 (Jan 28 - April 4 2020):</p> <ul style="list-style-type: none"> <li>- 587 SARS-CoV-2 +ve out of 3802 test results.</li> <li>- male sex independently associated with testing +ve (18 vs 13 % for women)</li> </ul> <p><b>Clinical factors and demographics more likely to testing +ve :</b></p> <ul style="list-style-type: none"> <li>- Adults, in particular <b>ages 40-64</b>, vs children (19 % in aged 40-64 vs 5 % in children)</li> <li>- People with <b>chronic kidney disease</b> (33 vs 14 % without chronic kidney disease), but no significant association with other chronic conditions.</li> <li>- <b>Obese people</b> (21 vs 13 % for people of normal weight)</li> <li>- <b>Active smoking at decreased odds of testing +ve</b> (11 vs 18 % in non-smokers)</li> <li>- <b>black people</b> vs white (62 vs 16 %)</li> <li>- <b>People living in urban areas</b> vs rural areas (26 vs 5-6% in rural areas)</li> <li>- <b>People living in deprived areas</b> (30 vs 8% in least deprived areas)</li> </ul> <p>-&gt; +ve SARS-CoV-2 test results in primary care cohort was associated with similar risk factors as for severe outcomes of COVID-19 in hospital settings, except for smoking.</p> <p>-&gt; Provides evidence of potential sociodemographic factors associated with a +ve test : deprivation, population density, ethnicity and chronic kidney disease.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Gastroenterology 15MAY2020	<b>Associations between Angiotensin Converting Enzyme Inhibitors and Angiotensin II Receptor Blocker Use, Gastrointestinal Symptoms, and Mortality among Patients with COVID-19</b>	Tan, Nian-Di et al. China <a href="#">gotopaper</a>	Therapeutic/ Clinic	<p><b>Retrospective cohort study</b> of consecutive patients with COVID-19. Among the <b>100 participants with hypertension</b>, 31 were classified as ACEI/ARB group and the remaining 69 were classified as non-ACEI/ARB group.</p> <p><b>Inpatient treatment with ACEI/ARB</b> was associated with <b>lower risk of digestive system involvement</b> and <b>lower risk of all-cause mortality</b> compared with ACEI/ARB non-users in COVID-19 patients with hypertension.</p> <p><b>Limitations:</b> small sample-size, possible unappreciated confounding effect.</p>
Science 15MAY2020	<b>Serology assays to manage COVID-19</b>	Krammer F., Simon V., USA <a href="#">gotopaper</a>	Public Health/Epidemiol	<p>Measurement of antibodies to SARS-CoV-2 will improve disease management if used correctly.</p> <p>This perspective article describes the serological assays available and discusses the potential applications, including:</p> <ul style="list-style-type: none"> <li>- understand the antibody responses mounted upon SARS-CoV-2 infection and vaccination;</li> <li>- inform on the prevalence of SARS-CoV-2 infection if different populations;</li> <li>- identification of donors for convalescent plasma therapy;</li> <li>- identify individuals who are immune (and the caveats concerning this point).</li> </ul> <p><b>With high-quality serological assays now available, the key challenge will be to apply and deploy these tests in a strategic manner.</b></p>
Clinical microbiology and infection 15MAY2020	<b>A multiple center clinical evaluation of an ultra-fast single-tube assay for SARS-CoV-2 RNA</b>	Wang, Ji and al. China <a href="#">gotopaper</a>	Diagnostic	<p>To evaluate the performance of an ultra-fast single-tube nucleic acid isothermal amplification detection assay for SARS-CoV-2 RNA using clinical samples from multiple centers.</p> <p>A reverse transcription recombinase-aided amplification (RT-RAA) assay for SARS-CoV-2 was conducted within 15 minutes at 39°C with portable instruments after addition of extracted RNA. The clinical performance of RT-RAA assay was evaluated using 947 clinical samples and the approved commercial real-time fluorescent RT-PCR (qRT-PCR) kits were used for parallel detection. The sensitivity and specificity of RT-RAA were compared and analyzed.</p> <p>CI<sup>o</sup>: with comparable sensitivity and specificity to the commercial qRT-PCR kits, RT-RAA assay for SARS-CoV-2 exhibited distinctive advantages of simplicity and rapidity in terms of operation and turn-around time.</p>
Science 15MAY2020	<b>Inferring change points in the spread of COVID-19 reveals the effectiveness of interventions</b>	Dehning et al., Germany <a href="#">gotopaper</a>	PublicHealth/Epidemiology	<p><b>By combining an established epidemiological model with Bayesian inference -&gt; analysis of the time dependence of the effective growth rate of new infections</b></p> <p>Focusing on COVID-19 spread in Germany, detection of change points in the effective growth rate that correlate well with the times of publicly announced interventions -&gt; Possibility to quantify the effect of interventions, and we can incorporate the corresponding change points into forecasts of future scenarios and case numbers.</p> <p>This code is freely available and can be readily adapted to any country or region.</p>
Nature 14MAY2020	<b>Proteomics of SARS-CoV-2-infected host cells reveals therapy targets</b>	Bojkova, Denisa et al. Germany <a href="#">gotopaper</a>	Therapeutic	<p><b>Identification of the host cell pathways</b> modulated by SARS-CoV-2 infection and inhibition of these pathways showed to prevent viral replication in human cells. A <b>human cell culture model for infection</b> with SARS-CoV-2 clinical isolate was established. Employing this system, the <b>SARS-CoV-2 infection profile</b> was determined by translato3 and proteome proteomics at different times after infection. These analyses revealed that <b>SARS-CoV-2 reshapes central cellular pathways, such as translation, splicing, carbon metabolism and nucleic acid metabolism. Small molecule inhibitors targeting these pathways prevented viral replication in cells.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Lancet 14MAY2020	Use of renin–angiotensin–aldosterone system inhibitors and risk of COVID-19 requiring admission to hospital: a case-population study	Abajo, Francisco J. De, et al. Spain <a href="#">gotopaper</a>	Therapeutic/ Clinic	<p><b>Case-population study</b> of consecutively selected hospitalized patients with a PCR-confirmed diagnosis of COVID-19, and randomly sampled ten patients per case, individually matched for age, sex, region, and date of admission to hospital as a reference group. <b>1139 cases</b> and <b>11 390 population controls</b>. Despite being matched on sex and age, a significantly higher proportion of cases had pre-existing cardiovascular disease (OR 1.98, 95% CI 1.62–2.41) and risk factors (1.46, 1.23–1.73) than did controls.</p> <p>The <b>current use of RAAS inhibitors is not associated with an increased risk of COVID-19 requiring admission to hospital</b> (including fatal cases and those admitted to an ICU) compared with other antihypertensive drugs. No substantial difference was observed between ACE inhibitors and angiotensin-receptor blockers, nor among short-term and long-term users. Sex, age, and background cardiovascular risk did not significantly affect the results, although use of RAAS inhibitors was associated with a reduced risk of COVID-19 requiring admission to hospital in patients with diabetes.</p> <p><b>Limitations:</b> different data sources to extract information from cases and controls ; cases and controls were recorded at different dates ; data on smoking and other lifestyle habits not collected; observational study, residual confounding due to unmeasured or unknown confounders cannot be ruled out.</p>
BMJ 14MAY2020	Hydroxychloroquine in patients with mainly mild to moderate coronavirus disease 2019: open label, randomised controlled trial	Tang, Wei et al. China <a href="#">gotopaper</a>	Therapeutic	<p><b>Multicentre, randomised, parallel, open label trial of hydroxychloroquine</b> (1200 mg daily for three days, then 800 mg daily) versus standard of care in 150 patients admitted to hospital with covid-19.</p> <p><b>No evidence to support an increase in the probability of negative conversion</b> of SARS-CoV-2 conferred by the addition of hydroxychloroquine administration to the current standard of care in patients admitted to hospital with mainly <b>persistent mild to moderate covid-19</b>. <b>Adverse events</b>, particularly gastrointestinal events, were <b>more frequently reported</b> in patients receiving <b>hydroxychloroquine</b>. <b>Limitations:</b> open label ; use of sequential envelopes for randomisation ; no patients at the early stage of disease ; 148/150 (99%) patients had mild to moderate disease ; underpowered sample size due to the lack of enough eligible patients to enrol.</p>
American journal of obstetrics and gynecology 14MAY2020	Clinical Characteristics of 46 Pregnant Women with a SARS-CoV-2 Infection in Washington State	Lokken EM et al, USA <a href="#">gotopaper</a>	Clinic	<p>Description of maternal disease and obstetrical outcomes – 6 hospital in Washington state <b>46 pregnant women</b> <u>Demographic:</u> median age: 29y – 6,5% in first &amp; 43,4% second &amp; 50% in third trimester 26,1% had at least one comorbidity: asthma – hypertension – diabetes - .... <b>28,6% were overweight and 35,7% obese</b> <u>Symptoms:</u> 93,5% had one – cough (70%) – fever (51%) – dyspnea – headache .... <u>Outcomes:</u></p> <ul style="list-style-type: none"> <li>- 15% categorized has severe disease: overweight/obese/comorbidity</li> <li>- 16% hospitalized &amp; 1 to ICU (all severe disease)</li> <li>- 8 delivered during the study period (median: 38,4 week)</li> <li>- 1 stillbirth: unknown etiology</li> </ul> <p>→ <b>higher risk group: chronic comorbidity / obese / overweight</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Clinica Chimica Acta 14MAY2020	<b>The underlying changes and predicting role of peripheral blood inflammatory cells in severe COVID-19 patients: a sentinel?</b>	Sun, Da-wei and al. China <a href="#">gotopaper</a>	Virology	<p>The underlying changes of peripheral blood inflammatory cells (PBICs) in COVID-19 patients are little known. Moreover, the risk factors for the underlying changes of PBICs and their predicting role in severe COVID-19 patients remain uncertain.</p> <p>This retrospective study includes two cohorts: the main cohort enrolling 45 patients of severe type serving as study group, and the secondary cohort enrolling 12 patients of no-severe type serving as control group. The PBICs analysis was based on blood routine and lymphocyte subsets. The inflammatory cell levels were compared among patients according to clinical classifications, disease-associated phases, as well as one-month outcomes.</p> <p>Results: the patients of severe type suffered from significantly decreased counts of lymphocytes, eosinophils, basophils, but increased counts of neutrophils. These PBICs alterations got improved in recovery phase, but persisted or got worse in aggravated phase.</p> <p>CI<sup>o</sup>: lymphopenia and eosinopenia may serve as predictors of disease severity and disease progression in COVID-19 patients, and enhancing the cellular immunity may contribute to COVID-19 treatment. Thus, PBICs might become a sentinel of COVID-19, and it deserves attention during COVID-19 treatment</p>
Journal of Allergy and Clinical Immunology 14MAY2020	<b>Complement activation in patients with COVID-19: a novel therapeutic target</b>	Cugno, Massimo et al. Italy <a href="#">gotopaper</a>	Therapeutic	<p>Preliminary data providing evidence of <b>complement activation</b> in patients with COVID-19 with <b>different degrees of respiratory failure</b>. Investigation of the <b>plasma levels of sC5b-9 and C5a</b> as markers of complement activation in 31 COVID-19 patients, compared with 27 healthy subjects.</p> <ul style="list-style-type: none"> <li>- <b>Plasma levels of sC5b-9</b> were <b>significantly higher</b> in the patients with moderate disease and those with severe disease <b>than in the healthy controls</b>, and <b>significantly higher</b> in the patients with <b>severe disease</b> than in those with moderate disease.</li> <li>- The <b>plasma levels of C5a</b> were <b>higher</b> in the patients with moderate disease and those with severe disease <b>than in the healthy</b> (P=0.0001 for both), with no statistically significant difference between the two patient groups.</li> <li>- The cohort of patients had <b>increased levels of acute-phase proteins and coagulation system abnormalities</b>.</li> </ul> <p>Complement activation may contribute to the development of lung and endothelial damage in patients. Possibility that the coronavirus may directly cause damage to endothelial cells.</p>
BMJ 14MAY2020	<b>Clinical efficacy of hydroxychloroquine in patients with covid-19 pneumonia who require oxygen: observational comparative study using routine care data</b>	Mahévas, Matthieu et al. France <a href="#">gotopaper</a>	Therapeutic	<p><b>Comparative study</b> that uses <b>real world data</b> collected from routine care to assess the <b>efficacy and safety of hydroxychloroquine</b> in a population of <b>181 patients</b> admitted to hospital with covid-19 hypoxaemic pneumonia.</p> <ul style="list-style-type: none"> <li>- Hydroxychloroquine treatment at 600 mg/day added to standard care was <b>not associated with a reduction of admissions to the intensive care unit or death</b> 21 days after hospital admission compared with standard care alone.</li> <li>- The <b>rate of survival without acute respiratory distress syndrome did not increase</b>.</li> <li>- Eight patients in the treatment group (10%) experienced <b>electrocardiographic modifications that required discontinuation of treatment</b>.</li> </ul> <p>The results of this study do not support its use in patients admitted to hospital with covid-19 who require oxygen. <b>Limitations:</b> observational data, centre effect not taken into account ; limited sample ; only patients admitted to hospital.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Clinical Infectious Disease 14MAY2020	<b>Risk Factors of Severe Disease and Efficacy of Treatment in Patients Infected with COVID-19: A Systematic Review, Meta-Analysis and Meta-Regression Analysis</b>	Zhang, John J. Y et al. Singapore <a href="#">gotopaper</a>	Therapeutic/ Clinic	<p><b>Systematic review and meta-analysis</b> on COVID-19 clinical features and/or treatment outcomes.</p> <p><b>45 studies</b> reporting <b>4203 patients</b> were included. Pooled rates of intensive care unit (ICU) admission, mortality and acute respiratory distress syndrome (ARDS) were 10.9%, 4.3% and 18.4%, respectively.</p> <ul style="list-style-type: none"> <li>- On meta-regression, <b>ICU admission</b> was <b>predicted by raised leukocyte count</b> (<math>p&lt;0.0001</math>), <b>raised alanine aminotransferase</b> (<math>p=0.024</math>), <b>raised aspartate transaminase</b> (<math>p=0.0040</math>), <b>elevated lactate dehydrogenase (LDH)</b> (<math>p&lt;0.0001</math>) and <b>increased procalcitonin</b> (<math>p&lt;0.0001</math>).</li> <li>- <b>ARDS</b> was <b>predicted by elevated LDH</b> (<math>p&lt;0.0001</math>), while <b>mortality</b> was <b>predicted by raised leukocyte count</b> (<math>p=0.0005</math>) and <b>elevated LDH</b> (<math>p&lt;0.0001</math>).</li> <li>- Treatment with <b>lopinavir-ritonavir</b> showed <b>no significant benefit in mortality and ARDS rates</b>. <b>Corticosteroids</b> were associated with a <b>higher rate of ARDS</b> (<math>p=0.0003</math>).</li> </ul> <p><b>Limitations:</b> possible selection bias (publications in English) ; only studies from Asia at the time of the literature search ; studies included were observational ; heterogeneity in the range of symptoms and comorbidities recorded in the different studies.</p>
Nature 14MAY2020	<b>Infection of dogs with SARS-CoV-2</b>	Sit, Thomas H. C Hong Kong <a href="#">gotopaper</a>	Fundamental research	<p><b>2/15 dogs from households with confirmed human COVID-19 cases</b> in Hong Kong were found to be infected (qRT-PCR, serology, viral genome sequencing, and virus isolation in 1 dog):</p> <ul style="list-style-type: none"> <li>- a 17yr-old male Pomeranian (SARS-CoV-2 RNA detected from 5 nasal swabs over 13-days).</li> <li>- a 2.5yr-old male German Shepherd dog (SARS-CoV-2 RNA on two occasions and virus isolated from nasal and oral swabs)</li> <li>- Both had antibody responses (plaque reduction neutralization assays).</li> <li>- <b>Viral genetic sequences from both dogs were identical to virus detected in respective human cases. Animals asymptomatic during quarantine.</b></li> </ul> <p><b>-&gt; These are instances of human-to-animal transmission of SARS-CoV-2.</b> Unclear whether infected dogs can transmit the virus to other animals or back to humans.</p>
Nature 14MAY2020	<b>Pathogenesis and transmission of SARS-CoV-2 in golden hamsters</b>	Sia, Sin Fun et al. Hong Kong <a href="#">gotopaper</a>	Animal model	<p>Pathogenesis and transmissibility of the SARS-CoV-2 in <b>golden Syrian hamsters</b> (intranasal infection):</p> <ul style="list-style-type: none"> <li>- viral antigens (immunohistochemistry) in nasal mucosa, bronchial epithelial cells and in areas of lung on 2 and 5 dpi, followed by rapid viral clearance and pneumocyte hyperplasia on 7 dpi.</li> <li>- Viral antigen found in duodenum epithelial cells and viral RNA in feces.</li> </ul> <p><b>-&gt; Efficient SARS-CoV-2 transmission from inoculated hamsters to naïve by direct contact and via aerosols.</b> Transmission via fomites less efficient.</p> <p><b>-&gt; Communicable period was short and correlated with detection of infectious virus but not viral RNA.</b></p> <p><b>-&gt; Inoculated and naturally-infected hamsters showed apparent weight loss, and all animals recovered with detection of neutralizing antibodies.</b></p>
Proc. Natl. Acad. Sci. U. S. A. 13MAY2020	<b>The airborne lifetime of small speech droplets and their potential importance in SARS-CoV-2 transmission</b>	Stadnytskyi et al. USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Speech droplets generated by asymptomatic carriers of SARS-CoV-2 likely to be a mode of disease transmission.</p> <p><b>-&gt; Highly sensitive laser light scattering observations have revealed that loud speech can emit thousands of oral fluid droplets per second.</b></p> <p><b>-&gt; In a closed, stagnant air environment, they disappear from the window of view with time constants in the range of 8 to 14 min (droplet nuclei of ca. 4 µm diameter, or 12- to 21-µm droplets prior to dehydration).</b></p> <p><b>There is therefore a substantial probability that normal speaking causes airborne virus transmission in confined environments.</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Science 13MAY2020	<b>Estimating the burden of SARS-CoV-2 in France</b>	Salje et al., France <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Using models applied to hospital and death data, we estimate the impact of the lockdown and current population immunity in France.</p> <p>-&gt; We find 3.6% of infected individuals are hospitalized and 0.7% die (0.001% in &lt;20 years of age (ya), 10.1% in &gt;80ya). The lockdown reduced the reproductive number from 2.90 to 0.67 (77% reduction). By 11 May 2020, when interventions are scheduled to be eased, we project that 4.4% (range: 2.8–7.2) of the population will have been infected.</p> <p>Population immunity appears insufficient to avoid a second wave if all control measures are released at the end of the lockdown</p>
Ann. Intern. Med. 13MAY2020	<b>Variation in False-Negative Rate of Reverse Transcriptase Polymerase Chain Reaction–Based SARS-CoV-2 Tests by Time Since Exposure</b>	Kucirka, Lauren M. and al. USA <a href="#">gotopaper</a>	Virology	<p>Tests for SARS-CoV-2 based on RT-PCR are being used to “rule out” infection among high-risk persons, such as exposed inpatients and health care workers. It is critical to understand how the predictive value of the test varies with time from exposure and symptom onset to avoid being falsely reassured by negative test results.</p> <p>Objective: To estimate the false-negative rate by day since infection.</p> <p>CI<sup>o</sup>: care must be taken in interpreting RT-PCR tests for SARS-CoV-2 infection—particularly early in the course of infection—when using these results as a basis for removing precautions intended to prevent onward transmission. If clinical suspicion is high, infection should not be ruled out on the basis of RT-PCR alone, and the clinical and epidemiologic situation should be carefully considered.</p>
N. Engl. J. Med. 13MAY2020	<b>Multiorgan and Renal Tropism of SARS-CoV-2</b>	Puelles, Victor G. et al Germany <a href="#">gotopaper</a>	Cellular tropism	<p><b>Autopsy series from 22 patients who died from Covid-19:</b></p> <ul style="list-style-type: none"> <li>- 77% had more than 2 coexisting conditions, and <b>greater coexisting conditions associated with SARS-CoV-2 tropism for kidneys.</b></li> <li>- <b>highest SARS-CoV-2 copies per cell = respiratory tract,</b></li> <li>- <b>lower viral copies per cell = kidneys, liver, heart, brain, and blood.</b></li> </ul> <p><b>Kidney tissue microdissection from 6 patients :</b></p> <ul style="list-style-type: none"> <li>- 3 = detectable SARS-CoV-2 <b>viral load in all kidney compartments</b> examined, with <b>preferential targeting of glomerular cells.</b></li> <li>-&gt; <b>SARS-CoV-2 organotropism beyond respiratory tract includes heart, liver, brain, and kidneys. Renal tropism is a potential explanation of commonly reported new clinical signs of kidney injury in Covid-19 patients</b></li> </ul>
The Journal of Molecular Diagnostics 13MAY2020	<b>Detection of SARS-CoV-2 is comparable in clinical samples preserved in saline or viral transport media</b>	Radbel, Jared and al. USA <a href="#">gotopaper</a>	Virology	<p>The availability of viral transport media (VTM) has become severely limited, contributing to delays in diagnosis and rationing of diagnostic testing.</p> <p>The phosphate buffered saline (PBS) may be a viable transport medium, as an alternative to VTM, for clinical qPCR testing. We assessed the intra- and inter-individual reliability of SARS-CoV-2 qPCR in clinical endotracheal secretion samples transported in VTM or PBS, evaluating the stability of the RT-qPCR signal for three viral targets (N gene, ORF1ab, and S gene) when samples were stored in these media at room temperature for up to 18 hours.</p> <p>Results: using PBS as a transport medium has high intra- and inter-individual reliability, maintains viral stability, and is comparable to VTM in the detection of the three SARS-CoV-2 genes through 18 hours of storage.</p> <p>CI<sup>o</sup>: PBS as a clinically useful medium for transporting and short-term preservation of specimens containing SARS-CoV-2 has the potential to increase testing capacity for SARS-CoV-2</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Lancet 13MAY2020	<b>An outbreak of severe Kawasaki-like disease at the Italian epicentre of the SARS-CoV-2 epidemic: an observational cohort study</b>	Verdoni L et al Italy <a href="#">gotopaper</a>	Clinic	<p>29 children with Kawasaki disease Group1(n=19): before SARS-CoV-2 outbreak Group2(n=10): after SARS-CoV-2 outbreak <u>Group1 versus group2:</u></p> <ul style="list-style-type: none"> <li>- Higher incidence in group2: 0,3 vs 10/month (p&lt;0,05)</li> <li>- Older group2: 3,0 vs 7,5 y (p&lt;0,05)</li> <li>- Abnormal echocardiogram: 60%(grp2) vs 10%(grp1), p&lt;0,05</li> <li>- More MAS group2: 50% vs 0%</li> </ul> <p>→ 30-fold increased incidence of KD in the past month</p> <p><b>→ SARS-CoV-2 outbreak is associated with high incidence of severe form of KD</b></p>
Nature Medicine 13MAY2020	<b>Infection of bat and human intestinal organoids by SARS-CoV-2</b>	Zhou, Jie et al. Hong Kong <a href="#">gotopaper</a>	Fundamental research	<p>Establishment and characterization of intestinal organoids derived from horseshoe bats (<i>Rhinolophus sinicus</i>) can recapitulate bat intestinal epithelium:</p> <ul style="list-style-type: none"> <li>- bat enteroids are fully susceptible to SARS-CoV-2 infection and robust viral replication.</li> <li>- human intestinal organoids also sustain active replication of SARS-CoV-2</li> </ul> <p><b>-&gt; First expandable organoid culture system of bat intestinal epithelium and evidence that SARS-CoV-2 can infect bat intestinal cells.</b></p> <p>-&gt; Robust SARS-CoV-2 replication in human intestinal organoids suggests that the human intestinal tract might be a transmission route of SARS-CoV-2.</p>
JAMA 13MAY2020	<b>SARS-CoV-2 Rates in BCG-Vaccinated and Unvaccinated Young Adult</b>	Hamiel et al., Israel <a href="#">gotopaper</a>	Vaccine	<p>The BCG vaccine was routinely administered to all newborns in Israel as part of the national immunization program between 1955 and 1982</p> <p>Since 1982, the vaccine has been administered only to immigrants from countries with high prevalence of tuberculosis. This change allowed comparison of infection rates and proportions with severe COVID-19 disease in 2 similar populations with differing BCG status: individuals born during the 3 years before and 3 years after cessation of the universal BCG vaccine program.</p> <p><b>-&gt; This study does not support the idea that BCG vaccination in childhood has a protective effect against COVID-19 in adulthood.</b></p>
Annals of Internal Medicine 12MAY2020	<b>Pharmacokinetics of Lopinavir and Ritonavir in Patients Hospitalized With Coronavirus Disease 2019 (COVID-19)</b>	Schoergenhofer, Christian et al. Austria <a href="#">gotopaper</a>	Therapeutic	<p><b>First pharmacokinetic data of lopinavir and ritonavir in patients hospitalized with COVID-19.</b> Series of 8 patients.</p> <ul style="list-style-type: none"> <li>- Lopinavir trough levels were approximately <b>2-fold higher</b> in this population <b>than in patients with HIV</b> receiving the same dose (7.1 µg/mL).</li> <li>- A correlation of drug concentrations with C-reactive protein, a downstream marker of IL-6, was observed.</li> </ul> <p>However, approximately <b>60- to 120-fold higher concentrations are required to reach the assumed EC50 at trough levels</b>, making effective treatment of COVID-19 with lopinavir and ritonavir at the currently used doses unlikely.</p> <p><b>Limitations:</b> only trough levels were quantified, more detailed pharmacokinetics not available ; no data on the half-maximal effective dose of lopinavir for SARS-CoV-2 in vivo.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 12MAY2020	<b>Respiratory disease in rhesus macaques inoculated with SARS-CoV-2</b>	Munster, Vincent J. et al. USA <a href="#">gotopaper</a>	Animal model	<p>SARS-CoV-2 causes respiratory disease in infected rhesus macaques, with disease lasting 8-16 days:</p> <ul style="list-style-type: none"> <li>- 8 adult rhesus macaques (4 males, 4 females, age 4-6 yrs) inoculated with combination of intranasal (0.5ml per nostril), intratracheal, oral and ocular of a <math>4 \times 10^5</math> TCID<sub>50</sub>/ml (<math>3 \times 10^8</math> genome copies/ml).</li> <li>- Pulmonary infiltrates visible in lung radiographs</li> <li>- High viral load in nose and throat swabs and bronchoalveolar lavages of all animals.</li> <li>- prolonged rectal shedding detected in 1 animal</li> </ul> <p><b>-&gt; Rhesus macaque recapitulates moderate disease with regard to virus replication, shedding, presence of pulmonary infiltrates, histological lesions and seroconversion.</b></p>
Nat Med 12MAY2020	<b>A serological assay to detect SARS-CoV-2 seroconversion in humans</b>	Amanat et al., USA <a href="#">gotopaper</a>	Diagnostic	<p><b>Describing a</b> a serological enzyme-linked immunosorbent assay for the screening and identification of human SARS-CoV-2 seroconverters.</p> <ul style="list-style-type: none"> <li>-&gt; based on reactivity to the immunogenic S protein of the virus, is relatively simple and quick in its execution and can be performed at biosafety level 2</li> <li>-&gt; there is no or only negligible cross-reactivity from human coronaviruses to SARS-CoV-2 in the tested individuals</li> <li>-&gt; strong seroconversion with ELISA AUC values in the 1:1,000 range after natural infection with SARS-CoV-2</li> </ul>
Cancer discovery 12MAY2020	<b>Impact of PD-1 blockade on severity of COVID-19 in patients with lung cancers</b>	Luo, Jia et al. USA <a href="#">gotopaper</a>	Therapeutic/ Clinic	<p>Analyses on <b>69 consecutive patients with lung cancers</b> who were diagnosed with COVID-19. Severity based on no or prior receipt of PD-1 blockade was examined.</p> <ul style="list-style-type: none"> <li>- Overall, the <b>severity of COVID-19 in patients with lung cancer was high</b>, including need for hospitalization in more than half of patients and death in nearly a quarter.</li> <li>- Prior PD-1 blockade was, as expected, associated with smoking status.</li> <li>- After adjustment for smoking status, <b>PD-1 blockade exposure was not associated with increased risk of severity of COVID-19</b>. PD-1 blockade does not appear to impact the severity of COVID-19 in patients with lung cancers.</li> </ul> <p>These initial results in patients with lung cancers support the <b>safety of PD-1 blockade treatment to achieve optimal cancer outcomes</b>.</p>
JAMA Intern Med 12MAY2020	<b>Development and Validation of a Clinical Risk Score to Predict the Occurrence of Critical Illness in Hospitalized Patients With COVID-19</b>	Liang W et al, China <a href="#">gotopaper</a>	Clinic	<p>Develop and validate a clinical score at admission for predicting critical illness</p> <p>Retrospective cohort (575 hospital in China)</p> <p><u>1590 patients with data were include for variable selection:</u></p> <ul style="list-style-type: none"> <li>- Mean age: 48,9y – 57,3% were men – 25,1% had at least 1 comorbidity</li> <li>- 72 variables entered in selection process (LASSO and logistic regression)</li> </ul> <p><u>10 variables were independently statistically significant predictors of critical illness:</u></p> <ul style="list-style-type: none"> <li>- CXR abnormality: OR: 3,39</li> <li>- Age OR: 1,03</li> <li>- Hemoptysis OR 4,53</li> <li>- Dyspnea OR 1,88</li> <li>- Unconsciousness OR 4,71</li> <li>- Number of comorbidities OR 1,60</li> <li>- Cancer history OR 4,07</li> <li>- Neutrophil to lymphocyte ratio OR 1,06</li> <li>- Lactate dehydrogenase OR: 1,002</li> <li>- Direct bilirubin OR: 1,15</li> </ul> <p><u>Validation:</u> cohort of 710 patients</p> <p>AUC of COVID-GRAM 0,88 IC<sub>95%</sub> [0,84 – 0,93]</p> <p><u>Limitations:</u> data are entirely from China</p> <p><b>→ risk score at the admission for predicting critical illness</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 11MAY2020	Association of Treatment With Hydroxychloroquine or Azithromycin With In-Hospital Mortality in Patients With COVID-19 in New York State	Rosenberg, Eli S. et al. USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Retrospective multicenter cohort study of 1438 hospitalized patients</b> who received both hydroxychloroquine and azithromycin, hydroxychloroquine alone, azithromycin alone, or neither.</p> <p>=&gt; Compared with patients receiving neither drug, there <b>were no significant differences in mortality</b> for patients receiving hydroxychloroquine + azithromycin (HR, 1.35 [95% CI, 0.76-2.40]), hydroxychloroquine alone (HR, 1.08 [95% CI, 0.63-1.85]), or azithromycin alone (HR, 0.56 [95% CI, 0.26-1.21]).</p> <p>=&gt; Compared with patients receiving neither drug, <b>cardiac arrest was significantly more likely in patients receiving hydroxychloroquine + azithromycin</b> (adjusted OR, 2.13 [95% CI, 1.12-4.05]), but not hydroxychloroquine alone (adjusted OR, 1.91 [95% CI, 0.96-3.81]) or azithromycin alone (adjusted OR, 0.64 [95% CI, 0.27-1.56]).</p> <p>=&gt; No significant differences in the relative likelihood of abnormal electrocardiogram findings.</p> <p><b>Limitations:</b> mortality limited to in-hospital death ; potential confounders such as inflammatory markers were not frequently measured ; confidence intervals for some of the findings are wide, reflecting limits in study power for some analyses</p>
Bio-design and manufacturing 11MAY2020	Development of a rapid test kit for SARS-CoV-2: an example of product design	Cui, Zhanfeng and al. China <a href="#">gotopaper</a>	Diagnostic	<p>The urgent need for large numbers of tests in field setting imposes constraints such as short test time and lack of access to specialist equipment, laboratories and skilled technicians to perform the test and interpret results.</p> <p>To meet these needs, an antigen test based on RT-LAMP with colorimetric readout was chosen. Direct use of swab sample with no RNA extraction was explored.</p> <p>After extensive experimental study, a rapid test kit has been fabricated to satisfy all design criteria</p>
J. Clin. Microbiol. 11MAY2020	Open Development and Clinical Validation Of Multiple 3D-Printed Nasopharyngeal Collection Swabs: Rapid Resolution of a Critical COVID-19 Testing Bottleneck	Callahan, Cody J. and al. USA <a href="#">gotopaper</a>	Diagnostic	<p>To address the shortage of the nasopharyngeal swabs, we designed and executed a translational-research program to allow immediate mass production by 3D printing.</p> <p>We validated four prototypes through an institutional review board (IRB)-approved clinical trial that involved 276 outpatient volunteers.</p> <p>Each participant was swabbed with a reference swab (the control) and a prototype, and SARS-CoV-2 reverse-transcriptase polymerase chain reaction (RT-PCR) results were compared.</p> <p>All prototypes displayed excellent concordance with the control. Contact information for ordering can be found at <a href="http://printedswabs.org">http://printedswabs.org</a></p>
Nature Medicine 11MAY2020	Real-time tracking of self-reported symptoms to predict potential COVID-19	Menni et al, UK <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>A total of 2,618,862 participants reported their potential symptoms of COVID-19 on a smartphone-based app.</p> <p>-&gt; Of 18,401 who had undergone a SARS-CoV-2 test, participants reporting loss of smell and taste was 65.03% in those with a positive test result and 21.72% in those with a negative test result (odds ratio = 6.74; 95% CI = 6.31–7.21).</p> <p>-&gt; A model combining symptoms to predict probable infection was applied to the data from all app users who reported symptoms (805,753) and predicted that 140,312 (17.42%) participants are likely to have COVID-19</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Journal of Allergy and Clinical Immunology 11MAY2020	<b>Safety and efficacy of early high-dose IV anakinra in severe COVID-19 lung disease</b>	Pontali, Emanuele et al. Italy <a href="#">gotopaper</a>	Therapeutic	<p><b>Pilot study of early use of high IV doses of anti-IL-1 anakinra in 5 patients</b> with severe/moderate COVID-19 with pulmonary involvement.</p> <p>-&gt;All five patients experienced <b>rapid resolution of systemic inflammation</b>, and <b>remarkable improvement of respiratory parameters</b>, with reduction of oxygen support requirement and early amelioration of chest CT scan abnormalities before discharge in 3 patients.</p> <p>-&gt; All patients were discharged 6 to 13 days after the start of anakinra.</p> <p>-&gt; No secondary infections or other adverse events were observed.</p> <p><b>Limitations:</b> non-controlled study ; small size ; short-term duration of the treatment ; variability of laboratory biomarkers.</p>
Clin. Infect. Dis. 11MAY2020	<b>Hydroxychloroquine in COVID-19 patients: what still needs to be known about the kinetics</b>	Martin-Blondel, G. et al. France <a href="#">gotopaper</a>	Therapeutic	<p>Aim: to determine whether or not the pharmacokinetics in systemic lupus erythematosus (SLE) patients can be applied to COVID-19 patients.</p> <p>Different dosage regimens were applied based on data that emerged: regimen 1 (200 mg x 3/day), regimen 2 (400 mg x 2 on day 1 followed by 200 mg x 3/day), regimen 3 (400 mg x 2 on day 1 followed by 400 mg x 1/day) and regimen 4 (600 mg x 2 followed by 400 mg x 1 /day).</p> <p>Blood samples (n=101) were collected from 57 COVID-19 patients for 7 days and concentrations were compared with simulated kinetic profiles.</p> <p>⇒ <b>Hydroxychloroquine exposure tends to be low</b> and in most instances lower than the values reported in SLE patients, in particular for the standard regimen of “200 mg x 3/day”.</p> <p>⇒ The <b>pharmacokinetic behavior in COVID-19 patients cannot be predicted by the SLE population or by rheumatoid arthritis patients</b>.</p>
European heart journal 11MAY2020	<b>Characteristics and clinical significance of myocardial injury in patients with severe coronavirus disease 2019</b>	Shi S and al, China <a href="#">gotopaper</a>	Clinic	<p>671 hospitalized patients COVID-19 Median age: 63 years Main comorbidities: hypertension (29,7%) –diabetes (14,5%) – CHD(9%) Case fatality rate 9,2%</p> <p><u>Death versus survivor group</u></p> <ul style="list-style-type: none"> <li>- Older and more often male (p&lt;0,001)</li> <li>- More comorbidities (p&lt;0,001)</li> <li>- More myocardial injury: 75,8% vs 9,7% (p&lt;0,001)</li> </ul> <p><u>Cardiac troponin I predicting in-hospital mortality:</u></p> <ul style="list-style-type: none"> <li>- AUC 0,92</li> <li>- Se 86% and Spe 86%</li> <li>- Single cut-off concentrations 73 µg/L</li> </ul> <p><u>Predictor of myocardial injury:</u></p> <ul style="list-style-type: none"> <li>- Older age – comorbidities</li> <li>- High level of CRP</li> </ul> <p>Limitation: small sample size, cause of death or myocardial injury underestimated</p> <p>→ <b>CtnI and CK-MB predict risk for in hospital mortality</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA ped 11MAY2020	<b>Characteristics and Outcomes of Children with Coronavirus Disease 2019 (COVID-19) Infection Admitted to US and Canadian Pediatric Intensive Care Units</b>	Shekerdeman LS and al, USA <a href="#">gotopaper</a>	Clinic	<p>46 PICU (March 14 and April 3) → 48 children (40 in US &amp; 6 in Canada)</p> <p>Median age: 13 years [4,2 – 16,6]</p> <p>Comorbidities (83%)</p> <p>Median PICU length of stay: 5 days</p> <p>Respiratory symptoms: 73 %</p> <p>→ 39/48 required ventilatory support: 21 non-invasively and 19 IMV</p> <p>Specific therapies (28/46): Hydroxychloroquine or hydroxy+azithro or remdesivir or tocilizumab</p> <p>Case fatality rate: 4,2% (2/48)</p> <p>→ <b>pre-hospital comorbidities = important factor</b></p>
Emerging Infectious Disease journal 08MAY2020	<b>Prolonged Persistence of SARS-CoV-2 RNA in Body Fluids</b>	Jiufeng Sun and al. <a href="#">gotopaper</a>	Virology	<p>To estimate the frequency and duration of detectable SARS-CoV-2 RNA in human body fluids. The prolonged persistence of virus RNA in various body fluids may guide the clinical diagnosis and prevention of onward virus transmission.</p> <p>43 patients with mild cases of COVID-19 - 490 specimens collected.</p> <p><b>Results:</b> through an AFT-based modeling study: persistent shedding of virus RNA in nasopharyngeal swab and feces samples. The estimated time until loss of virus RNA detection ranged from 45.6 days for nasopharyngeal swab samples to 46.3 days for feces samples in mild cases and from 48.9 days for nasopharyngeal swab samples to 49.4 days for feces samples in severe cases (longer than those for SARS-Cov and MERS-Cov).</p> <p><b>CI<sup>2</sup>:</b> results show prolonged persistence of SARS-CoV-2 RNA in hospitalized patients with COVID-19. Health professionals should consider these findings in diagnostic recommendations and prevention measures for COVID-19.</p>
Pediatric Blood Cancer 08MAY2020	<b>COVID-19 infection in children and adolescents with cancer in Madrid</b>	De Roja T and al, Spain <a href="#">gotopaper</a>	Clinic	<p>15 pediatric oncology patients</p> <p>Median age: 10,6 years [0,6 – 18,6]</p> <p>Hematological malignancy (73%) and solid tumor (27%)</p> <p>60% received chemotherapy in the 15 days before infection</p> <p><b>Symptoms:</b> fever (67%) – cough (40%) – asymptomatic (13%)</p> <p><b>Radiological finding:</b> 8/14 pathological findings</p> <p>2 patients received oxygen therapy</p> <p>Median hospital stays: 8 days</p> <p><b>All favorable outcome</b></p> <p>→ <b>prevalence among children with cancer in Madrid: 1,3%</b></p> <p>→ mild symptomatic and better prognosis than adults</p>
American Journal of Obstetrics & Gynecology MFM 08MAY2020	<b>Clinical course of severe and critical COVID-19 in hospitalized pregnancies: a US cohort study</b>	Rebecca Am, and al USA <a href="#">gotopaper</a>	Clinic	<p><b>64 pregnant women</b> hospitalized: 44 severe and 20 critical – no death</p> <p>Gestational age at admission: 30 ± 6 weeks</p> <p>Admission: 7 days after onset symptoms</p> <p>Majoration of dyspnea at day 8 and MV at day 9</p> <p>Median duration of hospital stays: 6 day for severe and 12 for critical</p> <p><b>Delivery preterm:</b> 75% of critical women</p> <ul style="list-style-type: none"> <li>- Severe: 37 ± 2</li> <li>- Critical: 32 ± 4</li> </ul> <p><b>Critical cases:</b> 95% required MV - 70% ARDS - 20% prone position</p> <p><b>Neonate:</b></p> <ul style="list-style-type: none"> <li>- 64% need ICU</li> <li>- One tested positive at 48-h without any symptoms</li> </ul> <p>→ clinical course not different from not pregnant women</p> <p>→ <b>pregnancy should not be considered an independent risk of factor</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Cell 08MAY2020	Host-viral infection maps reveal signatures of severe COVID-19 patients	Bost, Pierre; et al. Israel-France-China <a href="#">gotopaper</a>	Fundamental research	<p>Viral-Track, a new computational method to analyze host-viral infection maps :</p> <ul style="list-style-type: none"> <li>- <b>enables transcriptional sorting of infected vs bystander cells and reveals virus-induced expression</b> (scans unmapped scRNA-seq data for presence of viral RNA).</li> <li>- Applicable to multiple models of infection (HBV, HIV, VSV, etc)</li> <li>Applied to Bronchoalveolar-Lavage samples from severe vs mild COVID-19 patients, reveals: <ul style="list-style-type: none"> <li>- SARS-CoV-2 infects epithelial cells and alters immune landscape in severe patients.</li> <li>- <b>detected unexpected coinfection with hMPV</b> (human MetaPneumoVirus) mainly in monocytes, dampening interferon response.</li> </ul> </li> </ul> <p>-&gt; <b>robust technology for dissecting mechanisms of viral-infection and pathology.</b></p>
Lancet 08MAY2020	Triple combination of interferon beta-1b, lopinavir-ritonavir, and ribavirin in the treatment of patients admitted to hospital with COVID-19: an open-label, randomised, phase 2 trial	Hung, Ivan Fan-Ngai et al. China <a href="#">gotopaper</a>	Therapeutic	<p><b>Multicentre, prospective, open-label, randomised, phase 2 trial in 127 adults</b> with COVID-19 hospitalized in Hong Kong. Random assignement (2:1) to the <b>combination group (lopinavir-ritonavir + ribavirin+ interferon beta-1)</b> or to the <b>control group (lopinavir-ritonavir)</b>.</p> <p>=&gt; The <b>triple combination</b>, when <b>given within 7 days of symptom onset</b>, is <b>effective in suppressing the shedding of SARS-CoV-2</b>, not just in a nasopharyngeal swab, but in all clinical specimens, compared with lopinavir-ritonavir alone.</p> <p>=&gt; The <b>significant reductions in duration of RT-PCR positivity and viral load were associated with clinical improvement</b> as shown by the significant reduction in NEWS 2 and duration of hospital stay.</p> <p>=&gt; Subgroup comparison suggested interferon beta-1b to be a key component of the combination treatment.</p> <p><b>Limitations:</b> open label, no placebo group, confounded by a subgroup omitting interferon beta-1b within the combination group, no critically ill patients.</p>
Science 08MAY2020	A highly conserved cryptic epitope in the receptor-binding domains of SARS-CoV-2 and SARS-CoV	Yuan, Meng; et al. USA-China <a href="#">gotopaper</a>	Fundamental research	<p>Crystal <b>structure of CR3022 (neutralizing antibody from convalescent SARS-CoV infected patient) in complex with the receptor-binding domain of the SARS-CoV-2 spike</b> :</p> <ul style="list-style-type: none"> <li>- 3.1a resolution</li> </ul> <p>-&gt; <b>CR3022 targets a highly conserved epitope (conserved in SARS-CoV-2 and SARS-CoV) that is distal from the receptor binding site, and enables cross-reactive binding between SARS-CoV-2 and SARS-CoV.</b></p> <ul style="list-style-type: none"> <li>- CR3022 likely binds more tightly to SARS-CoV because its epitope contains a glycan absent in SARS-CoV-2.</li> <li>-&gt; Modeling showed this epitope only accessible when at least 2 of the 3 spike proteins are in a conformation competent to bind the receptor.</li> </ul>
Science 08MAY2020	Harnessing multiple models for outbreak management	Shea et al, USA <a href="#">gotopaper</a>	Public Health/Epidemio	<p>COVID-19 pandemic has triggered the development of several valuable models that can differ in various elements and provide disparate predictions, which could ultimately hinder intervention planning and response by policymakers.</p> <p>We advocate a more systematic approach, by merging two well-established research fields.</p> <ol style="list-style-type: none"> <li>1. Formal expert elicitation methods applied to multiple models to deliberately generate, retain, and synthesize valuable individual model ideas and share important insights during group discussions, while minimizing various cognitive biases.</li> <li>2. Using a decision-theoretic framework to capture and account for within- and between-model uncertainty as we evaluate actions in a timely manner to achieve management objectives.</li> </ol>



Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 08MAY2020	<b>Changes in SARS-CoV-2 Positivity Rate in Outpatients in Seattle and Washington State, March 1-April 16, 2020</b>	Randhawa et al, USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>Patient demographics: SARS-CoV-2 positivity rates were 8.2% in Washington State outpatient clinics, 8.4% in Seattle-area outpatient clinics, and 14.4% in Seattle EDs.</p> <p>The SARS-CoV-2 positivity rate was analysed by fitting penalized cubic regression splines to binomial testing data, and accounting for variation in the daily testing totals. SARS-CoV-2 positivity rate was 17.6% in the outpatient clinics and 14.3% in EDs at the peak period and 3.8% and 9.8%, respectively, at the end of the analysis period.</p> <p>SARS-CoV-2 infections in patients of Washington outpatient clinics and Seattle ED settings peaked in late March and have been declining.</p> <p>-&gt; This trajectory is aligned with local physical distancing and the "Stay Home, Stay Healthy" order announced March 23, 2020.</p>
Nature structural & molecular biology 07MAY2020	<b>Structural basis for the inhibition of SARS-CoV-2 main protease by antineoplastic drug carmofur</b>	Jin, Zhenming et al. China <a href="#">gotopaper</a>	Therapeutic	<p>The antineoplastic drug <b>carmofur is shown to inhibit the SARS-CoV-2 main protease (<math>M^{pro}</math>)</b>. The X-ray crystal structure of <math>M^{pro}</math> in complex with carmofur reveals that the carbonyl reactive group of carmofur is covalently bound to catalytic Cys145, whereas its fatty acid tail occupies the hydrophobic S2 subsite. <b>Carmofur inhibits viral replication in VeroE6 cells (<math>EC_{50} = 24.30 \mu M</math>)</b>. Carmofur has a favorable selectivity index (SI) of 5.36, but further optimization will be required to develop an effective drug.</p> <p>This study provides a basis for rational <b>design of carmofur analogs with enhanced inhibitory efficacy to treat COVID-19</b>.</p>
The Lancet Rheumatology 07MAY2020	<b>Interleukin-1 blockade with high-dose anakinra in patients with COVID-19, acute respiratory distress syndrome, and hyperinflammation: a retrospective cohort study</b>	Cavalli, Giulio et al. Italy <a href="#">gotopaper</a>	Therapeutic	<p><b>Retrospective cohort study</b> in adult patients with COVID-19, moderate-to-severe ARDS, and hyperinflammation, managed with non-invasive ventilation outside of the ICU and who received <b>standard treatment of hydroxychloroquine and lopinavir-ritonavir, with or without anakinra</b>.</p> <p><b>29 patients</b> received high-dose intravenous anakinra, <b>16 patients</b> comprised the comparison group for this study, and 7 patients received low-dose subcutaneous anakinra but treatment was interrupted after 7 days.</p> <p>At 21 days, <b>treatment with high-dose anakinra was associated with clinical improvement in 21 (72%)</b> of 29 patients versus 8 (50%) in the standard treatment group. At 21 days, <b>survival was 90% in the high-dose anakinra group</b> and 56% in the standard treatment group (<math>p=0.009</math>). Discontinuation of anakinra was not followed by inflammatory relapses.</p> <p><b>Limitations:</b> retrospective nature, relatively small size of the cohorts (particularly the historical comparator group), a more extended follow-up is also needed to assess long-term outcomes of treated patients.</p>
Journal of Clinical Virology 07MAY2020	<b>SARS-CoV-2 detection by direct rRT-PCR without RNA extraction</b>	Merindol, Natacha, and al. <a href="#">gotopaper</a>	Diagnostic	<p>There are many challenges associated with ramping up testing capacity, including shortage in the chain of supplies for extraction reagents. This situation called for alternatives protocols with similar sensitivity to ensure the continuity of testing in laboratories.</p> <p>-&gt; Comparison of sensitivity of 2 approved rRT-QPCR Assays with and without RNA extraction.</p> <p><b>Conclusion :</b> the two tests provided the same sensitivity. Direct rRT-PCR without RNA extraction is possible if samples are in UTM or molecular water; specimens collected in water should be screened rapidly. RNA extraction is necessary if samples are in saline water or Hanks medium.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 07MAY2020	<b>Necessity of a development and widespread distribution of COVID-19 medical treatments</b>	Bollyky, T. et al, US <a href="#">gotopaper</a>	HSS/Politic	<p>1- Equitable Distribution Plan for manufacturing capacity, financing, and distribution infrastructure necessary to produce sufficient quantities to meet global needs in a fair, public health-driven manner.</p> <p>2- Framework for Distribution =&gt;For flexible, trusted governance - take advantage of well-established international forums (e.g: G7) rather than build something new + central role of WHO in planning and coordinating the implementation of the framework - requires coordination of several institutions, donors, governments and pharmaceutical companies - COVID-19 access accelerator (ACT), recently launched and supported by the European Commission, should extend to other major contributors, members of the G20. - involve entities that develop vaccines, treatments and diagnostics and support group supply in LMICs (CEPI, Gavi, the Global Fund...)</p> <p>=&gt; Adequate, Predictable Financing Provide a funding mechanism to generate income for R&amp;D + deployment of vaccines and therapeutic products (advance purchase commitments (APC) for COVID-19 products + profiles of target products) Funds =&gt; mixture of national and philanthropic contributions, mobilized to raise additional funds on the capital markets Contribution from world leaders + subscription depending on payment capacity of countries. = low-income countries highly subsidized /free. Transparent regulatory pathway for approval of COVID-19 products → instill global confidence, reduce development costs and accelerate access to less profitable markets.</p>
The Lancet. Respiratory medicine 07MAY2020	<b>Tropism, replication competence, and innate immune responses of the coronavirus SARS-CoV-2 in human respiratory tract and conjunctiva: an analysis in ex-vivo and in-vitro cultures</b>	Hui, Kenrie P. Y.; et al. China <a href="#">gotopaper</a>	Fundamental research	<p>SARS-CoV-2 tissue and cellular tropism in ex-vivo cultures of human bronchus, lung, conjunctiva, and innate immune responses vs other coronavirus and influenza virus (H1N1).</p> <p>SARS-CoV-2 isolated from COVID-19 patients: - infected ciliated, mucus-secreting, and club cells of bronchial epithelium, type 1 pneumocytes in the lung, and the conjunctival mucosa. - In bronchus : replication similar to MERS-CoV, and higher than SARS-CoV, but lower than H1N1. - In lung: replication similar to SARS-CoV and H1N1, but lower than MERS-CoV. - In conjunctiva: replication greater than SARS-CoV. - <b>SARS-CoV-2 was a less potent inducer of proinflammatory cytokines than H5N1, H1N1pdm, or MERS-CoV.</b></p> <p>-&gt; <b>Conjunctival epithelium and conducting airways are potential portals of infection for SARS-CoV-2. SARS-CoV-2 replicated similarly to SARS-CoV in alveolar epithelium; but more extensively in bronchus.</b></p>
Cell 07MAY2020	<b>Coast-to-Coast Spread of SARS-CoV-2 during the Early Epidemic in the United States</b>	Fauver et al, USA <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>SARS-CoV-2 detected in all 50 states of USA. Data on sequencing of nine viral genomes from early reported COVID-19 patients, the majority of which from Washington State, combined with domestic and international travel patterns, showed:</p> <p>- SARS-CoV-2 transmission in Connecticut was likely driven by domestic introductions - the risk of domestic importation to Connecticut exceeded that of international importation by mid-March regardless of our estimated effects of federal travel restrictions</p> <p>→ Widespread transmission of SARS-CoV-2 within USA, need for critical surveillance</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Infect. Control Hosp. Epidemiol. MAY2020	<b>Effect of ambient air pollutants and meteorological variables on COVID-19 incidence</b>	Jiang et al, China <a href="#">gotopaper</a>	Public Health/Epidemiology	<p>The multivariate Poisson regression used to analyse correlation between COVID-19 incidence, eight air pollutants and three meteorological variables in three China's worst COVID-19 hit provinces.</p> <p>Daily COVID-19 incidence was positively associated with PM2.5 and humidity in all cities. The relative risk (RR) of PM2.5 was 1.036 (95% CI, 1.032 - 1.039), 1.059 (95% CI, 1.046 - 1.072) and 1.144 (95% CI, 1.12 - 1.169) for COVID-19 incidence per day in the three provinces. The RR of humidity was lower than that of PM2.5, difference ranging from 0.027 to 0.111. PM10 and temperature exhibited a negative correlation with daily COVID-19 incidence: the RR of PM10 ranged from 0.915 (95% CL, 0.896 - 0.934) to 0.961 (95% CL, 0.95 - 0.972) while that of temperature was 0.738 (95% CL, 0.717 - 0.759) to 0.969 (95% CL, 0.966 - 0.973).</p> <p>Data suggest that PM2.5/humidity and PM10/temperature could substantially increase and decrease the risk of COVID-19 incidence, respectively.</p>
Nature 07MAY2020	<b>The pathogenicity of SARS-CoV-2 in hACE2 transgenic mice</b>	Bao, Linlin; et al. China <a href="#">gotopaper</a>	Animal model	<p><b>Human ACE2 transgenic mice infected with SARS-CoV-2 :</b></p> <ul style="list-style-type: none"> <li>- SARS-CoV-2 intranasal inoculation at <math>10^5</math> TCID<sub>50</sub>/50 µL inoculum volume per mouse. (14 days observation)</li> <li>- 6-11 month-old, male and female WT (n=15) and hACE2 mice (n=19).</li> <li>- Typical histopathology: interstitial pneumonia with infiltration of significant macrophages and lymphocytes into the alveolar interstitium, and accumulation of macrophages in alveolar cavities.</li> <li>- <b>Weight loss observed in hACE2 mice (up to 8% at 5 dpi), not in WT mice.</b></li> <li>- <b>viral load detectable</b> (qRT-PCR) at 1, 3, 5 and 7 dpi (peak at 3 dpi) <b>in lungs of hACE2 mice but not in WT mice.</b></li> <li>- infectious virus isolated from lungs of hACE2 mice at 1, 3 dpi and 5 dpi (peak titers at 3 dpi), but not WT mice.</li> <li>- <b>Viral antigens detected</b> in bronchial epithelial cells, macrophages and alveolar epithelia.</li> </ul> <p>-&gt; <b>Confirmed pathogenicity of SARS-CoV-2 in hACE2 expressing mice and suggests that hACE2 was essential for infection and replication in mice.</b></p>
The Journal of Infectious Diseases 07MAY2020	<b>T cell subset counts in peripheral blood can be used as discriminatory biomarkers for diagnosis and severity prediction of COVID-19</b>	Jiang, Mei and al. China <a href="#">gotopaper</a>	Diagnostic	<p>Assessment of the significance of lymphocyte subsets detection in peripheral blood in the diagnosis and prognosis of Covid-19 disease.</p> <p>The counts of CD8+T and CD4+T cells can be used as diagnostic markers of COVID-19 and predictors of disease severity.</p>
The Lancet Psychiatry 07MAY2020	<b>COVID-19, unemployment, and suicide</b>	Kawohl And Nordt., Switzerland <a href="#">gotopaper</a>	Psy	<p><b>High scenario:</b> the worldwide unemployment rate would increase from 4-936% to 5-644%, which would be associated with an increase in suicides of about 9570 per year.</p> <p><b>Low scenario:</b> the unemployment would increase to 5-088%, associated with an increase of about 2135 suicides.</p>
New England Journal of Medicine 07MAY2020	<b>Observational Study of Hydroxychloroquine in Hospitalized Patients with Covid-19</b>	Geleris, Joshua et al. USA <a href="#">gotopaper</a>	Therapeutic	<p><b>Observational study</b> involving consecutive patients with Covid-19 admitted to a hospital, and <b>comparing outcomes in patients who received hydroxychloroquine with those in patients who did not.</b> The primary end point was a composite of intubation or death in a time-to-event analysis.</p> <p>Of the <b>1376 patients</b>, 811 (58.9%) received hydroxychloroquine and 565 (41.1%) did not. Hydroxychloroquine-treated patients were more severely ill at baseline than those who did not receive hydroxychloroquine.</p> <p>Overall, 346 patients (25.1%) had a primary end-point of respiratory failure. In the main analysis, there was <b>no significant association between hydroxychloroquine use and intubation or death</b> (hazard ratio, 1.04, 95% confidence interval, 0.82 to 1.32). Results were similar in multiple sensitivity analyses.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Nat Rev Immunology 06MAY2020	<b>Pathological inflammation in patients with COVID-19: a key role for monocytes and macrophages</b>	Merad et al., USA <a href="#">gotopaper</a>	Immuno	<p>The delay in production of type I interferon promotes the enhanced release of monocyte chemoattractants by alveolar epithelial cells leading to sustained recruitment of blood monocytes into the lungs.</p> <ul style="list-style-type: none"> <li>Monocytes differentiate into pro-inflammatory macrophages</li> <li>Activated natural killer (NK) cells and T cells further promote the recruitment and activation of monocyte-derived macrophages through the production of granulocyte-macrophage colony-stimulating factor (GM-CSF), tumour necrosis factor (TNF) and interferon-<math>\gamma</math> (IFN<math>\gamma</math>).</li> <li>Oxidized phospholipids (OxPLs) are accumulated in infected lungs and activate monocyte-derived macrophages through the Toll-like receptor 4 (TLR4)–TRAF6–NF-<math>\kappa</math>B pathway.</li> <li>It is possible that type I interferons induce the expression of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) entry receptors, enabling the virus to gain access to the cytoplasm of macrophages and to activate the NLRP3 inflammasome, leading to the secretion of mature IL-1<math>\beta</math> and/or IL-18.</li> <li>IL-1<math>\beta</math> can amplify activation of monocyte-derived macrophages in an autocrine or paracrine way, but it can also reduce type I interferon production in infected lungs.</li> <li>The engagement of Fc<math>\gamma</math> receptors (Fc<math>\gamma</math>Rs) by anti-spike protein IgG immune complexes can contribute to increased inflammatory activation of monocyte-derived macrophages.</li> </ul> <p><b>Conclusion:</b> Identifying the mechanisms that contribute to reduced type I interferon activity will be critical for the development of targeted immunomodulatory strategies in patients with COVID-19</p>
Antimicrobial agents and chemotherapy 06MAY2020	<b>Inhibition of SARS-CoV-2 infection by the cyclophilin inhibitor Alisporivir (Debio 025)</b>	Softic, Lauren et al. France, <a href="#">gotopaper</a>	Therapeutic	<p>Cyclophilins play a key role in the lifecycle of coronaviruses. <b>Alisporivir (Debio 025) is a non-immunosuppressive analogue of cyclosporin A with potent cyclophilin inhibition properties.</b> It has been administered to more than 1,800 patients with chronic hepatitis C virus infection in Phase 2 and 3 clinical trials, alone or in combination with pegylated interferon alpha and/or ribavirin.</p> <p>=&gt; Alisporivir reduced SARS-CoV-2 RNA production in a dose-dependent manner in VeroE6 cell line, with an <b>EC50 of 0.46<math>\pm</math>0.04 <math>\mu</math>M</b>.</p> <p>=&gt; Anti-SARS-CoV-2 effectiveness of alisporivir was confirmed by immunofluorescence.</p> <p>=&gt; Alisporivir did not inhibit SARS-CoV-2 entry into VeroE6 cells. Effect of alisporivir was preserved when the compound was added 3 h post-infection. These results suggest that <b>alisporivir inhibits a post-entry step of the SARS-CoV-2 life cycle.</b></p>
Journal of the American College of Cardiology 06MAY2020	<b>Association of Treatment Dose Anticoagulation with In-Hospital Survival Among Hospitalized Patients with COVID-19</b>	Paranjpe, Ishan et al. USA <a href="#">gotopaper</a>	Therapeutic	<p>Association between administration of in-hospital anticoagulation (AC) and survival in a large cohort of 2,773 hospitalized patients with COVID-19, among which 786 (28%) received systemic AC during their hospital course.</p> <p>=&gt; <b>Systemic AC may be associated with improved outcomes</b> among patients hospitalized with COVID-19. Potential benefits of systemic AC need to be weighed against the risk of bleeding and therefore should be individualized. The association of in-hospital AC and mechanical ventilation likely reflects reservation of AC for more severe clinical presentations.</p> <p>=&gt; <b>Association with AC and improved survival after adjusting for mechanical ventilation.</b></p> <p><b>Limitations:</b> observational study, unobserved confounding, unknown indication for AC, lack of metrics to further classify illness severity in the mechanically ventilated subgroup, and indication bias.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
BMC 06MAY2020	A single-center, retrospective study of COVID-19 features in children: a descriptive investigation	Ma H, and al China <a href="#">gotopaper</a>	Clinic	<p>50 children COVID19+ and 26 with PCR- and CT+ with history of exposure</p> <p><u>Symptoms:</u></p> <ul style="list-style-type: none"> <li>- More frequent: fever (64%) – cough (44%)</li> <li>- Less frequent: diarrhea (6%) – abdominal pain (4%) – rhinorrhea (16%)</li> <li>- Asymptomatic 12%</li> </ul> <p><u>Laboratory:</u></p> <ul style="list-style-type: none"> <li>- Lymphopenia (16%) – thrombocytopenia (14%)</li> <li>- Elevated CRP (20%) – anemia (12%)</li> </ul> <p>43/50 had abnormalities on CT:          - Ground glass opacity (64%)          29/50 &gt;1 CT which 65% had improved CT and 28% had more lesions</p> <p><b>At discharge: no association between changes in CT lesions</b>          →CT allow to detect COVID19 but do not evaluated the resolution of illness for children</p>
Cell host & microbe PREPRINT	Identification of human single-domain antibodies against SARS-CoV-2	Yanling Wu et al, China <a href="#">gotopaper</a>	Therapeutic	<p>SARS-CoV-2 spike protein, containing the receptor-binding domain (RBD) and S1 subunit involved in receptor engagement, is a potential therapeutic target.</p> <ul style="list-style-type: none"> <li>- <b>Development of a phage-displayed single-domain antibody library</b> by grafting naïve complementarity-determining regions (CDRs) into framework regions of a human germline immunoglobulin heavy chain variable region (IGHV) allele.</li> <li>- Panning this library against SARS-CoV-2 RBD and S1 subunit <b>identified fully human single-domain antibodies targeting five distinct epitopes on SARS-CoV-2 RBD with subnanomolar to low nanomolar affinities</b>. Some of these antibodies neutralize SARS-CoV-2 by targeting a cryptic epitope located in the spike trimeric interface.</li> </ul>
SCIENCE 06MAY2020	Development of an inactivated vaccine candidate for SARS-CoV-2	Qiang Gao et al, China <a href="#">gotopaper</a>	Vaccine	<p><b>Development of PiCoVacc Vaccine based on the inactivated SARS-CoV-2 virus (Sinovac vaccine currently in ph1)</b>          -&gt; Isolation of 11 SARS-CoV-2 strains COVID19 patients BALF (China, Italy, Switzerland, UK, and Spain) to developed preclinical in vitro neutralization and challenge models.          A strain from a Chinese patient inactivated with b-propionolactone was used for vaccine development (not causing severe disease).</p> <p><b>Immunogenicity in BAL/C mice:</b>          -&gt; Inactivated Chinese patient virus+adjuvant (PiCoVacc) was injected at day 0 and 7 at different doses (0ug, 1,5ug, 3ug, 6ug)          -&gt; SARS-CoV-2 S and RBD specific IgGs were developed very quickly in mice sera          -&gt;The dominant immunogen was shown to be the RBD (no response to N protein)          -&gt;Neutralizing antibodies against all the SARS-CoV-2strains were also produced</p> <p><b>Immunogenicity and protection in macaques</b>          -&gt; 3 immunizations at d 0, 7 and 14 with at different doses of PiCoVacc (3 or 6ug)          -&gt; Specific S IgG and neutralizing antibodies were induced from two weeks after vaccination          -&gt; macaques at day 22 after fist immunization: viral loads were shown to decrease.          -&gt; No detectable viral load in pharynx, crissum and lung in high dose vaccinated monkeys from 7 d after challenge.          no changes on serological markers (CD3+, CD4+ CD88+, TNF-a, IFN-g, IL2, IL4...) nor pathologies in heart, spleen kidney or lung were observed suggesting that PiCoVacc do not induce exacerbated T cell response nor organ pathology.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Pediatr Infect Dis J 6MAY2020	<b>The risk of children Hospitalized with severe COVID-19 in Wuhan</b>	Wang Y and al China <a href="https://doi.org/10.1097/INF.0000000000002739">https://doi.org/10.1097/INF.0000000000002739</a>	Clinic	<p>Retrospective case-control study – 8 severe children matched with 35 - Median age of severe cases: 5,06 years 2/8 had comorbidities</p> <p><u>Symptoms in both groups</u>: fever – cough – dyspnea – diarrhea/vomiting</p> <p><u>Laboratory</u>:</p> <ul style="list-style-type: none"> <li>-WBC higher in severe group</li> <li>-No difference for lymphocytes counts <math>p&gt;0,05</math></li> <li>-IL6 – IL10, D-dimer higher in severe group</li> </ul> <p>Hospital stay: 13,5 (severe) versus 11 days (non severe) Time for PCR turning negative: 10,5 (severe) versus 7,1 (<math>p&lt;0,05</math>)</p> <p><b>More lung segment involves in severe children</b> → unique risk factor for severe</p> <p>→ mild symptomatic (8 severe cases on 260 COVID-19) – rare mortality → <b>some factors associated with severity</b>: CT scan lesions – immune response (IL-6) – intravascular coagulation (D-dimer)</p>
Science 05MAY2020	<b>Rapid implementation of mobile technology for real-time epidemiology of COVID-19</b>	Drew et al, USA <a href="https://doi.org/10.1126/science.abc0473">https://doi.org/10.1126/science.abc0473</a>	Public Health/Epidemiology	<p>The Coronavirus Pandemic Epidemiology (COPE) consortium developed a Symptom Tracker mobile application, launched in the UK on March 24, 2020 and the US on March 29, 2020 garnering more than 2.8 million users as of May 2, 2020.</p> <p>This mobile application offers data on risk factors, herald symptoms, clinical outcomes, and geographical hot spots.</p> <p>This initiative offers critical proof-of-concept for the repurposing of existing approaches to enable rapidly scalable epidemiologic data collection and analysis which is critical for a data-driven response to this public health challenge.</p>
Autoimmunity reviews 05MAY2020	<b>Continuous hydroxychloroquine or colchicine therapy does not prevent infection with SARS-CoV-2: Insights from a large healthcare database analysis</b>	Gendelman, Omer et al, Israel <a href="https://doi.org/10.1016/j.autrev.2020.102566">https://doi.org/10.1016/j.autrev.2020.102566</a>	Therapeutic	<p><b>Retrospective study based on a large healthcare computerized database</b> including all patients that were screened for the SARS-CoV-2 in the study period from February 23rd 2020 to March 31st 2020.</p> <p>Comparison between subjects tested positive for SARS-CoV-2 and those found negative in terms of rate of <b>administration of hydroxychloroquine/ colchicine therapy</b>.</p> <ul style="list-style-type: none"> <li>- An overall sample of <b>14,520 subjects</b> were screened for SARS-CoV-2 infection and <b>1317 resulted positive</b>.</li> <li>- <b>No significant difference was found in terms of rates of usage of hydroxychloroquine or colchicine</b> between those who were found positive for SARS-CoV-2 and those who were found negative (0.23% versus 0.25% for hydroxychloroquine, and 0.53% versus 0.48% for colchicine, respectively).</li> </ul>
Cell 05MAY2020	<b>Structural Basis for Potent Neutralization of Betacoronaviruses by Single-domain Camelid Antibodies</b>	Daniel Wrapp and al, <a href="https://doi.org/10.1016/j.cell.2020.04.031">https://doi.org/10.1016/j.cell.2020.04.031</a>	Therapeutic	<p>Isolation of single-domain antibodies (VHHs) from a llama immunized with prefusion-stabilized coronavirus spikes.</p> <p>These VHHs neutralize MERS-CoV or SARS-CoV-1 S pseudotyped viruses, respectively. Crystal structures of these VHHs bound to their respective viral targets reveal two distinct epitopes, but both VHHs interfere with receptor binding. Cross-reactivity between the SARS-CoV-1 S-directed VHH and SARS-CoV-2 S.</p> <p>Cross-reactive VHH neutralizes SARS-CoV-2 S pseudotyped viruses as a bivalent human IgG Fc-fusion.</p> <p>=&gt; These data provide a <b>molecular basis for the neutralization of pathogenic betacoronaviruses by VHHs</b> and suggest that these molecules may serve as useful therapeutics during coronavirus outbreaks.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature Communications 04MAY2020	<b>A human monoclonal antibody blocking SARS-CoV-2 infection</b>	Wang, Chunyan et al, <a href="https://www.nature.com/articles/s41467-020-16256-y">https://www.nature.com/articles/s41467-020-16256-y</a>	Therapeutic	First report of a (human) monoclonal antibody that neutralizes SARS-CoV-2 (and SARS-CoV) in cell culture. 47D11 binds a conserved epitope on the spike RBD explaining its ability to cross-neutralize SARS-CoV and SARS-CoV-2, using a mechanism that is independent of receptor-binding inhibition.
Intensive Care Med 04MAY2020	<b>High risk of thrombosis in patients with severe SARS-CoV-2 infection: a multicenter prospective cohort study</b>	Helms J, and al France <a href="https://doi.org/10.1007/s00134-020-06062-x">https://doi.org/10.1007/s00134-020-06062-x</a>	Clinic	Multicentric study – 4 ICU in France – 150 patients with ARDS Historical prospective cohort → comparison of COVID to non-COVID by propensity score matching <u>At baseline</u> : >95% patients elevated had D-dimer and fibrinogen Median age = 63 years 64/150 thrombotic complications and <b>16,7% pulmonary embolisms</b> <u>COVID19 (=77) vs non-COVID19 (=145)</u> : - More thrombotic complication in COVID19 (11,7 vs 2,1%, $p<0,008$ ) Thrombotic complications despite prophylactic or therapeutic anticoagulation → large number of patients still intubated → under-estimated → monitoring anticoagulant treatment/ higher targets than usual?
Nature 04MAY2020	<b>Effect of non-pharmaceutical interventions to contain COVID-19 in China</b>	Lai, Shengjie et al, <a href="https://www.nature.com/articles/s41586-020-2293-x">https://www.nature.com/articles/s41586-020-2293-x</a>	Public Health/Epidemiology	<b>Modelling framework that uses daily travel networks to simulate different outbreak and intervention scenarios across China</b> , using epidemiological and anonymised human movement data. → Total of 114,325 COVID-19 cases (interquartile range 76,776 - 164,576) estimated in mainland China as of February 29, 2020. → Without non-pharmaceutical interventions (NPIs), the COVID-19 cases would likely have shown a 67-fold increase (interquartile range 44 - 94) by February 29, 2020, with the effectiveness of different interventions varying. → The early detection and isolation of cases was estimated to have prevented more infections than travel restrictions and contact reductions, but combined NPIs achieved the strongest and most rapid effect. → The lifting of travel restrictions since February 17, 2020 does not appear to lead to an increase in cases across China if the social distancing interventions can be maintained, even at a limited level of 25% reduction on average through late April.
Cell host & microbe 04MAY2020	<b>Heightened innate immune responses in the respiratory tract of COVID-19 patients</b>	Zhou, Zhuo; et al. China <a href="https://doi.org/10.1016/j.chom.2020.04.017">https://doi.org/10.1016/j.chom.2020.04.017</a>	Immunology	Metatranscriptomic seq. to profile immune signatures in bronchoalveolar lavage fluid of 8 COVID-19, 146 community-acquired pneumonia patients, and 20 healthy controls :  - Proinflammatory gene expression, especially chemokines, markedly elevated in COVID-19 vs community-acquired pneumonia patients and healthy controls, <b>suggesting SARS-CoV-2 causes hypercytokinemia</b> . - SARS-CoV-2 triggered <b>robust expression of IFN-inducible genes (ISGs) with immunopathogenic potential</b> (overrepresentation of genes involved in inflammation), unlike SARS-CoV which is thought to induce inadequate IFN. - <b>Estimations of immune cell populations, show increased activated dendritic cells and neutrophils.</b>
Science 04MAY2020	<b>Site-specific glycan analysis of the SARS-CoV-2 spike</b>	Watanabe, Yasunori, et al. UK - USA <a href="https://doi.org/10.1126/science.abb9983">https://doi.org/10.1126/science.abb9983</a>	Structural biology	<b>Revealing the glycan structures on a recombinant SARS-CoV-2 spike (S) glycoprotein immunogen by site-specific mass spectrometry.</b>  -SARS-CoV-2 S gene encodes 22 N-linked glycan sequons per protomer, which likely play a role in protein folding and immune evasion.  → <b>Glycosylation analysis enables detailed mapping of the glycan-processing states and signatures across the trimeric viral spike.</b> <b>Glycan profiling have implications in viral pathobiology as well as vaccine design for comparing immunogen integrity and monitoring manufacturing processes .</b>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature 04MAY2020	<b>Rapid reconstruction of SARS-CoV-2 using a synthetic genomics platform</b>	Thao, Tran Thi Nhu, et al. Switzerland - Germany - Russia <a href="https://www.nature.com/articles/s41586-020-2294-9">https://www.nature.com/articles/s41586-020-2294-9</a>	Fundamental research	<p>Accelerated yeast-based reverse genetics pipeline can genetically reconstruct diverse long RNA viruses, including Coronaviridae, Flaviviridae and Paramyxoviridae families :</p> <ul style="list-style-type: none"> <li>- Viral subgenomic fragments (from viral isolates, cloned viral DNA, clinical samples, or synthetic DNA) are reassembled in one step in <i>S. cerevisiae</i> using transformation associated recombination (TAR) cloning to maintain the genome as a yeast artificial chromosome (YAC). T7-RNA polymerase then used to generate infectious RNA and viable virus.</li> </ul> <p>-&gt; <b>Approach to generate SARS-CoV-2 is rapid</b> (1 week after receipt of synthetic DNA fragments) <b>and applicable to other emerging RNA viruses, and can provide infectious virus to health authorities and diagnostic labs without the need of access to clinical samples.</b></p> <p>- <b>Also possible to rapidly introduce sequence variations to functionally characterize phenotypic consequences of SARS-CoV-2 evolution in real-time.</b></p>
Autoimmunity reviews 03MAY2020	<b>Tocilizumab for the treatment of severe COVID-19 pneumonia with hyperinflammatory syndrome and acute respiratory failure: A single center study of 100 patients in Brescia, Italy</b>	Toniati, Paola et al, Italy <a href="https://doi.org/10.1016/j.autrev.2020.102568">https://doi.org/10.1016/j.autrev.2020.102568</a>	Therapeutic	<p>A <b>prospective series of 100 consecutive patients</b> admitted to a Hospital in Brescia (Italy) between March 9th and March 20th with confirmed COVID-19 pneumonia and ARDS requiring ventilatory support <b>were administered Tocilizumab</b> (TCZ, monoclonal antibody that targets the interleukin 6 receptor).</p> <p>The outcome measure was an improvement in ARDS assessed by means of the Brescia COVID Respiratory Severity Score.</p> <p>Out of 100 treated patients (88 M, 12 F; median age: 62 years), the <b>respiratory condition was improved or stabilized in 77 (77%) patients</b>, of whom 61 showed a significant clearing of diffuse bilateral opacities on chest x-ray and 15 were discharged from the hospital. Respiratory condition worsened in 23 (23%) patients, of whom 20 (20%) died.</p> <p>All the patients presented with lymphopenia and high levels of C-reactive protein (CRP), fibrinogen, ferritin and interleukin 6 (IL-6) indicating a HIS. During the 10-day follow-up, three cases of severe adverse events were recorded.</p>
Immunity 03MAY2020	<b>Detection of SARS-CoV-2-specific humoral and cellular immunity in COVID-19 convalescent individuals</b>	Ling et al., China <a href="https://www.sciencedirect.com/science/article/pii/S1074761320301813">https://www.sciencedirect.com/science/article/pii/S1074761320301813</a>	Diagnostic	<ol style="list-style-type: none"> <li>1. SARS-CoV-2-specific antibodies are detected in COVID-19 convalescent subjects.</li> <li>2. Most COVID-19 convalescent individuals have detectable neutralizing antibodies.</li> <li>3. Cellular immune responses to SARS-CoV-2 are found in COVID-19 convalescent subjects</li> <li>4. Neutralization antibody titers correlate with the numbers of virus-specific T cells.</li> </ol>
American journal of obstetrics and gynecology 03MAY2020	<b>Evidence for and against vertical transmission for SARS-CoV-2 (COVID-19)</b>	A, Amouroux; et al. France <a href="http://www.sciencedirect.com/science/article/pii/S000293782030524X">http://www.sciencedirect.com/science/article/pii/S000293782030524X</a>	Clinic	<p>Observations from 12 articles published from 10 February to 4 April 2020 reporting on 68 deliveries and 71 neonates with maternal infection in the third trimester of pregnancy:</p> <ul style="list-style-type: none"> <li>- <b>SARS-CoV-2 recovered (RT-PCR) from</b> nasal and throat swabs, sputum and feces of symptomatic patients including <b>neonates but not from vaginal swabs, amniotic fluid, placenta, cord blood, neonatal blood or breast milk.</b></li> <li>- <b>Neonatal infection was diagnosed within 48 hours of life in 4 cases.</b></li> </ul> <p>-&gt; <b>More complete evidence and reliable serological studies needed</b> before counselling pregnant women on the risk of congenital infection with SARS-CoV-2.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Int. J. Infect. Dis. 03MAY2020	<b>Viral kinetics of SARS-CoV-2 in asymptomatic carriers and presymptomatic patients</b>	Kim, Seong Eun; et al. South Korea <a href="https://doi.org/10.1016/j.ijid.2020.04.083">https://doi.org/10.1016/j.ijid.2020.04.083</a>	Virology	<p>71 laboratory-confirmed SARS-CoV-2 cases, identified 3 presymptomatic patients and 10 entirely asymptomatic infections:</p> <ul style="list-style-type: none"> <li>- <b>2 out of 3 incubation period patients (presymptomatic) had very high viral titer (Ct value &lt;20).</b></li> <li>- <b>In entirely asymptomatic carriers : median days to first negative RT-PCR was 4.5 (2.5–9) days and all reached a first Ct&gt;35 RT-PCR within 14 days after diagnosis.</b></li> </ul> <p><b>-&gt; COVID-19 patients may already be infectious before symptoms manifestation, and 14 days isolation after diagnosis may be sufficient in entirely asymptomatic cases.</b></p>
Gastroenterology 01MAY2020	<b>Gastrointestinal and Hepatic Manifestations of 2019 Novel Coronavirus Disease in a Large Cohort of Infected Patients From New York: Clinical Implications</b>	Kaveh H and al USA <a href="https://doi.org/10.1053/j.gastro.2020.05.010">https://doi.org/10.1053/j.gastro.2020.05.010</a>	Clinic	<p>1059 patients COVID-19 - 33% at least one gastrointestinal symptom</p> <p><b>GI symptom:</b> diarrhea (22%) – abdominal pain (7%) – nausea (16%)</p> <p>62% had biochemical liver injury</p> <p>GI manifestation and liver injury were associated with higher admission rate</p> <p><b>Multivariate analysis</b> → independent predictor of death or ICU admission</p> <ul style="list-style-type: none"> <li>- Liver injury at presentation OR:2,53</li> <li>- Older age OR:1,03</li> <li>- Tachypnea OR:1,73</li> <li>- Severe hypoxemia OR:1,47</li> </ul> <p>GI manifestation → no effect on the outcome</p> <p><b>→ COVID-19 patients had commonly GI manifestation</b></p> <p><b>Cohort study of hospitalized patients with coronavirus disease 2019.</b></p> <p>Among <b>90 patients given hydroxychloroquine, 53 received concomitant azithromycin.</b> Those receiving concomitant azithromycin had a greater median change in QT interval compared with those receiving hydroxychloroquine.</p> <p>Seven patients (19%) who received hydroxychloroquine monotherapy developed prolonged QTc of 500 milliseconds or more, and 3 patients (3%) had a change in QTc of 60 milliseconds or more. Of those who received concomitant azithromycin, 11 of 53 (21%) had prolonged QTc of 500 milliseconds or more and 7 of 53 (13 %) had a change in QTc of 60 milliseconds or more. Ten patients had hydroxychloroquine discontinued early because of potential adverse drug events, including intractable nausea, hypoglycemia, and <b>1 case of torsades de pointes.</b></p>
JAMA Cardiology 01MAY2020	<b>Risk of QT Interval Prolongation Associated With Use of Hydroxychloroquine With or Without Concomitant Azithromycin Among Hospitalized Patients Testing Positive for Coronavirus Disease 2019 (COVID-19)</b>	Mercuro, Nicholas J. Et al, USA <a href="https://jamanetwork.com/journals/jamacardiology/fullarticle/2765631">https://jamanetwork.com/journals/jamacardiology/fullarticle/2765631</a>	Therapeutic	<p>The spike protein of SARS-CoV-2 harbors a multiple arginine residues (multibasic) S1/S2 site.</p> <ul style="list-style-type: none"> <li>- The host cell protease furin cleaves the SARS-CoV-2 spike protein at the S1/S2 site.</li> <li>- <b>Cleavage at the S1/S2 site is essential for spike-driven cell-cell fusion and viral entry into lung cells.</b></li> </ul> <p><b>-&gt; Suggests that acquisition of a S1/S2 multibasic cleavage site was essential for SARS-CoV-2 infection of humans and identify furin as a potential target for therapeutic intervention.</b></p>
Molecular Cell 01MAY2020	<b>A Multibasic Cleavage Site in the Spike Protein of SARS-CoV-2 Is Essential for Infection of Human Lung Cells</b>	Hoffmann, Markus; et al. Germany <a href="https://doi.org/10.1016/j.molcel.2020.04.022">https://doi.org/10.1016/j.molcel.2020.04.022</a>	Fundamental research	
New England Journal of Medicine 01MAY2020	<b>Cardiovascular Disease, Drug Therapy, and Mortality in Covid-19</b>	Mehra M, and al USA <a href="https://doi.org/10.1056/NEJMoa2007621">https://doi.org/10.1056/NEJMoa2007621</a>	Clinic	<b>RETRACTED</b>

Journal and date	Title	Authors and link	Field of expertise	Key facts
New England Journal of Medicine 01MAY2020	<b>Early Detection of Covid-19 through a Citywide Pandemic Surveillance Platform</b>	Chu, Helen Y. and al. USA <a href="https://www.nejm.org/doi/full/10.1056/NEJMc2008646">https://www.nejm.org/doi/full/10.1056/NEJMc2008646</a>	Public Health/Epidemiology	<p>The Seattle Flu Study is a multi-institutional, community-wide pandemic surveillance platform that was established in November 2018.</p> <p>-&gt; Persons enrolled online and were sent kits, by rapid-delivery services, for home collection of a midnasal swab; samples were returned by mail.</p> <p>-&gt;Persons reporting symptoms of respiratory illness provided informed consent for testing to identify influenza and other respiratory pathogens.</p> <p>-&gt;The first Covid-19 case detected through the Seattle Flu Study, in a specimen collected on February 24, 2020, was the first documented U.S. case of community transmission at the time.</p> <p><b>Conclusion:</b> widespread implementation of simple methods that are scalable and require minimal interaction for collection of samples from persons who may not seek clinical care is critical for early detection of community cases. An ubiquitous, community-based sampling for respiratory illnesses appears as an essential infrastructure for early detection and mitigation of future pandemics</p>
Clin. Infect. Dis 01MAY2020	<b>Early detection of SARS-CoV-2 antibodies in COVID-19 patients as a serologic marker of infection</b>	Zhao, Rongqing and al. <a href="https://doi.org/10.1093/cid/ciaa523">https://doi.org/10.1093/cid/ciaa523</a>	Diagnostic	<p><b>A COVID-19/SARS-CoV-2 S1 serology ELISA kit was developed.</b> The overall accuracy rate reached 97.3%. The assay was able to detect SARS-CoV-2 antibody on day one after the onset of COVID-19 disease. SARS-CoV-2 antibodies were detected in 28 out of 276 asymptomatic medical staff and one out of five nucleic acid test-negative “Close contacts” of COVID-19 patient.</p> <p><b>Conclusion:</b> the assays developed here can screen medical staff, in-coming patients, passengers and people who are in close contact with the confirmed patients to identify the “innocent viral spreaders”, protect the medical staff and stop the further spreading of the virus.</p>
J. Clin. Virol. 01MAY2020	<b>A RT-PCR Assay for the Detection of Coronaviruses from Four Genera</b>	Xiu, Leshan and al. China <a href="https://www.sciencedirect.com/science/article/pii/S1386653220301335">https://www.sciencedirect.com/science/article/pii/S1386653220301335</a>	Diagnostic	<p>A better understanding of the natural hosts and genetic diversity of CoVs are needed to help mitigate these threats.</p> <p>Objective: to design and evaluate a molecular diagnostic tool for detection and identification of all currently recognized and potentially future emergent CoVs from the Orthocoronavirinae subfamily.</p> <p><b>Conclusion:</b> a semi-nested, reverse transcription RT-PCR assay capable of detecting and identifying all previously recognized CoVs, including SARS-CoV-2, and potentially any emergent CoVs in this subfamily.</p>
Science 01MAY2020	<b>SARS-CoV-2 productively infects human gut enterocytes</b>	Lamers, Mart M.; et al. Netherlands <a href="https://doi.org/10.1126/science.abc1669">https://doi.org/10.1126/science.abc1669</a>	Fundamental research	<p>Infection of human small intestinal organoids (hSIOs) grown from primary gut epithelial stem cells to investigate intestine as another viral target organ :</p> <ul style="list-style-type: none"> <li>- hSIOs enterocytes were readily infected by SARS-CoV and SARS-CoV-2 (confocal- and electron-microscopy) and significant titers of infectious viral particles detected.</li> <li>- SARS-CoV-2 infected airway and gut organoids.</li> <li>- mRNA expression analysis revealed <b>strong induction of a generic viral response program. SARS-CoV-2 induced a stronger interferon response than SARS-CoV in HIOs.</b></li> </ul> <p>-&gt; <b>intestinal epithelium supports SARS-CoV-2 replication, and data imply that human organoids represent faithful experimental models to study of coronavirus infection and biology.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Emerging Microbes & Infections 01MAY2020	<b>Kinetics of SARS-CoV-2 specific IgM and IgG responses in COVID-19 patients</b>	Sun et al., China <a href="https://www.tandfonline.com/doi/full/10.1080/22221751.2020.1762515#">https://www.tandfonline.com/doi/full/10.1080/22221751.2020.1762515#</a>	Immuno	<ul style="list-style-type: none"> <li><b>Kinetic steps:</b> <ul style="list-style-type: none"> <li>- A total of 130 blood samples from 38 COVID-19 patients were analyzed.</li> <li>- Study showed that the seropositive rates of N-IgM, N-IgG, S-IgM and S-IgG antibody responses in non-ICU (intensive care unit) patients gradually increased within 1-3 weeks after the onset.</li> <li>- N-IgM and S-IgM reached a peak in the second week, while N-IgG and S-IgG antibodies continued to increase in the third week.</li> <li>- The joint detection of N-IgM, N-IgG, S-IgM, and S-IgG antibodies, could detect up to 75% of infections in the first week and the joint detection of N-IgM+N-IgG, or N-IgG+S-IgG could detect up to 94.7% of infections in the second week. Finally, in the third weeks after symptom onset, seropositive rates for N-IgG and S-IgG reached 100%.</li> </ul> </li> <li><b>Comparison between ICU and non-ICU patients</b> <ul style="list-style-type: none"> <li>- Most ICU patients had higher N-IgG than S-IgG after the symptom onset</li> <li>- ICU patients had SARS-CoV-2 nucleic acid positive days of 31.0, whereas non-ICU patients had SARS-CoV-2 nucleic acid positive days of 13. Therefore, a continuous increase of N-IgG may indicate disease progression towards more severe illness.</li> <li>- S-IgG in ICU patients was significantly lower than non-ICU patients by 2 weeks after the onset</li> </ul> </li> </ul> <p><b>Conclusion:</b> Intensive care unit monitoring the kinetics of S-IgG should help to predict prognosis.</p>
Science 01MAY2020	<b>Structural basis for inhibition of the RNA-dependent RNA polymerase from SARS-CoV-2 by remdesivir</b>	Yin, Wanchao; et al. China <a href="https://doi.org/10.1126/science.abc1560">https://doi.org/10.1126/science.abc1560</a>	Structural biology	<p>Cryo-EM structure of the SARS-CoV-2 RNA-dependent RNA polymerase (RdRp) in the apo form (unbound) (2.8 Å) or in complex with a 50-base template-primer RNA and the active form of Remdesivir (2.5 Å).</p> <ul style="list-style-type: none"> <li>- Structure comparison and sequence alignment suggest that mode of substrate RNA recognition and Remdesivir inhibition of RdRp is highly conserved in diverse RNA viruses</li> <li>-&gt; <b>providing a basis for designing broad spectrum antiviral drugs based on nucleotide analogs and a template structure for modeling/modifying existing nucleotide drugs (ex. EIDD-2801).</b></li> <li>-&gt; <b>Complex structure reveals the partial double-stranded RNA template is inserted into the central channel of the RdRp where Remdesivir is covalently incorporated into the primer strand at the first replicated base pair and terminates chain elongation.</b></li> </ul>
Gastroenterology 01MAY2020	<b>Taste Changes (Dysgeusia) in COVID-19: A systematic review and metaanalysis</b>	Aziz, Muhammad; et al. USA <a href="https://doi.org/10.1053/j.gastro.2020.05.003">https://doi.org/10.1053/j.gastro.2020.05.003</a>	Clinic	<p>Systematic review (case series/case-control/ cohort studies) (January 1st - April 21st, 2020) reporting on ageusia/dysgeusia:</p> <ul style="list-style-type: none"> <li>- 4 single-nation studies, 1 multinational study from Europe = total of 817 patients included.</li> <li>-&gt; <b>Prevalence : almost half of patients (49.8%) with COVID-19 have altered taste sensation across the five studies.</b></li> </ul> <p><b>Limitations :</b> lack of data comparing ageusia/dysgeusia in COVID-19 +ve vs -ve patients, or severe COVID-19.</p> <p>In contexts of lack of diagnostic tests (ex. developing world), <b>distinctive clinical features like ageusia/dysgeusia can be useful in identifying suspected COVID-19 patients.</b></p>
New England Journal of Medicine 01MAY2020	<b>Renin–Angiotensin–Aldosterone System Inhibitors and Risk of Covid-19</b>	Reynolds H, et al USA <a href="https://doi.org/10.1056/NEJMoa2008975">https://doi.org/10.1056/NEJMoa2008975</a>	Clinic	<p>Relation between previous treatment that act on the RAAS and the likelihood of a positive test or the likelihood of severe illness?</p> <p>Five class of antihypertensive medication examined. Estimated a propensity score for the likelihood of treatment with each medication class</p> <ul style="list-style-type: none"> <li>-&gt; 12594 patients were tested</li> <li>-&gt; 5894 patients positive COVID19 which 17% had severe illness</li> <li>-&gt; 2573(/5894) had HTA which 24,6% had severe illness</li> </ul> <p><b>No association between medication examined and increased likelihood of a positive test or in the risk of severe Covid19.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
NEJM 01MAY2020	<b>Inhibitors of the Renin–Angiotensin–Aldosterone System and Covid-19</b>	Jarcho, John A.; et al. USA <a href="https://doi.org/10.1056/NEJMe2012924">https://doi.org/10.1056/NEJMe2012924</a>  Mehra MR, et al. USA <a href="https://doi.org/10.1056/NEJMoa2007621">https://doi.org/10.1056/NEJMoa2007621</a>  Mancia, Giuseppe; et al. Italy <a href="https://doi.org/10.1056/NEJMoa2006923">https://doi.org/10.1056/NEJMoa2006923</a>  Reynolds HR, et al. USA <a href="https://doi.org/10.1056/NEJMoa2008975">https://doi.org/10.1056/NEJMoa2008975</a>	Clinic	<p>Analysis of 3 <i>N Engl J Med</i> articles, as clinicians weigh alleged harm of continuing RAAS inhibitor medications (like ACE inhibitors and angiotensin-receptor blockers (ARBs)) often prescribed to patient with hypertension, diabetes, and coronary artery disease (high risk groups for severe COVID-19):</p> <ul style="list-style-type: none"> <li>- Mehra et al. database study, 8910 hospitalised Covid-19 patients, 11 countries : <b>ACE inhibitors nor ARBs associated with increased risk in-hospital death. Secondary analysis restricted to hypertension patients (for whom ACE inhibitor or ARB would be indicated) also did not show harm.</b></li> <li>- Mancia et al. case–control study, 6272 confirmed SARS-CoV-2 patients in Lombardy (Italy) vs 30,759 matched controls : <b>ACE inhibitors nor ARBs associated with likelihood of SARS-CoV-2 infection. In severe/fatal infections vs matched controls, no association between these drugs and severe Covid-19.</b></li> <li>- Reynolds et al. electronic health records, 12,594 people in New York University -&gt; 5894 tested +ve, of which 1002 had severe illness (admission to ICU/mechanical ventilation/death): <b>no +ve association for drug classes, ACE inhibitors and ARBs, for a +ve test result or severe illness.</b></li> </ul> <p><b>-&gt; none of the 3 studies showed evidence of harm with continued use of ACE inhibitors and ARBs.</b></p>
JAMA Cardiology 30APR2020	<b>Assessment of QT Intervals in a Case Series of Patients With Coronavirus Disease 2019 (COVID-19) Infection Treated With Hydroxychloroquine Alone or in Combination With Azithromycin in an Intensive Care Unit</b>	Bessiere, Francis et al, France <a href="https://jamanetwork.com/journals/jamacardiology/fullarticle/2765633">https://jamanetwork.com/journals/jamacardiology/fullarticle/2765633</a>	Therapeutic	<p>40 consecutive patients with COVID-19 confirmed by positive RT-PCR results on respiratory samples admitted to the ICU who received hydroxychloroquine with or without were included.</p> <ul style="list-style-type: none"> <li>-&gt; 30 patients (75%) required invasive mechanical ventilation and 25 (63%) received vasoactive drugs.</li> <li>-&gt; Hydroxychloroquine with or without azithromycin was given to 18 (45%) and 22 patients (55%), respectively. 20 patients (50%) also received other treatments favoring QT prolongation in the ICU.</li> <li>⇒ Most patients (37 [93%]): increase in QTc. Prolonged QTc was observed in 14 patients (36%) after a duration of antiviral treatment of 2 to 5 days. No ventricular arrhythmia, including torsades de pointes.</li> <li>⇒ Among patients treated with hydroxychloroquine and azithromycin, 6 of 18 (33%) developed an increase in QTc of 500 milliseconds or greater vs 1 of 22 (5%) of those treated with hydroxychloroquine alone (P = .03).</li> <li>⇒ The antiviral treatment ceased before completion for 7 patients (17.5%) following ECG abnormalities and in 10 (25%) for acute renal failure.</li> </ul>
Nature 30APR2020	<b>A SARS-CoV-2 protein interaction map reveals targets for drug repurposing</b>	Gordon, David E et al, <a href="https://www.nature.com/articles/s41586-020-2286-9#Abs1">https://www.nature.com/articles/s41586-020-2286-9#Abs1</a>	Therapeutic	<p><b>26 of the 29 SARS-CoV-2 proteins in human cells were cloned, tagged and expressed, and the human proteins physically associated with each were identified</b> using affinity-purification mass spectrometry (AP-MS)</p> <ul style="list-style-type: none"> <li>-&gt; Identification of 332 high-confidence SARS-CoV-2-human protein-protein interactions (PPIs).</li> <li>-&gt; Among these, 66 druggable human proteins or host factors targeted by 69 compounds (29 FDA-approved drugs, 12 drugs in clinical trials, and 28 preclinical compounds).</li> <li>-&gt; Screening a subset of these in multiple viral assays identified <b>two sets of pharmacological agents that displayed antiviral activity</b>: inhibitors of mRNA translation and predicted regulators of the Sigma1 and Sigma2 receptors. Further studies of these host factor targeting agents, including their combination with drugs that directly target viral enzymes, could lead to a therapeutic regimen to treat COVID-19.</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Jama Psychiatry 30APR2020	<b>Mental Health in the Coronavirus Disease 2019 Emergency—The Italian Response</b>	De Girolamo et al., Italy <a href="https://jamanetwork.com/journals/jamapsychiatry/fullarticle/2765557">https://jamanetwork.com/journals/jamapsychiatry/fullarticle/2765557</a>	Psy	<b>Experience of mental health services and the lessons learned in Italy</b> -> During epidemic: challenges have occurred in the management of health services -> Psychiatric wards have been reorganized to admit patients with COVID-19, and many physicians and nurses have been diverted to wards managing patients with COVID-19 -> Facilities for patients with psychiatric needs have been temporarily closed -> Patient confined in the facilities with very limited or no leave Considerable stresses <b>Need for a leadership position in the psychosocial management of disasterlike situations.</b>
Lancet 30APR2020	<b>Obesity could shift severe COVID-19 disease to younger ages</b>	Kass, David A.; et al. USA <a href="https://doi.org/10.1016/S0140-6736(20)31024-2">https://doi.org/10.1016/S0140-6736(20)31024-2</a>	Clinic	Correlation of body-mass index (BMI) vs age in COVID-19 patients admitted to ICU at 5 US university hospitals, 265 patients (58% male) : - <b>significant inverse correlation between age and BMI -&gt; younger individuals admitted to hospital more likely obese.</b> - no difference by sex -> <b>In populations with high prevalence of obesity, COVID-19 will affect younger populations more than previously reported.</b>
Viruses 30APR2020	The SARS-CoV-2 Exerts a Distinctive Strategy for Interacting with the ACE2 Human Receptor	Brielle, Esther S.; et al. Israel <a href="https://doi.org/10.3390/v12050497">https://doi.org/10.3390/v12050497</a>	Computational Structural Biology	Using molecular dynamics simulations to compare interaction between human ACE2 receptor and spike protein (SARS-CoV, SARS-CoV-2, and HCoV-NL63): - SARS-CoV and SARS-CoV-2 have comparable binding affinities achieved by balancing energetics and dynamics.  -> <b>SARS-CoV-2–ACE2 complex -&gt; higher number of interacting residues larger, larger interface area, decreased interface residue fluctuations relative to the SARS-CoV–ACE2 complex.</b>  -> <b>Data implies therapeutic challenge attributed to the enhanced rigidity of the COVID-19 RBD relative to that of SARS-2002.</b>
Cell 30APR2020	<b>Genomic Epidemiology of SARS-CoV-2 in Guangdong Province, China</b>	Lu, Jing; et al. China <a href="http://www.sciencedirect.com/science/article/pii/S0092867420304864">http://www.sciencedirect.com/science/article/pii/S0092867420304864</a>	Phylogenetic analysis	Extensive early surveillance in densely populated Guangdong, China's, 1.6 million tests from 30 Jan - 19 March, resulted in 1,388 reported RNA-positive SARS-CoV-2. 53 genomes generated from infected individuals (combination of metagenomic sequencing and tiling amplicon approaches) indicated:  - <b>most infections were due to virus importation to Guangdong, and chains of local transmission limited in size and duration, (clustering uncertain due to low virus genetic variation early in the pandemic).</b>  -> <b>national travel restrictions and province's large-scale intensive surveillance and intervention measures helped reduce/ eliminate transmission chains.</b> - vigilance still required following recent increase in COVID-19 cases imported to China from other countries.
Nature 29APR2020	<b>Massively multiplexed nucleic acid detection using Cas13</b>	Ackerman, Cheri M. and al. USA <a href="http://www.ncbi.nlm.nih.gov/pubmed/32349121">http://www.ncbi.nlm.nih.gov/pubmed/32349121</a>	Diagnostic	Combinatorial Arrayed Reactions for Multiplexed Evaluation of Nucleic acids (CARMEN), a platform for scalable, multiplexed pathogen detection. The combination of CARMEN and Cas13 detection allowed to develop a multiplexed assay that simultaneously differentiates all 169 human-associated viruses with ≥10 published genome sequences and rapidly incorporated an additional crRNA to detect the causative agent of the 2020 COVID-19 pandemic.  CARMEN-Cas13 further enables comprehensive subtyping of influenza A strains and multiplexed identification of dozens of HIV drug-resistance mutations.  <b>Conclusion:</b> Scalable, highly-multiplexed CRISPR-based nucleic acid detection shifts diagnostic and surveillance efforts from targeted testing of high-priority samples to comprehensive testing of large sample sets.



Journal and date	Title	Authors and link	Field of expertise	Key facts
International journal of antimicrobial agents 29APR2020	<b>Global coronavirus disease 2019: what has daily cumulative index taught us?</b>	Lai C, and al China <a href="http://www.sciencedirect.com/science/article/pii/S092485792030159X">http://www.sciencedirect.com/science/article/pii/S092485792030159X</a>	Public Health/Epidemiology	<p>Rapid increase of COVID19 cases in short time = insufficiency of healthcare system and negatively affect patient's outcome?</p> <p><b>Daily cumulative index (DCI)</b> = cumulative cases/number of days between the first reported case and March 6. DCI significantly positively correlated with incidence (aRR:1,01 [1,00 – 1,02]) After adjustment of HCl or HAQI, DCI associated with mortality per 1,000,000 (aRR:1,02 [1,01 – 1,03])</p> <p>Higher level of healthcare performance is associated with higher incidence → ability to detect?</p> <p>→ Reduction of DCI:</p> <ul style="list-style-type: none"> <li>- slow the increasing number of COVID19</li> <li>- improve outcome in COVID-19 patients</li> </ul>
Science 29APR2020	<b>Changes in contact patterns shape the dynamics of the COVID-19 outbreak in China</b>	Zhang J, and al China <a href="https://doi.org/10.1126/science.abb8001">https://doi.org/10.1126/science.abb8001</a>	Public Health/Epidemiology	<p>Analysis of contact surveys data in Wuhan and Shanghai before and during the outbreak → construction of a model Social distancing → ↓ 7-8-fold daily contact / most interaction restricted to household</p> <p>The model consider potential age differences in susceptibility to infection <u>Susceptibility to SARS-CoV-2 infection increase with age:</u></p> <ul style="list-style-type: none"> <li>- 0-14 years: OR=0,34 [0,24 – 0,49] (compared to 15-64y)</li> <li>- &gt; 65 years: OR=1,47 [1,12 – 1,92] (compared to 15-64y)</li> </ul> <p><u>Model to study the impact of social distancing and school closures:</u></p> <ul style="list-style-type: none"> <li>- social distancing alone is sufficient to control the outbreak</li> <li>- School closures → reduce peak incidence (40-60%) and delay the epidemic / impact on the disease dynamic and hospital surge capacity</li> </ul> <p>→ Refining age-specific estimates of susceptibility to infection → to evaluating the impact of interventions put in place.</p>
PNAS 29APR2020	<b>Effective treatment of severe COVID-19 patients with tocilizumab</b>	Xu, Xiaoling et al, China <a href="https://doi.org/10.1073/pnas.2005615117">https://doi.org/10.1073/pnas.2005615117</a>	Therapeutic	<p><b>Retrospective study with tocilizumab</b>, an IL6R inhibitor, in treatment of 21 patients with severe and critical COVID-19.</p> <p><b>Clinical data showed that the symptoms, hypoxymia, and CT opacity changes were improved immediately after the treatment with tocilizumab</b> in most of the patients, suggesting that tocilizumab could be an efficient therapeutic for the treatment of COVID-19. No obvious adverse reactions were observed.</p> <p>Limitations: limited number of patients, single observation study.</p>
The Lancet 29APR2020	<b>Remdesivir in adults with severe COVID-19: a randomised, double-blind, placebo-controlled, multicentre trial</b>	Wang et al., China <a href="https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)31022-9/fulltext?utm_campaign=tlc-coronavirus20&amp;utm_source=twitter&amp;utm_medium=social">https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)31022-9/fulltext?utm_campaign=tlc-coronavirus20&amp;utm_source=twitter&amp;utm_medium=social</a>	Therapeutic	<p>-&gt; 237 patients were enrolled and randomly assigned to a treatment group (158 to remdesivir and 79 to placebo)</p> <p>-&gt; Remdesivir use was not associated with a difference in time to clinical improvement (hazard ratio 1.23 [95% CI 0.87–1.75])</p> <p>-&gt; Although not statistically significant, patients receiving remdesivir had a numerically faster time to clinical improvement than those receiving placebo among patients with symptom duration of 10 days or less (hazard ratio 1.52 [0.95–2.43])</p> <p>-&gt; Adverse events were reported in 102 (66%) of 155 remdesivir recipients versus 50 (64%) of 78 placebo recipients. Remdesivir was stopped early because of adverse events in 18 (12%) patients versus four (5%) patients who stopped placebo early.</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
J Infect Dis 29APR2020	<b>Effect of Convalescent Plasma Therapy on Viral Shedding and Survival in COVID-19 Patients</b>	Zeng, Qing-Lei et al, China <a href="https://doi.org/10.1093/infdis/jiaa228">https://doi.org/10.1093/infdis/jiaa228</a>	Therapeutic	<p>6 COVID-19 subjects with respiratory failure received <b>convalescent plasma</b> at a median of <b>21.5 days</b> after first detection of viral shedding.</p> <p>All tested <b>negative for SARS-CoV-2 RNA by 3 days after infusion</b>, and <b>5 died eventually</b>.</p> <p>In conclusion, convalescent plasma treatment can discontinue SARS-CoV-2 shedding but cannot reduce mortality in critically end-stage COVID-19 patients, and treatment should be initiated earlier.</p>
Cancer discovery 28APR2020	<b>Patients with cancer appear more vulnerable to SARS-COV-2: a multi-center study during the COVID-19 outbreak</b>	Dai M, and al USA <a href="https://doi.org/10.1158/2159-8290.CD-20-0422">https://doi.org/10.1158/2159-8290.CD-20-0422</a>	Clinic	<p>Patients with cancer are more vulnerable to infections 195 and 536 patients with and without cancer respectively matched.</p> <p>More in-hospital infection and smoking history in cancer group</p> <p><u>Patient with cancer had higher:</u></p> <ul style="list-style-type: none"> <li>- <b>ICU admission</b> [OR:2,84 (1,59 – 5,08)]</li> <li>- <b>Death rate</b> [OR:2,34 (1 15 – 4,77)]</li> <li>- <b>Having one or more severe/critical symptom</b> [OR:2,79 (1,74 – 4,41)]</li> <li>- <b>Changes of needing MV</b></li> </ul> <p>Hematological, lung, metastatic cancer → higher rates of severe events</p> <p>→ <b>patients with cancer tend to have more severe outcomes</b></p>
Critical Care 28APR2020	<b>Clinical determinants for fatality of 44,672 patients with COVID-19</b>	Deng G, and al China <a href="https://doi.org/10.1186/s13054-020-02902-w">https://doi.org/10.1186/s13054-020-02902-w</a>	Clinic Letter	<p>Meta-analysis, confirmed cases series in China of 44672 patients</p> <p>Mortality rate all age → 2,3%</p> <p>CFR increase with age to 14,8 % in patients &gt; 80 years</p> <p><u>Risk factors for fatality:</u></p> <ul style="list-style-type: none"> <li>- Male [RR:1,67 (1,47 – 1,89)]</li> <li>- Cardiovascular disease [RR:6,75 (5,4 – 8,43)]</li> <li>- HTA [HR:4,48 (3,69 – 5,85)]</li> <li>- Diabetes [RR:4,43 (3,49 – 5,61)]</li> <li>- Cancer [RR:2,92 (1,34 – 6,41)]</li> <li>- Respiratory disease [RR:3,43 (2,42 – 4,87)]</li> </ul> <p><b>More intensive surveillance for male and those with comorbidities</b></p>
Natl Sci Rev 28APR2020	<b>Plasma Metabolomic and Lipidomic Alterations Associated with COVID-19</b>	Di Wu; Shu, et al. China <a href="https://doi.org/10.1093/nsr/nwaa086">https://doi.org/10.1093/nsr/nwaa086</a>	Metabolite and lipid alterations	<p><b>Targeted metabolomic and lipidomic analyses of plasma from cohort of COVID-19 patients to identify plasma biomarkers associated with COVID-19:</b></p> <ul style="list-style-type: none"> <li>- malic acid of the TCA cycle and carbamoyl phosphate of urea cycle <b>reveal altered energy metabolism and hepatic dysfunction</b>, respectively.</li> <li>- <b>carbamoyl phosphate profoundly down-regulated in fatal patients compared with mild patients.</b></li> <li>- <b>guanosine monophosphate (GMP)</b> (mediated by GMP synthase, CD39 and CD73) <b>significantly altered between healthy subjects vs COVID-19 patients, and between the mild vs fatal groups.</b></li> </ul>
Public Health 28APR2020	<b>Examining the Effect of Social Distancing on the Compound Growth Rate of SARS-CoV-2 at the County Level (United States) Using Statistical Analyses and a Random Forest Machine Learning Model</b>	Cobb J and al, USA <a href="http://www.sciencedirect.com/science/article/pii/S0033350620301219">http://www.sciencedirect.com/science/article/pii/S0033350620301219</a>	Public Health/Epidemiology	<p>Trends among US counties and COVID 19 growth rate in relation to existence of shelter in place (SIP) orders <b>Machin learning</b></p> <p><b>Limiting gatherings to &lt; 10 people reduced growth rate by 6,6%</b></p> <p><b>SIP → reduction of 7,8% versus counties with no SIP</b> <b>SIP orders and limitation gathering were additive</b></p> <p>Features predicting the effect of SIP:</p> <ul style="list-style-type: none"> <li>- Population / Longitude / Population per square</li> </ul> <p>→ SIP was effective → Counties with large population or high population density: benefit the most from a SIP</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Cell 28APR2020	<b>Trained immunity: a tool for reducing susceptibility and severity of SARS-CoV-2 infection</b>	Neteal et al., The Netherlands <a href="https://www.cell.com/pb-assets/products/coronavirus/CELL_11391_S5.pdf">https://www.cell.com/pb-assets/products/coronavirus/CELL_11391_S5.pdf</a>	Fundamental research	<p>-&gt; Long-term boosting of innate immune responses, also termed 'trained immunity', by certain live vaccines (BCG, oral polio vaccine, measles) induces heterologous protection against infections, through epigenetic, transcriptional and functional reprogramming of innate immune cells.</p> <p><b>-&gt; We propose that induction of trained immunity by whole microorganism vaccines may represent an important tool for reducing susceptibility and severity to SARS-CoV-2.</b></p>
Cell PreProof	<b>Clinically Applicable AI System for Accurate Diagnosis, Quantitative Measurements and Prognosis of COVID-19 Pneumonia Using Computed Tomography</b>	Zhang et al., China <a href="https://www.cell.com/pb-assets/products/coronavirus/CELL_CELL-D-20-00656.pdf">https://www.cell.com/pb-assets/products/coronavirus/CELL_CELL-D-20-00656.pdf</a>	Diagnostic	<p>Using a large computed Tomography (CT) database from 4,154 patients, we developed an AI system that can diagnose NCP and differentiate it from other common pneumonia and normal controls.</p> <p>-&gt; Identification of important clinical and biochemical markers of multiple organs that correlated with the NCP lesion properties</p> <p>-&gt; Together with the clinical data, this AI system was able to provide accurate clinical prognosis</p>
Science Advances 27APR2020	<b>Squalene-based multidrug nanoparticles for improved mitigation of uncontrolled inflammation</b>	Dormont, Flavio et al, France <a href="https://doi.org/10.1126/sciadv.aaz5466">https://doi.org/10.1126/sciadv.aaz5466</a>	Therapeutic	<p>Development of <b>multidrug nanoparticles for the mitigation of uncontrolled inflammation</b>.</p> <p>The nanoparticles are made by conjugating squalene, an endogenous lipid, to adenosine, an endogenous immunomodulator, and then encapsulating <math>\alpha</math>-tocopherol, a natural antioxidant. This resulted in <b>high drug loading, biocompatible, multidrug nanoparticles</b>.</p> <p>By exploiting the vascular endothelial barrier dysfunction at sites of acute inflammation, these <b>multidrug nanoparticles could deliver the therapeutic agents in a targeted manner</b> and conferred a significant survival advantage to treated animals in lethal models of endotoxemia.</p> <p><b>-&gt; Selectively delivering adenosine and antioxidants together could serve as a novel approach for the treatment of acute inflammation with reduced-side effects and high therapeutic potential.</b></p>
European Journal of Clinical Microbiology & Infectious Diseases 27APR2020	<b>Viral RNA load as determined by cell culture as a management tool for discharge of SARS-CoV-2 patients from infectious disease wards</b>	La Scola, Bernard and al. France <a href="https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7185831/">https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7185831/</a>	Virology	<p>It is of paramount importance to define when a treated patient can be considered as no longer contagious.</p> <p>Correlation between successful isolation of virus in cell culture and Ct value of quantitative RT-PCR targeting E gene suggests that patients with Ct above 33–34 using our RT-PCR system are not contagious and thus can be discharged from hospital care or strict confinement for non-hospitalized patients.</p>
Microbial Pathogenesis 27APR2020	<b>Design of a peptide-based subunit vaccine against novel coronavirus SARSCoV-2</b>	Parismita Kalita et al., India-Japan <a href="http://www.sciencedirect.com/science/article/pii/S0882401020305234">http://www.sciencedirect.com/science/article/pii/S0882401020305234</a>	Vaccines	<p>Multi-peptide subunit-based epitope vaccine against COVID-19 containing and adjuvant, CTL, HTL and B Cell epitopes</p> <p>-&gt; Epitopes were selected from known SARS-CoV-2 antigenic proteins (Nucleocapside, membrane, Spike) by using <i>in silico</i> prediction tools.</p> <p>-&gt; Identification of 6 high immunogenic epitopes targeting HTL, 18 targeting CTL, 9 targeting B-cells.</p> <p>- Epitopes were linked together to build a 566 aa subunit vaccine</p> <p>- Human b-defensine 1 (68aa): as adjuvant</p> <p>Extensive bioinformatics analysis suggest that the vaccine is immunogenic, non-toxic, non-allergenic, thermostable, with the capability to elicit a humoral and cell-mediated immune response.</p> <p>The binding modes, dynamics, and stability of the vaccine-TLR3 complex were validated by using molecular dynamics simulation studies.</p> <p>Estimation of the half-life of the vaccine :</p> <ul style="list-style-type: none"> <li>- 30 h in mammalian reticulocytes (in vitro)</li> <li>- &gt; 20 h in yeast (in vivo)</li> <li>- &gt; 10 h in E. coli (in vivo), suggesting that the construct is stable in vivo</li> </ul> <p>-&gt; Probability of showing good protective efficacy and safety against SARS-CoV-2 infection in humans</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Eurosurveillance 27APR2020	<b>Coronavirus disease (COVID-19) – impact on vaccine preventable diseases</b>	Hungerdord et al., UK <a href="https://doi.org/10.2807/1560-7917.ES.2020.25.18.2000756">https://doi.org/10.2807/1560-7917.ES.2020.25.18.2000756</a>	HSS/Politic	<p>COVID-19 has caused an unintentional drop of the European's delivery of routine immunisations of 2020 birth cohorts + catch up campaigns+ immunisation of at risk groups because of:</p> <ol style="list-style-type: none"> <li>1. Isolation + COVID-19 illness in families with newborn children</li> <li>2. Disruption of vaccine supplies</li> <li>3. Healthcare staffing issues</li> <li>4. Difficulty to launch a prevention campaign for adults at risk</li> </ol> <p><b>Impact:</b></p> <ul style="list-style-type: none"> <li>• Increase in number of sensitive children facing winter illnesses</li> <li>• The poorest populations are disproportionately affected</li> <li>• Risk of outbreaks of diseases such as measles...</li> </ul> <p><b>To deal with this impact,</b></p> <ul style="list-style-type: none"> <li>- ensure consideration + resources for delivery of routine vaccination</li> <li>- Vaccination rates should be monitored by analysts and immunization teams to identify hotspots = vaccine coverage surveillance + modeling.</li> </ul> <p>This will help face and not allow the Covid to increase the disability and mortality from vaccine preventable diseases.</p>
The Lancet. Infectious diseases 27APR2020	<b>Epidemiology and transmission of COVID-19 in 391 cases and 1286 of their close contacts in Shenzhen, China: a retrospective cohort study</b>	Qifang B and al, China/USA <a href="http://www.sciencedirect.com/science/article/pii/S1473309920302875">http://www.sciencedirect.com/science/article/pii/S1473309920302875</a>	Public Health/Epidemiology	<p>391 cases and 1286 close contact</p> <p>Cases were older - Most case mild or moderate and 9% severe</p> <p>Median incubation period: 4,8 days</p> <p>All those develop symptoms will do within 14 days</p> <p>Median time recovery: 20,8 days</p> <p>Contact tracing reduced isolation period by 1,9 days</p> <p>Higher risk of infection:</p> <ul style="list-style-type: none"> <li>- Household contact (OR6,27)</li> <li>- Contact travelling with a case (OR:7,06)</li> </ul> <p>Secondary attack = 11,2%</p> <p>Children likely to be infected (7,4%) than adults (6,6%)</p> <p>→ Isolation and contact tracing reduce the R and time during which cases are infectious</p> <p>→ children similar risk → analyses for transmission and control</p>
Inter J Infectious Disease 27APR2020	<b>Coronavirus disease 2019 in pregnancy</b>	Xu Q and al, China <a href="http://www.sciencedirect.com/science/article/pii/S1201971220302800">http://www.sciencedirect.com/science/article/pii/S1201971220302800</a>	Clinic	<p>Pregnant woman (28) compared to non-pregnant woman (54)</p> <p><b>Time from illness to admission: shorter pregnant women</b></p> <p><u>Laboratory:</u> significantly</p> <ul style="list-style-type: none"> <li>- Higher leukocyte in pregnant women (10 vs 2 x10<sup>9</sup>/L)</li> <li>- Higher CRP (17 vs 14 mg/dl)</li> </ul> <p>75% pregnant received antiviral vs 100% non-pregnant</p> <p><b>No association between</b></p> <ul style="list-style-type: none"> <li>- pregnancy and virus clearance time</li> <li>- pregnancy and LOS</li> <li>- pregnancy and severity of disease</li> </ul> <p>Median gestational age: 38 [IQR:36,5 – 39]</p> <p><b>None of neonates had a positive result for SARS-CoV-2</b></p> <p>→ no vertical transmission – good outcomes for both group</p>
Pathogens 26APR2020	<b>Emergence of Drift Variants That May Affect COVID-19 Vaccine Development and Antibody Treatment</b>	Takahiko Koyama et al., USA <a href="https://doi.org/10.3390/pathogens9050324">https://doi.org/10.3390/pathogens9050324</a>	Vaccines	<p>&gt; Twenty-one virus variants affecting T-cell epitopes identified.</p> <p>&gt; Twelve virus variants affecting B-cell epitopes of spike protein (S), nucleocapsid protein (N), and membrane protein (M) identified</p> <p>A variant replacing 23403A&gt;G in the S protein B-Cell epitope has been frequently observed in European countries (Netherlands, Switzerland, and France) although it is not yet known if it is the predominant sub-strain.</p> <p>This change involves a substitution of a large acidic residue D (aspartic acid) into small hydrophobic residue G (glycine), meaning differences in both size and hydrophobicity in the middle of the epitope and compromising the action of vaccines against wt S protein&gt; Genetic drift</p> <p>&gt; Vaccine design and convalescent plasma antibody treatment might require specific considerations to accommodate the drift</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet. Infectious diseases 27APR2020	<b>Epidemiology and transmission of COVID-19 in 391 cases and 1286 of their close contacts in Shenzhen, China: a retrospective cohort study</b>	Qifang B and al, China/USA <a href="http://www.sciencedirect.com/science/article/pii/S1473309920302875">http://www.sciencedirect.com/science/article/pii/S1473309920302875</a>	Public Health/Epidemiology	<p><b>391 cases and 1286 close contact</b></p> <p>Cases were older - Most case mild or moderate and 9% severe Median incubation period: 4,8 days All those develop symptoms will do within 14 days Median time recovery: 20,8 days Contact tracing reduced isolation period by 1,9 days Higher risk of infection:</p> <ul style="list-style-type: none"> <li>- Household contact (OR6,27)</li> <li>- Contact travelling with a case (OR:7,06)</li> </ul> <p>Secondary attack = 11,2% Children likely to be infected (7,4%) than adults (6,6%)</p> <p>→ Isolation and contact tracing reduce the R and time during which cases are infectious → children similar risk → analyses for transmission and control</p>
An International Journal of Obstetrics & Gynaecology 27APR2020	<b>Vaginal delivery in SARS-CoV-2 infected pregnant women in Northern Italy: a retrospective analysis</b>	Ferrazzi E and al, Italy <a href="https://doi.org/10.1111/1471-0528.16278">https://doi.org/10.1111/1471-0528.16278</a>	Clinic	<p>42-woman COVID-19 who delivered during study period <u>Diagnosis COVID:</u> 10 before delivered / 27 in delivery room /5 within 36h after 19 woman diagnosed pneumonia which 4 admitted ICU 24 cases vaginal delivery – 10 cesareans related to COVID19 <b>Cesareans related to pneumonia (p=0,024)</b> <b>3 neonates positive for SARS-CoV:</b></p> <ul style="list-style-type: none"> <li>- 2 newborns of women diagnosed after delivery</li> <li>- 1 newborn after vaginal delivery: gastrointestinal and respiratory symptoms → ICU with 24h of MV</li> </ul> <p>→ vaginal delivery is appropriated → cesarean: women with sever symptoms</p>
Clinical microbiology and infection 25APR2020	<b>Umifenovir treatment is not associated with improved outcomes in patients with coronavirus disease 2019: A retrospective study</b>	Lian, Ningfang et al, China <a href="http://www.sciencedirect.com/science/article/pii/S1198743X20302342">http://www.sciencedirect.com/science/article/pii/S1198743X20302342</a>	Therapeutic	<p><b>Retrospective study</b>, 81 COVID-19 patients included, with 45 in umifenovir group and 36 in control group. Baseline clinical, laboratory characteristics were comparable between two groups.</p> <p><b>Umifenovir treatment did not shorten the negativity time of SARS-CoV-2, or the length of hospital stay in non-ICU hospitalized patients with COVID-19.</b> No severe side effect was found in umifenovir treatment.</p> <p><b>Limitations :</b> single center, retrospective study with a small sample size ; pharyngeal swabs were not collected every day due to the limited medical resources, and pathogenic nucleic acids were not quantified as well ; only included patients with moderate and severe COVID-19, so the effectiveness of umifenovir in mild and critical patients cannot be confirmed in this study</p>
Cell 24APR2020	<b>SARS-CoV-2 receptor ACE2 is an interferon-stimulated gene in human airway epithelial cells and is detected in specific cell subsets across tissues</b>	Ziegler et al., USA <a href="https://www.cell.com/cell/fulltext/S0092-8674(20)30500-6">https://www.cell.com/cell/fulltext/S0092-8674(20)30500-6</a>	Fundamental research	<p>-&gt; Meta-analysis of human, primate &amp; mouse scRNA-seq for putative SARS-CoV-2 targets -&gt; Type II pneumocytes, nasal secretory cells &amp; absorptive enterocytes are <i>ACE2+TMPRSS2+</i> -&gt; Interferon &amp; influenza increase <i>ACE2</i> in human nasal epithelia and lung tissue -&gt; Mouse <i>Ace2</i> is not upregulated by interferon, raising implications for disease models</p>
Infection, Genetics and Evolution 24APR2020	<b>Emerging genetic diversity among clinical isolates of SARS-CoV-2: Lessons for today</b>	Sheikh, Javaid Ahmad; et al. India-Germany-UK <a href="https://doi.org/10.1016/j.meegid.2020.104330">https://doi.org/10.1016/j.meegid.2020.104330</a>	Phylogenetics	<p>Machine learning approaches to analyse genome sequences of 257 available SARS-CoV-2 clinical isolates :</p> <ul style="list-style-type: none"> <li>- <b>At least 5 different clades of SARS-CoV-2, great deal of genetic diversity emerging among clinical isolates.</b></li> <li>- <b>Every continent appears to have multiple introductions of different viral strains</b> (no geographical clustering unlike previous pandemics).</li> <li>- 5' terminal of viral genome more prone to mutations compared to 3' end.</li> <li>- <b>ORF1ab, spike, ORF3a and E proteins most prone to mutations.</b></li> <li>- <b>RBD of spike protein is a mutational hotspot (major driver of diversity).</b></li> <li>- <b>Greater transmission/mortality in the Lombardy region (Italy), compared to other European countries or African continent or China, could not be correlated to any specific molecular divergence pattern.</b></li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Annals of the rheumatic diseases 24APR2020	<b>Clinical course of coronavirus disease 2019 (COVID-19) in a series of 17 patients with systemic lupus erythematosus under long-term treatment with hydroxychloroquine</b>	Mathian, Alexis et al, France <a href="https://ard.bmj.com/content/early/2020/04/24/ard.2020-217566.long">https://ard.bmj.com/content/early/2020/04/24/ard.2020-217566.long</a>	Therapeutic	<p><b>Observational study</b> with the aim to follow the clinical course of COVID-19 in <b>patients with systemic lupus erythematosus (SLE) who received long-term treatment with HCQ</b> (17 patients).</p> <p>-&gt; Main comorbidities were obesity and chronic kidney disease. The duration of HCQ treatment prior to COVID-19 was relatively long, with a median (range) of 7.5 (0.5–29.8) years. Twelve (71%) patients were also treated with prednisone, and seven (41%) with an immunosuppressant.</p> <p>-&gt; This case series does not allow to draw conclusions on the incidence rate and severity of COVID-19 in SLE. However, it gives a first clinical picture of the course of this infection in patients with SLE treated with HCQ.</p> <p>Based on the observation that <b>most of the patients with SLE in this study received long-term treatment with HCQ, having blood concentrations of the drug within therapeutic range, is that HCQ does not seem to prevent COVID-19, at least its severe forms, in patients with SLE.</b></p>
Cell 24APR2020	<b>Clinically Applicable AI System for Accurate Diagnosis, Quantitative Measurements and Prognosis of COVID-19 Pneumonia Using Computed Tomography</b>	Zhang, Ket al., China <a href="https://www.cell.com/pb-assets/products/coronavirus/CELL_CELL-D-20-00656.pdf">https://www.cell.com/pb-assets/products/coronavirus/CELL_CELL-D-20-00656.pdf</a>	Diagnostics	<p>Many COVID-19 patients develop pneumonia called novel coronavirus pneumonia (NCP) and rapidly progress to respiratory failure. However, rapid diagnosis and identification of high-risk patients for early intervention are challenging.</p> <p>Development of an AI system that can diagnose NCP and differentiate it from other common pneumonia and normal controls.</p> <p>The AI system can assist radiologists and physicians in performing a quick diagnosis and is able to identify important clinical markers that correlated with the NCP lesion properties.</p> <p>It provides accurate clinical prognosis that can aid clinicians to consider appropriate early clinical management and allocate resources appropriately.</p> <p>This AI system has been made available globally to assist the clinicians to combat COVID-19.</p>
J Med Virol 24APR2020	<b>SARS-CoV-2 can be detected in urine, blood, anal swabs and oropharyngeal swabs specimens</b>	Peng, Liang and al. China <a href="https://onlinelibrary.wiley.com/doi/abs/10.1002/jmv.25936">https://onlinelibrary.wiley.com/doi/abs/10.1002/jmv.25936</a>	Diagnostics	<p>To assess the presence of the SARS-CoV-2 ribonucleic acid (RNA) in urine and blood specimens, and anal and oropharyngeal swabs.</p> <p>SARS-CoV-2 RNA was present in all 4 specimen types, though not all specimen types were positive simultaneously.</p> <p>CI*: SARS-CoV-2 can infect multiple systems, including the urinary tract. Testing different specimen types may be useful for monitoring disease changes and progression, and for establishing a prognosis</p>
JAMA network open, 24APR2020	<b>Effect of High vs Low Doses of Chloroquine Diphosphate as Adjunctive Therapy for Patients Hospitalized With Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) Infection: A Randomized Clinical Trial</b>	Borba, Mayla Gabriela Silva et al, Brazil, <a href="https://doi.org/10.1001/jamanetworkopen.2020.8857">https://doi.org/10.1001/jamanetworkopen.2020.8857</a>	Therapeutic	<p>Parallel, double-masked, randomized, phase IIb clinical trial <b>81 adult patients with severe acute respiratory syndrome SARS-CoV-2 infection</b></p> <p>High-dosage CQ (ie, <b>600 mg CQ twice daily for 10 days</b>) versus low-dosage CQ (ie, <b>450 mg twice daily on day 1 and once daily for 4 days</b>)</p> <p>At Day 13, 6 of 40 patients (15.0%) in the low-dose group had died, compared with 16 of 41 patients (39.0%) in the high-dose group. Prolongation of QTc interval was observed in 4 of 36 patients (11.1%) in the low-dose group and 7 of 37 patients (18.9%) in the high-dose group. In addition, 2 patients in the high-dose group (2.7%) experienced ventricular tachycardia. The trial was stopped.</p> <p><b>The preliminary findings of this study suggest that the higher CQ dosage should not be recommended for critically ill patients with COVID-19</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Nature Medicine 24APR2020	<b>The QT interval in patients with COVID-19 treated with hydroxychloroquine and azithromycin</b>	Chorin, Ehud et al, USA <a href="https://doi.org/10.1038/s41591-020-0888-2">https://doi.org/10.1038/s41591-020-0888-2</a>	Therapeutic	<p>Charts review and corrected QT (QTc) interval follow-up in a consecutive <b>cohort of 84 patients receiving HY</b> (400mg daily on D1, then 200mg daily from D2 to D5)/<b>AZ</b> (500mg per day for 5 days).</p> <ul style="list-style-type: none"> <li><b>QTc significantly prolonged.</b></li> <li>In a subset of nine (11%) of those patients, the QTc was severely prolonged to &gt;500 ms, a known marker of high risk of malignant arrhythmia and sudden cardiac death. Five of the nine patients had a normal QTc at baseline.</li> <li><b>No torsades de pointes</b> events recorded for any patients, including those with a severely prolonged QTc.</li> </ul> <p>=&gt; Suggest that the <b>QTc should be followed repeatedly in patients with COVID-19 who are treated with HY/AZ</b>, particularly in those with co-morbidities and in those who are treated with other QT-prolonging medications.</p>
JAMA Pediatrics 24APR2020	<b>Mental Health Status Among Children in Home Confinement During the Coronavirus Disease 2019 Outbreak in Hubei Province, China</b>	Xinyan Xie et al., China <a href="https://jamanetwork.com/journals/jamapediatrics/fullarticle/2765196">https://jamanetwork.com/journals/jamapediatrics/fullarticle/2765196</a>	Psy	<p>Investigation of depressive and anxiety symptoms among students in Hubei province, China.</p> <p>Restricted to home for a mean (SD) of 33.7 days -&gt; A total of 403 students (22.6%) and 337 students (18.9%) reported depressive and anxiety symptoms, respectively. -&gt; Students in Wuhan: significantly higher CDI-S scores than those in Huangshi + greater risk of depressive symptoms -&gt; Students who were slightly or not worried about being affected by COVID-19 had significantly lower CDI-S scores than those who were quite worried, with a decreased risk of depressive symptoms. Those who were not optimistic about the epidemic, compared with those who were quite optimistic, had significantly higher CDI-S scores, with an increased risk of depressive symptoms.</p> <p><b>There was no significant association between demographic characteristics and anxiety symptoms.</b></p>
Nat Com 24APR2020	<b>Neutralization of SARS-CoV-2 spike pseudotyped virus by recombinant ACE2-Ig</b>	Lei, Changhai et al, China <a href="https://doi.org/10.1038/s41467-020-16048-4">https://doi.org/10.1038/s41467-020-16048-4</a>	Therapeutic	<p>A recombinant protein was generated by connecting the extracellular domain of human ACE2 to the Fc region of the human immunoglobulin IgG1. A fusion protein containing an ACE2 mutant with low catalytic activity is also used in this study. The fusion proteins are then characterized.</p> <ul style="list-style-type: none"> <li>Both fusion proteins have a <b>high binding affinity for the receptor-binding domains of SARS-CoV and SARS-CoV-2</b> and exhibit <b>desirable pharmacological properties in mice.</b></li> <li>Moreover, the <b>fusion proteins neutralize virus pseudotyped with SARS-CoV or SARS-CoV-2 spike proteins in vitro.</b></li> </ul> <p>=&gt; As these fusion proteins exhibit cross-reactivity against coronaviruses, they have potential applications in the diagnosis, prophylaxis, and treatment of SARS-CoV-2.</p>
Clinica Chimica Acta 23APR2020	<b>Highly sensitive detection of SARS-CoV-2 RNA by multiplex rRT-PCR for molecular diagnosis of COVID-19 by clinical laboratories</b>	Shige, Takayuki and al. Japan <a href="https://www.sciencedirect.com/science/article/pii/S0009898120301789">https://www.sciencedirect.com/science/article/pii/S0009898120301789</a>	Diagnostics	<p>Three genes are used for multiplex rRT-PCR: the Sarbecovirus specific E gene, the SARS-CoV-2 specific N gene, and the human ABL1 gene as an internal control.</p> <p>CI*: The proposed multiplex rRT-PCR methodology enable highly sensitive detection of SARS-CoV-2 RNA, reducing reagent use and cost, and time required by clinical laboratory technicians.</p>
Journal of Thrombosis and Haemostasis 23APR2020	<b>Direct oral anticoagulant plasma levels striking increase in severe COVID-19 respiratory syndrome patients treated with antiviral agents. The Cremona experience</b>	Testa, Sophie et al, Italy <a href="https://doi.org/10.1111/jth.14871">https://doi.org/10.1111/jth.14871</a>	Therapeutic	<p>32 patients with <b>COVID 19 and direct oral anticoagulants (DOACs) eligible for antiviral therapy</b> (lopinavir, ritonavir or darunavir)</p> <p>DOAC stopped in 20, and continued in 12. On average, <b>C-trough DOAC levels were 6.14 times higher</b> during hospitalization than in pre-hospitalization period</p> <p><b>Physicians should consider withholding DOACs from patients with SARS-CoV-2 and replacing them with alternative parenteral antithrombotic strategies</b> for as long as antiviral agents are deemed necessary and until discharge</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
BMJ 23APR2020	<b>Covid-19: Two thirds of healthcare workers who have died were from ethnic minorities</b>	Rimmer et al., UK <a href="https://doi.org/10.1136/bmj.m1621">https://doi.org/10.1136/bmj.m1621</a>	HSS/Politic	<p>'- Two thirds of healthcare workers who have died from covid-19 were from an ethnic minority background, and at least half were not born in the UK : causes could be biological, medical, or sociological =&gt; need for a concerted effort to seek explanations and solutions.</p> <p>Notable absence of deaths occurred among certain staff groups:</p> <p>'- Deaths notably in surgery (five cases), general practice (four), emergency medicine (two. No anaesthetists or intensivists. =&gt;Better use of PPE?</p> <p>Need for a central registry of deaths among health and social care workers</p>
Canadian Journal of Political Science 23APR2020	<b>Sociodemographic and psychological correlates of compliance with the Covid-19 public health measures in France</b>	Bouraurd et al., France <a href="https://doi.org/10.1017/S0008423920000335">https://doi.org/10.1017/S0008423920000335</a>	HSS/Politic	<p>In order to face the Covid-19 pandemic, the French government adopted health measures at an unprecedented scale to slow the spread of the virus.</p> <p>However, compliance by citizens cannot be taken for granted.</p> <p>Individual panel data =&gt; personality and demographic characteristics are relevant predictors of compliance, more than ideology, trust (towards the government and scientists), and emotional reactions.</p> <p>Main characteristics that lead to complying with the measures are :</p> <ul style="list-style-type: none"> <li>- Age (older people)</li> <li>- Sex (women)</li> <li>- Conscientiousness</li> </ul> <p>Characteristics with no impact on behavior :</p> <ul style="list-style-type: none"> <li>- Education</li> <li>- Extraversion</li> <li>- Neuroticism</li> <li>- Ideological extremity</li> </ul>
Nature Medicine 23APR2020	<b>SARS-CoV-2 entry factors are highly expressed in nasal epithelial cells together with innate immune genes</b>	Sungnak, Waradon; et al. UK-France-Netherlands-Germany-USA <a href="https://doi.org/10.1038/s41591-020-0868-6">https://doi.org/10.1038/s41591-020-0868-6</a>	Fundamental research	<p>Tropism analysis from single cell RNA-seq datasets from multiple tissues from healthy human donors (Human Cell Atlas tissue consortium):</p> <ul style="list-style-type: none"> <li>- <b>ACE2 expressed in cells from multiple tissues at generally low levels.</b></li> <li>- <b>TMPRSS2 highly expressed with a broader distribution, suggesting that ACE2, rather than TMPRSS2, may be a limiting factor for intal viral entry stage.</b></li> <li>- <b>ACE2 and TMPRSS2 highest co-expression in nasal secretory epithelial cells (nasal goblet and ciliated cells), co-expressed with genes involved in innate immunity.</b></li> </ul> <p>Expression of viral receptor genes used by other coronaviruses and influenza viruses (ANPEP used by HCoV-22944, DPP4 used by MERS-CoV45, ST6GAL1, ST3GAL4 important for influenza viruses) show :</p> <ul style="list-style-type: none"> <li>- <b>expression distribution coincided with viral transmissibility based on a comparison to R0 -&gt; upper airway in viruses with higher R0/infectivity (SARS-CoV-2, influenza) vs lower airway/lung parenchyma for MERS-CoV</b></li> </ul> <p>-&gt; All data provided as a user-friendly an open resource: <a href="http://www.covid19cellatlas.org">www.covid19cellatlas.org</a></p>
Radiology 23APR2020	<b>Acute Pulmonary Embolism in COVID-19 Patients on CT Angiography and Relationship to D-Dimer Levels</b>	Leonard-Lorant I and al, France <a href="https://doi.org/10.1148/radiol.2020201561">https://doi.org/10.1148/radiol.2020201561</a>	Clinic/Radiology	<p>106 patients COVID19+ and CT angiograms</p> <p><b>32 (30%) positive for pulmonary embolus:</b></p> <ul style="list-style-type: none"> <li>- Higher D-dimer levels (6110 vs 1920, p&lt;0,01)</li> <li>- More in ICU (75% vs 32ù, p&lt;0,01)</li> <li>- Treated more often with LWMH (78% vs 23%)</li> </ul> <p>D-dimer levels &gt;2660µg/L :</p> <ul style="list-style-type: none"> <li>- Sensitivity 100%</li> <li>- Specificity 67%</li> </ul>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Emerging Infectious Disease 23APR2020	<b>Population-Based Estimates of Chronic Conditions Affecting Risk for Complications from Coronavirus Disease, United States</b>	Mary L. Adams; USA <a href="https://doi.org/10.3201/eid2608.200679">https://doi.org/10.3201/eid2608.200679</a>	Demographic / Risk assessment	<p>USA demographic analysis to infer risk for complications from COVID-19 due to chronic conditions: (Based on publicly available 2017 Behavioral Risk Factor Surveillance System (BRFSS) data from telephone surveys of 444,649 randomly selected adults (&gt;18 years of age) in the 50 states and District of Columbia).</p> <p><b>-&gt; 45.4% of US adults are potentially at increased risk of complications because of cardiovascular disease, diabetes, respiratory disease, hypertension, or cancer.</b></p> <p>- Rates increased by age: 19.8% for 18–29 years of age, 80.7% for persons &gt;80 years of age, and varied by state, race/ethnicity, health insurance status, and employment.</p>
Analytical chemistry 23APR2020	<b>Rapid and sensitive detection of anti-SARS-CoV-2 IgG using lanthanide-doped nanoparticles-based lateral flow immunoassay</b>	Chen, Zhenhua and al. China <a href="https://pubs.acs.org/doi/abs/10.1021/acs.analchem.0c00784">https://pubs.acs.org/doi/abs/10.1021/acs.analchem.0c00784</a>	Diagnostics	<p>Simple and rapid immunodiagnostic method based on lateral flow immunoassay (LFIA) that uses lanthanide-doped polystyrene nanoparticles (LNPs) to detect anti-SARV-CoV-2 IgG in human serum.</p> <p>The results of the validation experiment met the requirements for clinical diagnostic reagents</p> <p>Conclusion: this new assay can achieve rapid and sensitive detection of anti-SARS-CoV-2 IgG in human serum and allow positive identification in suspicious cases; it can also be useful for monitoring the progression COVID-19 and evaluating patients' response to treatment.</p>
Science of the Total Environment journal 22APR2020	<b>COVID-19 outbreak: Migration, effects on society, global environment and prevention</b>	Chakraborty et al., India <a href="https://doi.org/10.1016/j.scitotenv.2020.138882">https://doi.org/10.1016/j.scitotenv.2020.138882</a>	HSS/Politic	<p>Economic impact:</p> <ul style="list-style-type: none"> <li>- Threat of high inflation and high unemployment as a result of lack of productivity and increased expenditures</li> <li>- For each month there will be an approximate loss of 2% points in annual GDP growth</li> <li>- The tourism sector alone faces an output decrease as high as 50% to 70%</li> </ul> <p>Global environment :</p> <ul style="list-style-type: none"> <li>-&gt; Non-functioning of industries: decrease of industrial waste emission, recovery of ecosystems and revival of ozone layer.</li> <li>-&gt;Deforestation linked to disease outbreaks.</li> <li>-&gt;Population growth: increasing sources of pollution + deforestation = exposing populations to new pathogens</li> </ul> <p>The global strategy for COVID-19 prevention and control:</p> <ul style="list-style-type: none"> <li>• Global threat that requires a global response involving all countries – in the short term: Restricting mass gatherings + research for new drugs/vaccines/prevention</li> </ul> <ul style="list-style-type: none"> <li>• In the long run: <ul style="list-style-type: none"> <li>o Forestation/Respecting wildlife habitats.</li> <li>o Control of population growth</li> <li>o Global ban on wildlife trade</li> </ul> </li> </ul>
Science 22APR2020	<b>Structure-based design of antiviral drug candidates targeting the SARS-CoV-2 main protease</b>	Dai, Wenhao et al, China <a href="https://doi.org/10.1126/science.abb4489">https://doi.org/10.1126/science.abb4489</a>	Therapeutic	<p>The main protease (Mpro) of SARS-CoV-2 is a key enzyme that plays a pivotal role in mediating viral replication and transcription.</p> <p><b>Two lead compounds (11a and 11b) targeting Mpro were designed and synthesized.</b></p> <p>Both exhibited excellent inhibitory activity and <b>potent anti-SARS-CoV-2 infection activity</b>. The X-ray crystal structures of SARS-CoV-2 Mpro in complex with 11a or 11b, both determined at 1.5 Å resolution, showed that the <b>aldehyde groups of 11a and 11b are covalently bound to Cys145 of Mpro</b>.</p> <p>Both compounds showed <b>good PK properties in vivo</b>, and 11a also exhibited low toxicity, suggesting that these compounds are promising drug candidates.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
European Journal of Epidemiology 22APR2020	<b>Vaccine confidence in the time of COVID-19</b>	Harrison et al., US <a href="https://doi.org/10.1007/s10654-020-00634-3">https://doi.org/10.1007/s10654-020-00634-3</a>	HSS/Politic	<p><b>Rushing for a vaccine in the Covid19 epidemic will not solve the problem of vaccine hesitation among the population.</b></p> <p>Need to re-imagine the culture of public health more broadly than the delivery of vaccine /technology</p> <p><b>4 points to consider:</b></p> <p>1- Vaccination technologies presented as a technical/ objective solution to the problem of prevention and health of the population. The broader view of public health (social/political equality...) has been eroded: idea that vaccines could obviate need for broader social and environmental policies kept prevention in clinical hands.</p> <p>2- The success, in the first years, of preventive mass vaccination programs ≠ a timeless public acceptance of vaccination as a preventive strategy. Result of a complex set of circumstances.</p> <p>3- Reluctance to vaccinate: symptom of a greater desire to ignore threats because they are not bothersome or do not constitute an emergency (yet).</p> <p>4- Essential ethical dilemma of public health: tension between autonomy and state power</p> <p>Oppose a logic of "care": capacity of the States to provide the biomedical resources but also the economic and social resources making it possible to fairly protect the life of the citizens against health disasters.</p> <p>=&gt; A broader confidence in vaccines or any pre-emptive measure depends on widespread public trust in preventive health directives and involves strong care and social solidarity structures.</p>
The Lancet Infectious diseases 22APR2020	<b>No SARS-CoV-2 detected in amniotic fluid in mid-pregnancy</b>	Yu, Nan; et al. China <a href="https://doi.org/10.1016/S1473-3099(20)30320-0">https://doi.org/10.1016/S1473-3099(20)30320-0</a>	Clinic	<p>2 pregnant women diagnosed with COVID-19 at first trimester :</p> <ul style="list-style-type: none"> <li>- In the second trimester, both positive for SARS-CoV-2 total antibodies in serum and negative for SARS-CoV-2 RNA in throat swabs.</li> <li>- Amniotic fluid : negative (RT-PCR) and SARS-CoV-2 IgM and IgG negative in both patients.</li> <li>- In serum : both IgG positive, and only case 1 tested positive for IgM.</li> </ul> <p><b>-&gt; No SARS-CoV-2 detected in the amniotic fluid of both women diagnosed with COVID-19 in early stage of pregnancy.</b></p> <p>- <b>Limit</b> : Only 2 patients, sensitivity (RNA is much less stable in amniotic fluid than is DNA), lack of cord blood.</p>
Eur J Neurol 22APR2020	<b>Acute-onset smell and taste disorders in the context of Covid-19: a pilot multicenter PCR-based case-control study</b>	Beltran-Corbellini A and al, Spain <a href="https://doi.org/10.1111/ene.14273">https://doi.org/10.1111/ene.14273</a>	Clinic	<p>Multicenter study – cases (79) controls (40) study</p> <p><i>Controls: historical group of season influenza patients</i></p> <p>Basal characteristics: no difference between group</p> <p><i>Cases: 31(39%) with new onset smell or taste disorders (STD):</i></p> <p>for 35.5% initial symptoms</p> <p>Now-set of STD more frequent in cases than controls:</p> <ul style="list-style-type: none"> <li>- Adjusted OR: 21,4 [2.77 – 165.4]</li> </ul> <p>No difference for gender/smoking habits/severity between STD or not in case group.</p> <p>Increased frequency of STD in young patients</p> <p><b>STD more frequent among COVID-19 patients</b></p> <p><u>Limitations:</u> historical controls – lack of comparison with others virus – self reported questionnaire</p>
Radiology 22APR2020	<b>Acute Pulmonary Embolism Associated with COVID-19 Pneumonia Detected by Pulmonary CT Angiography</b>	Grillet F and al, France <a href="https://doi.org/10.1148/radiol.2020201544">https://doi.org/10.1148/radiol.2020201544</a>	Clinic/Radiology	<p>100 patients COVID19+ and severely ill</p> <p>Mean age: 66 years and 70% males</p> <p>23% had acute pulmonary embolus:</p> <ul style="list-style-type: none"> <li>- More frequent in critical care unit (p&lt;0,01)</li> <li>- Longer delay from symptoms onset to CT (12 d)</li> </ul> <p>Requirement of mechanical ventilation was associated with pulmonary embolus (OR=3,8, p=0,049)</p> <p>Extent of lesions was not associated with pulmonary embolus.</p> <p><b>Contrast enhanced CT rather for these patients</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Science of the Total Environment journal 22APR2020	<b>COVID-19 outbreak: Migration, effects on society, global environment and prevention</b>	Chakraborty, I. al., India <a href="https://doi.org/10.1016/j.scitotenv.2020.138882">https://doi.org/10.1016/j.scitotenv.2020.138882</a>	HSS/Politic	<p><b>Economic impact:</b></p> <ul style="list-style-type: none"> <li>- Threat of high inflation and high unemployment as a result of lack of productivity and increased expenditures</li> <li>- For each month there will be an approximate loss of 2% points in annual GDP growth</li> <li>- The tourism sector alone faces an output decrease as high as 50% to 70%</li> </ul> <p><b>Global environment :</b></p> <ul style="list-style-type: none"> <li>-Non-functioning of industries: decrease of industrial waste emission, recovery of ecosystems and revival of ozone layer.</li> <li>-Deforestation linked to disease outbreaks.</li> <li>-Population growth: increasing sources of pollution + deforestation = exposing populations to new pathogens</li> </ul> <p><b>The global strategy for COVID-19 prevention and control:</b></p> <ul style="list-style-type: none"> <li>• Global threat that requires a global response involving all countries – in the short term: Restricting mass gatherings + research for new drugs/vaccines/prevention</li> </ul> <ul style="list-style-type: none"> <li>• In the long run: <ul style="list-style-type: none"> <li>o Forestation/Respecting wildlife habitats.</li> <li>o Control of population growth</li> <li>o Global ban on wildlife trade</li> </ul> </li> </ul>
Gastroenterology, 21APR2020	<b>Characteristics and prognosis of patients with inflammatory bowel disease during the SARS-CoV-2 pandemic in the Basque Country (Spain)</b>	Rodríguez-Lago, Iago et al, Espagne, <a href="https://doi.org/10.1053/j.gastro.2020.04.043">https://doi.org/10.1053/j.gastro.2020.04.043</a>	Clinic Gastroenterology	<p><b>Patients (N=40) with inflammatory bowel disease (IBD) and a positive test for SARS-CoV-2</b> from 5 sites as for the 8th April 2020. Mean age: 59 (range 18 – 90)</p> <p><b>28% under immunomodulator (28%) and 18% under biologic monotherapy.</b></p> <p>Most frequent symptoms: fever (77%) and cough (67%), with 21% reporting diarrhea</p> <p>No patient was admitted to the ICU</p> <p><b>Two deaths</b> were reported (5%):</p> <ul style="list-style-type: none"> <li>a 86-year-old male with diabetes, prostate adenocarcinoma and ulcerative proctitis on mesalamine</li> <li>a 77-year-old male with dementia and left-sided ulcerative colitis under mesalamine and methotrexate.</li> </ul> <p><b>patients with IBD and COVID have a good overall prognosis</b></p>
Ophthalmology 21APR2020	<b>Ocular Findings and Proportion with Conjunctival SARS-COV-2 in COVID-19 Patients</b>	Zhou, Yunyun and al. China <a href="https://www.sciencedirect.com/science/article/pii/S016164202030405X">https://www.sciencedirect.com/science/article/pii/S016164202030405X</a>	Virology	<p>Study of 121 patients</p> <p>SARS-CoV-2 RNA could be detected in the conjunctival swabs of 2.5% (3/121) patients.</p> <p>-&gt;Eight patients (6.6%) had ocular symptoms: itching, redness, tearing, discharge, and foreign body sensation.</p> <p>-&gt;Two patients without ocular symptoms tested positive for conjunctival SARS-CoV-2.</p> <p>The appearance of ocular symptoms or the result of conjunctival SARS-CoV-2 detection was not significantly correlated with the duration of disease. The proportion with a positive result for SARS-CoV-2 RNA was significantly different between the conjunctival and nasopharyngeal specimens. These findings may help to improve precaution practices during the COVID-19 pandemic.</p>
Gastroenterology 22APR2020	<b>Prevalence and Characteristics of Gastrointestinal Symptoms in Patients with SARS-CoV-2 Infection in the United States: A Multicenter Cohort Study</b>	Walker D.R and al, USA <a href="https://doi.org/10.1053/j.gastro.2020.04.045">https://doi.org/10.1053/j.gastro.2020.04.045</a>	Clinic	<p>Multicenter study (9 centers) - <b>318 patients COVID+ 61,3%= reported at least 1 gastrointestinal symptom:</b> anorexia (34,8%), diarrhea (33,7%) and nausea (26,4%)</p> <p>Lost of smell/taste more frequent in gastrointestinal symptoms group (p&lt;0,05)</p> <p><b>No difference</b> in patients with gastrointestinal symptom and those without for:</p> <ul style="list-style-type: none"> <li>- Laboratory results</li> <li>- Rates of clinical deterioration</li> <li>- ICU admission, mechanical ventilation, mortality</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet Psychiatry 21APR2020	<b>Suicide risk and prevention during the COVID-19 pandemic</b>	Gunnel et al., UK <a href="https://doi.org/10.1016/S2215-0366(20)30171-1">https://doi.org/10.1016/S2215-0366(20)30171-1</a>	Psy	<p>-&gt;Suggestions that suicide rates will rise</p> <p>-&gt;Many people vulnerable to mental health problems and suicidal behaviour</p> <p>-&gt;Need for timely public health responses: list of actions presented in the publication</p> <p><b>Mental health consequences are likely to be present for longer and peak later than the actual pandemic.</b></p>
Emerging Infectious Disease journal 21APR2020	<b>Possible Bat Origin of Severe Acute Respiratory Syndrome Coronavirus 2</b>	Susanna K.P. Lau; et al. Hong Kong, China <a href="https://doi.org/10.3201/eid2607.200092">https://doi.org/10.3201/eid2607.200092</a>	Virology	<p>Phylogenetic analysis :</p> <ul style="list-style-type: none"> <li>- SARS-CoV-2 <b>genome closest to that of SARS-related coronaviruses (SARSr-CoVs) from horseshoe bats, and receptor-binding domain (RBD) closest to that of pangolin viruses.</b></li> <li>- <b>Potential recombination sites identified around the RBD region</b></li> <li>- none of existing SARSr-CoVs represents its immediate ancestor.</li> </ul> <p>--&gt; <b>SARS-CoV-2 probably a novel recombinant virus (genome backbone evolved from Yunnan bat virus—like SARSr-CoVs and RBD from pangolin virus—like SARSr-CoVs).</b></p> <p>Its origin and direct ancestral viruses not identified.</p>
BMJ 21APR2020	<b>Viral load dynamics and disease severity in patients infected with SARS-CoV-2 in Zhejiang province, China, January-March 2020: retrospective cohort study</b>	Zheng S and al, China <a href="https://doi.org/10.1136/bmj.m1443">https://doi.org/10.1136/bmj.m1443</a>	Clinic/Virology	<p><b>Retrospective study – hospitalized patients only</b></p> <p><b>3497 samples collected from 96 patients COVID-19</b></p> <p>Samples: serum/respiratory/stool/urine</p> <p>Duration of virus significantly longer in stool samples</p> <p><u>Respiratory samples:</u></p> <ul style="list-style-type: none"> <li>- median duration of virus in severe disease was significantly longer than in mild disease (14 days, 10-21 days; P=0.04)</li> <li>- patients with severe disease: significantly higher viral loads.</li> <li>- Letter shedding peak in severe group</li> </ul> <p>Other samples: no difference</p> <p>No effect of the antiviral treatment on viral load/duration</p> <p><u>Factors associated significantly with duration of virus:</u></p> <ul style="list-style-type: none"> <li>- glucocorticoid &gt; 10 days in severe group</li> <li>- men</li> <li>- &gt; 60 years</li> </ul> <p><u>Limitations:</u> small sample size / viral load influenced by many factors</p>
Thrombosis and haemostasis, 21APR2020	<b>COVID-19-Related Severe Hypercoagulability in Patients Admitted to Intensive Care Unit for Acute Respiratory Failure</b>	Spiezia, Luca et al, Italy <a href="https://www.ncbi.nlm.nih.gov/pubmed/32316063">https://www.ncbi.nlm.nih.gov/pubmed/32316063</a>	Clinic Hematology	<p>Evaluation of coagulation abnormalities via traditional tests and <b>whole blood thromboelastometry profiles</b> in a group of <b>22 patients</b> with COVID and acute respiratory failure due to COVID-19 (mean age 67 ± 8 years, M:F 20:2) compared to 44 controls.</p> <p>Cases showed significantly <b>higher fibrinogen and D-dimer plasma levels</b> versus healthy controls</p> <p><b>Markedly hypercoagulable thromboelastometry profiles</b> in COVID-19 patients,</p> <p>COVID-19 patients with acute respiratory failure present a <b>severe hypercoagulability rather than consumptive coagulopathy</b></p>
CDC Morbidity and Mortality Weekly Report 20APR2020	<b>Cleaning and Disinfectant Chemical Exposures and Temporal Associations with COVID-19 — National Poison Data System, United States, January 1, 2020–March 31, 2020</b>	Chang et al., USA <a href="https://www.cdc.gov/mmwr/volumes/69/wr/mm6916e1.htm?_cid=mm6916e1_wt&amp;con=tribAff">https://www.cdc.gov/mmwr/volumes/69/wr/mm6916e1.htm?_cid=mm6916e1_wt&amp;con=tribAff</a>	Public Health/Epidemiology	<p>-&gt;To assess whether there might be a possible association between COVID-19 cleaning recommendations from public health agencies and the media and the number of chemical exposures reported to the National Poison Data System (NPDS)</p> <p>-&gt; During January–March 2020, poison centers received 45,550 exposure calls related to cleaners (28,158) and disinfectants (17,392), representing overall increases of 20.4% and 16.4% from January–March 2019 (37,822) and January–March 2018 (39,122), respectively.</p> <p><b>Although NPDS data do not provide information showing a definite link between exposures and COVID-19 cleaning efforts, there appears to be a clear temporal association with increased use of these products.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Antimicrobial agents and chemotherapy 20APR2020	<b>Nafamostat mesylate blocks activation of SARS-CoV-2: New treatment option for COVID-19</b>	Hoffmann, Markus et al, Germany <a href="https://doi.org/10.1128/AAC.00754-20">https://doi.org/10.1128/AAC.00754-20</a>	Therapeutic	<p>The SARS-24 CoV-2 spike protein (S) is inserted into the viral envelope and mediates viral entry into cells. For this, the S protein depends on the cellular enzyme transmembrane protease serine 2 (TMPRSS2), which cleaves and thereby activates the S protein. <b>Serine protease inhibitors gabexate mesylate (FOY), nafamostat mesylate (Futhan) along with camostat mesylate were tested for inhibition of SARS-CoV-2 infection of lung cells.</b> All compounds are approved for human use in Japan.</p> <p>Gabexate mesylate slightly inhibited SARS-CoV-2 S-driven host cell entry while <b>camostat mesylate robustly suppressed entry</b>. Notably, <b>nafamostat mesylate</b>, which is FDA-approved for indications unrelated to coronavirus infection, <b>inhibited SARS-CoV-2 S-mediated entry into host cells with roughly 15-fold higher efficiency than camostat mesylate</b>, with an EC50 in the low nanomolar range. Moreover, <b>nafamostat mesylate blocked SARS-CoV-2 infection of human lung cells with markedly higher efficiency than camostat mesylate</b> while both compounds were not active against vesicular stomatitis virus infection, as expected.</p>
J of Emerg Microb and Inf 20APR2020	<b>Different longitudinal patterns of nucleic acid and serology testing results based on disease severity of COVID-19 patients</b>	Yongchen et al., China <a href="https://www.tandfonline.com/doi/full/10.1080/22221751.2020.1756699">https://www.tandfonline.com/doi/full/10.1080/22221751.2020.1756699</a>	Diagnostic	<p>-&gt; Detailed timeline of nucleic acid testing results for throat or anal samples along with the anti-SARS-CoV-2 IgM and IgG responses in 21 individuals infected with SARS-CoV-2, including 11 non-severe COVID-19 patients, 5 severe COVID-19 patients and 5 asymptomatic carriers.</p> <p>-&gt; Seroconversion was observed in 100% (17/17) of symptomatic patients during the observation period</p> <p>-&gt; Did not identify a strong association of seroconversion and disease severity in our cohort</p>
Metabolism: clinical and experimental 19APR2020	<b>Obesity as a risk factor for greater severity of COVID-19 in patients with metabolic associated fatty liver disease</b>	Zheng, Kenneth I and al China <a href="http://www.sciencedirect.com/science/article/pii/S0026049520301086">http://www.sciencedirect.com/science/article/pii/S0026049520301086</a>	Clinic	<p><b>Sixty six COVID-19 patients with metabolic associated fatty liver disease (MAFLD)</b></p> <p>n=45 with Body mass index (BMI) &gt; 25 and n=21 with BMI &lt; 25 Mean age :47 years and 74,2% female</p> <p>Results : <b>BMI &gt; 25 was related to more severe COVID 19</b> (adjusted-OR 6.32 95%CI 1.16 -34.54, p =0.033) even after adjusting for age, sex, smoking, diabetes, hypertension, and dyslipidaemia</p>
Clin Inf Dis 19APR2020	<b>Profile of RT-PCR for SARS-CoV-2: a preliminary study from 56 COVID-19 patients</b>	Xiao et al., China <a href="https://academic.oup.com/cid/article/doi/10.1093/cid/cia460/5822175">https://academic.oup.com/cid/article/doi/10.1093/cid/cia460/5822175</a>	Diagnostic	<p>Dynamics profile of SARS-CoV-2 from 56 recovered COVID-19 patients</p> <p>-&gt; Virus shedding was up to 6 weeks after onset of symptoms</p> <p>-&gt; Longest duration between RT-PCR test for SARS-CoV-2: 42 days after onset of symptoms.</p> <p>-&gt; Median duration between onset of symptom to nucleic acid conversion: 24 days</p> <p>-&gt; In first 3 weeks after symptoms onset, majority results of RT-PCR for SARS-CoV-2 were positive. From week 3 after symptoms onset, number of negative RT-PCR results increased.</p> <p>-&gt; The positive rate of RT-PCR test results was highest at week 1 (100%), followed by 89.3%, 66.1%, 32.1%, 5.4% and 0% at week 2, week 3, week 4, week 5 and week 6 respectively.</p>
Clinical Therapeutic 19APR2020	<b>Association between clinical manifestations and prognosis in patients with COVID-19</b>	Yu T and al, China <a href="https://doi.org/10.1016/j.clinthera.2020.04.009">https://doi.org/10.1016/j.clinthera.2020.04.009</a>	Clinic	<p>Multicenter study – 95 patients COVID-19+</p> <p><b>73 had pneumonia (CT findings)</b>, significantly:</p> <ul style="list-style-type: none"> <li>- <b>Older</b></li> <li>- <b>Higher BMI</b>, ASAT and LDH levels</li> <li>- Lower lymphocyte and platelet count</li> </ul> <p>ARDS (n=24) and non-ARDS (n=71)</p> <p><u>Independent risk factors associated with ARDS:</u></p> <ul style="list-style-type: none"> <li>- High systolic blood pressure (OR:1.04, p=0.025)</li> <li>- High LDH level (OR:1.01, p=0.021)</li> </ul> <p><u>Association with pneumonia exacerbation (n=19):</u></p> <ul style="list-style-type: none"> <li>- High BMI (OR: 1.28, p=0.017)</li> <li>- Tobacco smoking (OR: 16.13, p=0.032)</li> </ul> <p><u>Limitations:</u> exacerbation based on CT scan findings</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Clin Inf Dis 19APR2020	<b>Antibody Detection and Dynamic Characteristics in Patients with COVID-19</b>	Xiang et al., China <a href="https://academic.oup.com/cid/article/doi/10.1093/cid/ciaa461/5822173">https://academic.oup.com/cid/article/doi/10.1093/cid/ciaa461/5822173</a>	Diagnostic	<p><b>ELISA based on the recombinant nucleocapsid protein of SARS-CoV-2</b> Seroconversion of specific IgM and IgG antibodies were observed as early as the 4<sup>th</sup> day after symptom onset.</p> <p>In confirmed patient: <b>IgM:</b> Sensitivity, 77.3% Specificity, 100% PPV, 100% NPV, 80.0% Consistency rate : 88.1% <b>IgG:</b> Sensitivity, 83.3.3% Specificity, 95.0% PPV, 94.8% NPV, 83.8% Consistency rate : 88.9 %</p> <p>In patients with suspected COVID-19, sensitivity, specificity, PPV, NPV, and consistency rate of IgM were 87.5% (21/24), 100%, 100%, 95.2%, and 96.4%, and those of IgG were 70.8% (17/24), 96.6%, 85.0%, 89.1%, and 88.1%.</p> <p><b>-&gt; Both antibodies performed well in serodiagnosis for COVID-19 rely on great specificity.</b></p>
Med 18APR2020	<b>Efficacy and safety of lopinavir/ritonavir or arbidol in adult patients with mild/moderate COVID-19: an exploratory randomized controlled trial</b>	Li, Y et al, China <a href="https://marlin-prod.literatumonline.com/pb-assets/products/coronavirus/MEDJ1.pdf">https://marlin-prod.literatumonline.com/pb-assets/products/coronavirus/MEDJ1.pdf</a>	Therapeutics	<p><b>Exploratory randomized (2:2:1) controlled trial</b> assessing the efficacy and safety of <b>lopinavir/ritonavir (LPV/r) or arbidol monotherapy</b> for treating patients with mild/moderate COVID-19. 86 patients with mild/moderate COVID-19 enrolled.</p> <p>LPV/r and arbidol <b>did not shorten the time of positive-to-negative conversion of COVID-19 nucleic acid in respiratory specimens</b> (9.0 vs. 9.1 vs. 9.3 days), <b>nor did they improve the symptoms of COVID-19 or pneumonia on lung CT imaging at 7 days and 14 days.</b> More patients treated with LPV/r progressed from mild/moderate to severe/critical status than patients from the other two groups. Adverse events occurred in the treatment groups.</p> <p>Limitations: small sample size, no severely or critically ill patients, or patients at increased risk of poor outcome with many comorbidities, not completely blinded.</p>
European urology 18APR2020	<b>Coronavirus Disease 2019 Pneumonia in Immunosuppressed Renal Transplant Recipients: A Summary of 10 Confirmed Cases in Wuhan, China</b>	Zhu L and al, China <a href="http://www.sciencedirect.com/science/article/pii/S0302283820302141">http://www.sciencedirect.com/science/article/pii/S0302283820302141</a>	Clinic	<p>Controls: 10 family members <b>10 kidney transplant patients + COVID-19 pneumonia:</b></p> <ul style="list-style-type: none"> <li>- Classical symptoms: fever, cough, shortness of breath, ....</li> <li>- 100%: lymphopenia and elevated CRP</li> <li>- 50% had temporally increase of serum creatinine</li> <li>- Abnormalities on chest CT scan</li> <li>- <b>8/10 were severe or critical cases and 1 died</b></li> </ul> <p><u>Versus controls:</u></p> <ul style="list-style-type: none"> <li>- <b>Transplants patients more severe</b></li> <li>- <b>Much longer time to become negative for SARS-CoV-2</b> (median time: 28,4 d)</li> <li>- <b>Reduce fatal severe pneumonia:</b> suppressing the hyperimmune response</li> </ul>
Science 17APR2020	<b>Comparative pathogenesis of COVID-19, MERS, and SARS in a nonhuman primate model</b>	Rockx, Barry; et al. Netherlands <a href="https://doi.org/10.1126/science.abb7314">https://doi.org/10.1126/science.abb7314</a>	Fundamental research	<p>Cynomolgus macaques inoculated with SARS-CoV-2 or MERS-CoV.</p> <ul style="list-style-type: none"> <li>- SARS-CoV-2 causes COVID-19-like disease in macaques: virus excreted from nose and throat in the absence of clinical signs, detected in type I and II pneumocytes in foci of diffuse alveolar damage and in ciliated epithelial cells of nasal, bronchial, and bronchiolar mucosae.</li> <li>- lung lesions typically more severe with SARS-CoV-2 than in MERS-CoV infection, where virus was detected mainly in type II pneumocytes.</li> </ul> <p><b>-&gt; Cynomolgus macaques provide a new infection model to test preventive and therapeutic strategies.</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
J Thromb Haemost 17APR2020	<b>The procoagulant pattern of patients with COVID-19 acute respiratory distress syndrome</b>	Ranucci M and al, Italy <a href="https://doi.org/10.1111/jth.14854">https://doi.org/10.1111/jth.14854</a>	Clinic	<p><b>16 patients COVID-19 pneumonia and ARDS in ICU</b> 94% were male and 31% were obese D-Dimer, IL-6 and fibrinogen = <b>higher than upper limit</b> <b>Association between IL-6 and fibrinogen levels</b> Clot firmness higher than normal <u>Follow-up:</u> - Significant decrease of D-dimere and fibrinogen - Significant prolongation of the aPTT Pro-coagulant profile of COVID-19 ARDS patients and its normalization after an increased thromboprophylaxis. <u>Limitation:</u> lack of data on thrombin generation and fibrinolysis. Further studies: best prophylaxis and treatment ?</p>
Circulation research 17APR2020	<b>Association of Inpatient Use of Angiotensin Converting Enzyme Inhibitors and Angiotensin II Receptor Blockers with Mortality Among Patients With Hypertension Hospitalized With COVID-19</b>	Zhang, Peng et al, China <a href="https://doi.org/10.1161/CIRCRESAHA.120.317134">https://doi.org/10.1161/CIRCRESAHA.120.317134</a>	Therapeutics	<p><b>Retrospective, multi-center study including 1128 adult patients with hypertension diagnosed with COVID-19</b>, including <b>188 taking ACEI/ARB</b> (ACEI/ARB group; median age 64 [IQR 55-68] years; 53.2% men) <b>and 940 without using ACEI/ARB</b> (non-ACEI/ARB group; median age 64 [IQR 57-69]; 53.5% men).</p> <p>Among hospitalized COVID-19 patients with hypertension, inpatient <b>use of ACEI/ARB was associated with lower risk of all-cause mortality compared with ACEI/ARB non-users.</b></p> <p>Limitations: hospital only, modest sample size, retrospective study.</p>
Gut 17APR2020	<b>Covid-19 and immunomodulation in IBD</b>	Neurath, Germany <a href="https://gut.bmj.com/content/early/2020/04/16/gutjnl-2020-321269">https://gut.bmj.com/content/early/2020/04/16/gutjnl-2020-321269</a>	Immunology	<p><b>Results/ recommendations:</b></p> <ul style="list-style-type: none"> <li>No evidence for an increased risk or aggravated outcomes in patients with IBD in the context of covid-19</li> <li>However, covid-19 risks situation comprise older patients with IBD with comorbidities as well as patients suffering from malnutrition</li> <li>Experimental covid-19 treatment with hydroxychloroquine or remdesivir may increase the risks for drug-drug interactions with established IBD medications.</li> <li>Currently available recommendations for patients with IBD are : <ul style="list-style-type: none"> <li>Continue current treatment if disease is stable and discuss suitable medicine if disease has flared</li> <li>Use of mesalamine should be continued and should not increase the risk of infection.</li> <li>Corticosteroid use can be continued, but be cautious of possible side effects.</li> <li>A new prescription of immunosuppressant or increase in dose of an ongoing immunosuppressant is not recommended in epidemic areas.</li> <li>Use of biologics such as the antitumour necrosis factors infliximab or adalimumab should be continued.</li> <li>If infliximab infusion is not accessible, switching to adalimumab injection at home should be considered.</li> <li>Vedolizumab use can be continued due to the specificity of the drug for the intestine.</li> <li>Ustekinumab use can be continued, but starting ustekinumab requires infusion centre visits and therefore should be discussed before initiation of therapy.</li> <li>Enteral nutrition might be used if biologics are not accessible</li> <li>Tofacitinib should not be newly prescribed in epidemic areas unless there are no other alternatives.</li> </ul> </li> </ul>
Nature Biotechnology 16APR2020	<b>CRISPR-Cas12-based detection of SARS-CoV-2</b>	Broughton, James P. and al. USA <a href="https://doi.org/10.1038/s41587-020-0513-4">https://doi.org/10.1038/s41587-020-0513-4</a>	Diagnostic	<p>Development of a rapid (&lt;40 min), easy-to-implement and accurate CRISPR-Cas12-based lateral flow assay for detection of SARS-CoV-2 from respiratory swab RNA extracts.</p> <p>Validation using contrived reference samples and clinical samples, including 36 patients with COVID-19 infection and 42 patients with other viral respiratory infections.</p> <p>The CRISPR-based DETECTR assay provides a visual and faster alternative to SARS-CoV-2 real-time RT-PCR assay, with 95% positive predictive agreement and 100% negative predictive agreement.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
NEJM 16APR2020	<b>Not a Perfect Storm — Covid-19 and the Importance of Language</b>	Brandt M. et al., USA <a href="https://doi.org/10.1056/NEJMp2005032">https://doi.org/10.1056/NEJMp2005032</a>	HSS/Politic	<p>Metaphors we use to describe disease shape our experience of illness : has an impact on the approach taken against the Covid =&gt; misleading our approach. Covid characterized as a "perfect storm"="fierce storm arising from a rare combination of adverse meteorological factors"</p> <p>(randomness/volatility =&gt; reactive, disempowering). Vs developing and implementing preventive strategies to prepare for pandemics</p> <p>What is the impact of the language facing Pandemics? - The use of language eludes important conversation about our responsibility for emerging zoonoses + effects on the most vulnerable people. - The force of language minimizes our capacity to anticipate and prevent the next epidemics: long term investments in disease tracking and surveillance, scientific research and public health infrastructure</p> <p><b>Conclusion:</b> Covid is not a simple natural event. It is also the result of human actions. =&gt; Be conscientious about our language and its implications as a first step for reforms and preventive measures to strengthen our health infrastructure to face future disease outbreaks.</p>
Cell 16APR2020	<b>Development of CRISPR as an antiviral strategy to combat SARSCoV-2 and influenza</b>	Abbott, T et al, USA <a href="https://www.cell.com/pb-assets/products/coronavirus/CELL_CELL-D-20-00736.pdf">https://www.cell.com/pb-assets/products/coronavirus/CELL_CELL-D-20-00736.pdf</a>	Therapeutics	<p>A CRISPR-Cas13-based strategy, PAC-MAN (Prophylactic Antiviral CRISPR in huMAN cells), for viral inhibition can <b>effectively degrade RNA from SARS-CoV-2 sequences</b> and live influenza A virus (IAV) in human lung epithelial cells. CRISPR RNAs (crRNAs) targeting conserved viral regions were designed and screened, and <b>functional crRNAs targeting SARS-CoV-2 were identified</b>. The bioinformatic analysis showed a group of only six crRNAs can target more than 90% of all coronaviruses.</p> <p>With the development of a safe and effective system for respiratory tract delivery, PAC-MAN has the potential to become an important pan-coronavirus inhibition strategy.</p>
Plos One 16APR2020	<b>Mental health problems and social media exposure during COVID-19 outbreak</b>	Gao et al., China <a href="https://journals.plos.org/plosone/article?id=10.1371/journal.pone.0231924">https://journals.plos.org/plosone/article?id=10.1371/journal.pone.0231924</a>	Psy	<p><b>Social media exposure (SME) +++ during Covid-19.</b> Study on 4872 participants from 31 provinces and autonomous region -&gt; High prevalence of mental health problems -&gt; frequently SME was positively associated with high odds of anxiety (OR = 1.72, 95%CI: 1.31–2.26) and combination of depression and anxiety (CDA) (OR = 1.91, 95%CI: 1.52–2.41) compared with less SME.</p> <p>-&gt; Need to combat with “infodemic” while combating during public health emergency</p>
International journal of antimicrobial agents 16APR2020	<b>Can post-exposure prophylaxis for COVID-19 be considered as one of outbreak response strategies in long-term care hospitals?</b>	Lee, Sun Hee et al, Republic of Korea <a href="https://doi.org/10.1016/j.ijantimicag.2020.105988">https://doi.org/10.1016/j.ijantimicag.2020.105988</a>	Therapeutics	<p>After a large COVID-19 exposure event in a long-term care hospital (LTCH) in Korea, <b>PEP using hydroxychloroquine (HCQ)</b> was conducted to <b>211 persons</b> including 189 patients and 22 careworkers, with baseline negative PCR tests for COVID-19 (oral, dose of 400mg daily until the completion of 14 days of quarantine).</p> <p>=&gt; PEP was completed in 184 (97.4%) patients and 21 (95.5%) careworkers <b>without serious adverse events</b>. =&gt; At the end of 14 days of quarantine, <b>follow-up PCR tests were all negative</b>.</p> <p>Limitations: - no control group. - 92 hospital staff showed negative results by RT-PCR after 14-day quarantine even though they did not receive PEP, however this group was considered at low risk exposure.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Respiratory Research 15APR2020	<b>Prognostic value of NT-proBNP in patients with severe COVID-19</b>	Gao L and al, China <a href="https://doi.org/10.1186/s12931-020-01352-w">https://doi.org/10.1186/s12931-020-01352-w</a>	Clinic	<p><b>54 patients – 2 groups:</b> high and low NT-proBNP</p> <p>NT-proBNP at admission</p> <p><u>High group significantly:</u></p> <ul style="list-style-type: none"> <li>- Older</li> <li>- More comorbidities</li> <li>- Pro inflammatory</li> <li>- Lymphopenia</li> <li>- <b>Higher risk of death</b></li> </ul> <p>Cut-off of NT-proBNP for predicting <b>in hospital death</b>: 88,64 pg/ml (sensitivity: 100% - specificity: 66,67%)</p> <p>AUC for in hospital death = 0,909</p> <p><b>NT-proBNP = independent risk factors for in hospital death</b> (after adjusting)</p>
ACS nano 15APR2020	<b>Rapid Detection of COVID-19 Causative Virus (SARS-CoV-2) in Human Nasopharyngeal Swab Specimens Using Field-Effect Transistor-Based Biosensor</b>	Seo, Giwan and al Rep of Korea <a href="https://doi.org/10.1021/acsnano.0c02823">https://doi.org/10.1021/acsnano.0c02823</a>	Diagnostic	<p>Technology based on a field-effect transistor (FET)-based biosensing device for detecting SARS-CoV-2 in clinical samples. The sensor was produced by coating graphene sheets of the FET with a specific antibody against SARS-CoV-2 spike protein.</p> <p>The performance of the sensor was determined using antigen protein, cultured virus, and nasopharyngeal swab specimens from COVID-19 patients.</p> <p>CI*: the device is a highly sensitive immunological diagnostic method for COVID-19 that requires no sample pretreatment or labeling.</p>
Nature Medicine 15APR2020	<b>Temporal dynamics in viral shedding and transmissibility of COVID-19</b>	He, Xi; et al. China <a href="https://doi.org/10.1038/s41591-020-0869-5">https://doi.org/10.1038/s41591-020-0869-5</a>	Virology	<p>Pemporal viral shedding (94 patients with lab-confirmed COVID-19) and modeling of COVID-19 infectiousness profiles (separate 77 infector–infectee transmission pairs):</p> <ul style="list-style-type: none"> <li>- highest viral load in throat swabs was at the time of symptom onset.</li> <li>- 44% of secondary cases were infected during the infectors' presymptomatic stage, in settings with substantial household clustering, active case finding and quarantine outside the home.</li> </ul> <p>-&gt; control measures should be adjusted to account for substantial presymptomatic transmission.</p>
Journal of Biomolecular Structure and Dynamics 15APR2020	<b>Reverse vaccinology approach to design a novel multi-epitope vaccine candidate against COVID-19: an in silico study</b>	Maryam Enayatkhani et al. Iran <a href="https://doi.org/10.1080/07391102.2020.1756411">https://doi.org/10.1080/07391102.2020.1756411</a>	Vaccine	<p>3 known antigenic proteins of SARS-CoV-2 (Nucleocapsid, ORF3a, and Membrane protein)</p> <p>-&gt; used to predict <i>in silico</i> the potential immunogenic B and T-cell epitopes.</p> <p>-&gt; Prediction of best tertiary structure of selected epitopes docking TLR4 and HLA-A + Evaluation of the end the stability of complex of these receptors with the selected epitopes , by molecular Dynamic simulation methods.</p> <p>Antigenicity of the designed antigenic sequence -&gt; predicted by bioinformatic methods.</p> <p><b>The designed protein sequences without adjuvant were sufficient to produce an immune response. The allergenicity of the sequence was also predicted and this vaccine was not recognized as an allergen. As a conclusion, the engineered epitope could be considered as a possible vaccine candidate against COVID-19</b></p>
Nat Med 15APR2020	<b>Temporal dynamics in viral shedding and transmissibility of COVID-19</b>	He et al., China <a href="https://www.nature.com/articles/s41591-020-0869-5">https://www.nature.com/articles/s41591-020-0869-5</a>	Public Health/Epidemiology	<p>94 COVID-19 patients:</p> <ul style="list-style-type: none"> <li>- Highest Viral loads in Throat swabs <b>at time of symptom onset</b></li> <li>- Estimation: 44% (95% confidence interval, 25–69%) of secondary cases were infected during the index cases' <b>presymptomatic stage</b></li> </ul> <p>-&gt; Infectiousness started from 2.3 days (95% CI, 0.8–3.0 days) before symptom onset and peaked at 0.7 days (95% CI, –0.2–2.0 days) before symptom onset</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
ACS nano 14APR2020	<b>Computational Design of ACE2-Based Peptide Inhibitors of SARS-CoV-2</b>	Han, Yanxiao et al, USA <a href="https://doi.org/10.1021/acsnano.0c02857">https://doi.org/10.1021/acsnano.0c02857</a>	Therapeutics	<p>Design of <b>peptide inhibitors against the SARS-CoV-2</b>, mostly formed by two sequential self-supporting alpha-helices (bundle) extracted from the protease domain (PD) of angiotensin-converting enzyme 2 (ACE2), which bind to the SARS-CoV-2 receptor binding domains.</p> <p>Molecular dynamics simulations revealed that the alpha-helical peptides maintain their secondary structure and provide a <b>highly specific and stable binding (blocking) to SARS-CoV-2</b>. To provide a multivalent binding to the SARS-CoV-2 receptors, many such peptides could be attached to the surfaces of nanoparticle carriers.</p> <p>The proposed peptide inhibitors could provide simple and efficient therapeutics against the COVID-19 disease.</p>
Kidney International 14APR2020	<b>Renal histopathological analysis of 26 postmortem findings of patients with COVID-19 in China</b>	Su et al., China <a href="https://www.kidney-international.org/article/S0085-2538(20)30369-0/fulltext">https://www.kidney-international.org/article/S0085-2538(20)30369-0/fulltext</a>	Clinic	<p>Analyzing <b>kidney abnormalities in 26 autopsies</b> -&gt; Patients: respiratory failure associated with multiple organ dysfunction syndrome as the cause of death.</p> <p>9/26: <u>clinical signs</u> of kidney injury that included increased serum creatinine and/or new-onset proteinuria.</p> <p><u>Light microscopy</u>: diffuse proximal tubule injury with loss of brush border, non-isometric vacuolar degeneration, and even frank necrosis. + Occasional hemosiderin granules and pigmented casts. + Prominent erythrocyte aggregates obstructing the lumen of capillaries without platelet or fibrinoid material. + Absence of evidence of vasculitis, interstitial inflammation or hemorrhage.</p> <p><u>Electron microscopic</u>: clusters of coronavirus particles with distinctive spikes in the tubular epithelium and podocytes.</p> <p>ACE2 was found to be upregulated in patients with COVID-19, and immunostaining with SARS-CoV nucleoprotein antibody was positive in tubules.</p>
Journal of Autoimmunity 14APR2020	<b>Assessing ACE2 expression patterns in lung tissues in the pathogenesis of COVID-19</b>	Li, Guoping; et al. China <a href="https://doi.org/10.1016/j.jaut.2020.102463">https://doi.org/10.1016/j.jaut.2020.102463</a>	Fundamental research	<p>Data mining analysis (6 independent studies) of ACE2 expression in healthy population compared to patients with underlying diseases (chronic obstructive pulmonary diseases, asthma patients, smokers) :</p> <ul style="list-style-type: none"> <li>- no difference in ACE2 lung expression in healthy vs patients with chronic airway disease, suggesting no difference in susceptibility to SARS-CoV-2 infection.</li> <li>- long-term smokers have significantly greater ACE2 expression than healthy non-smokers (small airway epithelium), suggesting a risk factor for COVID-19.</li> <li>- ACE2 expression dramatically increased between 12-24h post SARS-CoV infection (airway epithelial cells), suggesting a role of ACE2 in post-infectious regulation.</li> </ul> <p>- In SARS-CoV infected cells, ACE2 expression significantly correlated with activation of neutrophils, NK cells, Th17 cells, Th2 cells, Th1 cells, dendritic cells and production of IL-1, IL-10, IL-6 and IL-8 (healthy non-smokers).</p> <p>Protein-protein regulation networks before and after infection identify:</p> <ul style="list-style-type: none"> <li>- ribosomal protein RPS3 plays a key role in viral replication.</li> <li>- non-receptor protein kinase SRC has a role in macrophage mediated innate immunity and cytokine release.</li> </ul> <p><b>Working hypothesis</b> -&gt; SARS-CoV-2 infection increases ACE2 expression, which affects RPS3 and SRC activity, two key hub genes involved in viral replication and inflammatory responses.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Disaster medicine and public health preparedness 14APR2020	<b>Public Education and Electronic Awareness of the New Coronavirus (COVID-19): Experiences from Iran</b>	Peyravi, M. et al, Iran <a href="https://doi.org/10.1017/dmp.2020.94">https://doi.org/10.1017/dmp.2020.94</a>	HSS/Politic	<p>When WHO declared a global health emergency, the Iranian Red Crescent Society and Ministry of Health took measures for public awareness (13 measures).</p> <ul style="list-style-type: none"> <li>- Training on preventive measures and how to deal with infection and exposure to patients are the most important steps to cope with COVID-19.</li> <li>- Take into account : new educational technologies and applications and the capacity of national and private media.</li> <li>- Necessity to develop content related to the individuals' ages. Adapt also to illiterate and disabled.</li> <li>- Messages adapted to certain special public groups (drivers, bakeries, ...) or social activities (shopping, ...) more impactful than some general advice.</li> </ul> <p><b>In the cyberspace</b></p> <ul style="list-style-type: none"> <li>- Effective and timely use of cyberspace=&gt; acceptance and dissemination</li> <li>- Better effectiveness of E-training than ordinary training.</li> <li>- Infographics (humor, animation and kids-friendly themes) viewed more and possibly higher effectiveness than monolog lectures.</li> <li>- Exponential growth in sharing the materials</li> </ul> <p><b>Conclusion :</b> to manage the virus, policymakers /organizations should provide innovative, unified and applied educational content to all people.</p>
BMJ 14APR2020	<b>COVID-19: why we need a national health and social care service</b>	Pollock, A. et al, UK <a href="https://doi.org/10.1136/bmj.m1465">https://doi.org/10.1136/bmj.m1465</a>	HSS/Politic	<p>Social services in the UK: most privatized and fragmented in Western world.</p> <p>High proportion of NHS workforce currently off work+pay very low. Sector short 120 000 workers. Emergency legislation in the UK curtailed rights to social care services of elderly, ill and disabled people in community/residential settings contrary to international law and common sense.</p> <p>Lack of access to, i.e., support after hospital discharge or mental health services: =&gt; more health crises &amp; hospital admissions + essential care workers taking time off to care for family members.</p> <p><b>Conclusion:</b> Universal integrated health and social care service =&gt; bring all services and staff under government control.</p> <p>=&gt; social care delivered by a trained and properly equipped workforce with decent terms of service. + mandate collection of data quantifying effect of COVID on social care sector.</p>
Ear, nose, & throat journal 13APR2020	<b>Clinical Presentation of COVID-19: A Systematic Review Focusing on Upper Airway Symptoms</b>	Lovato A and al, Italy <a href="https://doi.org/10.1177/0145561320920762">https://doi.org/10.1177/0145561320920762</a>	Clinic	<p>5 retrospective studies and cohort studies Quality of evidence = level 4 (low) <b>1556 patients: 57,5% males</b> <b>Mortality: 2,4% - ICU admission: 7,3%</b> <u>Upper airways symptoms:</u></p> <ul style="list-style-type: none"> <li>- Pharyngodynia: 12,4%</li> <li>- Nasal congestion: 3,7%</li> <li>- Rhinorrhea: 4% (1 study)</li> </ul> <p><b>None of the studies reported olfactory or gustative dysfunction</b></p> <p>Rest symptoms: same other study (fever, cough, fatigue) Alteration chest CT: 83% → bilateral++++ Severe cases: older, lymphopenia, radiologic abnormalities <u>Limits:</u> only hospitalized patients → not full clinical spectrum of COVID-19 / olfactory disorders could have been underestimated</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Mayo Clinic Proceedings 13APR2020	<b>ST-segment Elevation, Myocardial Injury, and Suspected or Confirmed COVID-19 Patients: Diagnostic and Treatment Uncertainties</b>	Bennett et al., USA <a href="https://doi.org/10.1016/j.jmayocp.2020.04.005">https://doi.org/10.1016/j.jmayocp.2020.04.005</a>	Diagnostic	<p>-&gt; For patients with COVID-19, the evaluation can be challenging due to reports of STE without obstructive coronary disease, which creates diagnostic and management challenges</p> <p>-&gt; Institutions to define acute cardiac care pathways which balance the risks of complicating COVID-19 patients from invasive therapies and unnecessary contrast exposure versus the potential benefit if the patient is experiencing a MI from acute coronary occlusion</p>
International Forum of Allergy & Rhinology 12APR2020	<b>Association of chemosensory dysfunction and Covid-19 in patients presenting with influenza-like symptoms</b>	Carol Y and al, USA <a href="https://doi.org/10.1002/air.22579">https://doi.org/10.1002/air.22579</a>	Clinic	<p><b>Cross sectional study – question survey – 2 groups</b> <b>All with influenza like symptoms</b> 59 COVID-19 positive and 203 COVID-19 negative Hospital admission low and comparable between groups</p> <p><u>Smell and taste loss</u>: more frequent in COVID-19 group</p> <ul style="list-style-type: none"> <li>- 68% and 71% versus 16% and 17% respectively</li> <li>- Largest magnitude of association with COVID-19</li> </ul> <p><b>Independently associated with COVID-19:</b></p> <ul style="list-style-type: none"> <li>- Anosmia: OR 10,9 [5,08 – 23,5]</li> <li>- Taste: OR 10,2 [4,74 – 22,1]</li> </ul> <p>Improvement of olfaction and taste that correlated with clinical resolution of illness.</p>
Travel Med. Infect. Dis. 11APR2020	<b>Clinical and microbiological effect of a combination of hydroxychloroquine and azithromycin in 80 COVID-19 patients with at least a six-day follow up: A pilot observational study</b>	Gautret, Philippe et al, France <a href="https://doi.org/10.1016/j.tmaid.2020.101663">https://doi.org/10.1016/j.tmaid.2020.101663</a>	Therapeutic	<p><b>Uncontrolled non-comparative observational study</b> in a cohort of <b>80 relatively mildly infected inpatients</b> treated with a combination of <b>hydroxychloroquine and azithromycin</b> over a period of at least three days.</p> <p>All patients <b>improved clinically</b> except one 86 year-old patient who died, and one 74 year-old patient still in intensive care. A <b>rapid fall of nasopharyngeal viral load</b> was noted, with 83% negative at Day7, and 93% at Day8. <b>Virus cultures from patient respiratory samples were negative in 97.5% of patients at Days5.</b></p> <p><b>Limitations:</b> descriptive pilot study in only 80 patients with relatively mild clinical presentation. No analytic approach to account for possible confounds including notably the severity of illness.</p>
The Lancet 11APR2020	<b>Centring sexual and reproductive health and justice in the global COVID-19 response</b>	Matthew J Harris et al., UK <a href="https://doi.org/10.1016/S0140-6736(20)30801-1">https://doi.org/10.1016/S0140-6736(20)30801-1</a>	HSS/Politic	<p>COVID-19 + existing sexual &amp; reproductive health inequities =&gt; women, girls and vulnerable populations' health, wellbeing and economic stability disproportionately impacted.</p> <p>1) COVID-19 = Increased risks for women</p> <ul style="list-style-type: none"> <li>- Women's risk factors of contracting COVID-19 may be higher = 70% of the global health and social care workforce worldwide,</li> <li>- Potential pregnancy-related complications</li> </ul> <p>2) Impact on sexual/reproductive health care</p> <ul style="list-style-type: none"> <li>- Disruption/Diversion of resources away from essential sexual/reproductive health care for COVID</li> <li>- Restrictive global policies that target vulnerable populations (Protecting Life in Global Health Assistance + migration policies of deterrence)</li> </ul> <p><b>Solutions:</b></p> <ul style="list-style-type: none"> <li>- Additional resources for sexual/reproductive health care + increase of telemedicine</li> <li>- Sex-disaggregated mortality and morbidity surveillance for COVID-19 research</li> <li>- Community driven efforts: recognize inequitable power structures + collaborative response</li> <li>- Eliminate legal/policy restrictions to sexual/reproductive health care.</li> </ul>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Journal of Clinical Virology 11APR2020	<b>Dynamic profile of RT-PCR findings from 301 COVID-19 patients in Wuhan, China: a descriptive study</b>	Xiao et al., China <a href="https://www.sciencedirect.com/science/article/pii/S1386653220300883?via%3Dihub">https://www.sciencedirect.com/science/article/pii/S1386653220300883?via%3Dihub</a>	Diagnostic	301 patients: -> median period between symptoms presence and positive SARS-CoV-2 RT-PCR results was 16 days -> median period between symptoms presence and an effective negative SARS-CoV-2 RT-PCR result was 20 days Although two consecutive negative results were confirmed in 70 patients, 30% of them had positive viral test results for the third time. Using specimens from nasal swabs to run the RT-PCR test showed a higher positive rate than using specimens from throat swabs.
Science 10APR2020	<b>Structure of the RNA-dependent RNA polymerase from COVID-19 virus</b>	Gao, Yan; et al. China - Australia <a href="https://doi.org/10.1126/science.abb7498">https://doi.org/10.1126/science.abb7498</a>	Structural biology	Cryo-EM structure of SARS-CoV-2 full-length RNA-dependent RNA polymerase (RdRp, also named nsp12) in complex with cofactors nsp7 and nsp8 (2.9-Å resolution) reveal:  - Conserved architecture of nsp12 with polymerase core of SARS-CoV, and resolution of a newly identified $\beta$ -hairpin domain at its N terminus.  - Comparative modeling reveals how remdesivir binds to nsp12 polymerase, its primary antiviral drug target.  -> provides basis for design of new antiviral therapeutics /cocktails targeting viral RdRp (nsp12).
Emerg. Infect. Dis. 10APR2020	<b>Aerosol and Surface Distribution of Severe Acute Respiratory Syndrome Coronavirus 2 in Hospital Wards, Wuhan, China, 2020</b>	Zhen-Dong Guo; et al. China <a href="https://doi.org/10.3201/eid2607.200885">https://doi.org/10.3201/eid2607.200885</a>	Virology	Samples taken from <b>potentially contaminated objects and air from an ICU</b> (housed 15 patients with severe COVID-19) and a general ward (housed 24 patients with milder COVID-19): - SARS-CoV-2 contamination greater in ICU than general ward. - Almost all positive samples concentrated in the contaminated areas. - Virus widely distributed on floors, computer mice, trash cans, sickbed handrails in both the ICU and general wards. - Virus-laden aerosols mainly concentrated near and downstream from patients, up to 4 m.  -> Atricter protective measures should be taken by medical staff working in the ICUs then general wards. -> Aerosol distribution in the general ward indicate transmission distance of SARS-CoV-2 might be 4 m
NEJM 10APR2020	<b>Compassionate Use of Remdesivir for Patients with Severe Covid-19</b>	Grein, Jonathan et al, USA <a href="https://doi.org/10.1056/NEJMoa2007016">https://doi.org/10.1056/NEJMoa2007016</a>	Therapeutic	<b>63 compassionate use of remdesivir for COVID patients</b> with either an oxygen saturation of 94% or less while breathing ambient air or need for oxygen support: 40 patients (75%) received the full 10-day course of remdesivir, 10 (19%) received 5 to 9 days of treatment, and 3 (6%) fewer than 5 days of treatment. During a median follow-up of 18 days, <b>36 patients (68%) had an improvement in oxygen-support class</b> , including 17 of 30 patients (57%) receiving mechanical ventilation who were extubated. <b>A total of 25 patients (47%) were discharged, and 7 patients (13%) died</b>  <b>Point of attention</b> : Measurement of efficacy will require ongoing randomized, placebo-controlled trials
Journal of Clinical Virology 10APR2020	<b>Clinical characteristics and risk assessment of newborns born to mothers with COVID-19</b>	Yang, Pu et al Chine <a href="https://doi.org/10.1016/j.jcv.2020.104356">https://doi.org/10.1016/j.jcv.2020.104356</a>	Clinic	Case report of <b>7 newborns</b> delivered by <b>SARS-CoV-2 infected pregnant women</b>  The current data show that the infection of <b>SARS-CoV-2 in late pregnant women does not cause adverse outcomes in their newborns</b>
Euro Surv 9APR2020	<b>Excess cases of influenza-like illnesses synchronous with coronavirus disease (COVID-19) epidemic, France, March 2020</b>	Boëlle, Pierre-Yves et al, France <a href="https://doi.org/10.2807/1560-7917.ES.2020.25.14.2000326">https://doi.org/10.2807/1560-7917.ES.2020.25.14.2000326</a>	HSS/Political	<b>Comparison of data from the Sentinelles network</b> monitors influenza-like illnesses (ILI) and acute respiratory infections (ARI) in general practice in France and <b>official COVID 19 reported cases in early March 2020 from the Santé Publique France</b>  It is estimated that <b>760 (95% CrI: 219–1,706) of acute respiratory infections</b> consultations in those older than 65 years in two regions of France (BFC and GRE) <b>could have been caused by COVID-19 during week 10.</b>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Clinical infectious disease 9APR2020	<b>Factors associated with prolonged viral RNA shedding in patients with COVID-19</b>	Xu K and al, China <a href="https://doi.org/10.1093/cid/ciaa351">https://doi.org/10.1093/cid/ciaa351</a>	Clinic	<p>Retrospective study – Two hospital – <b>113 patients</b></p> <p>Median age: 52 years – 58,4% were male</p> <p>28,3% were diagnosed as severe illness</p> <p>Median hospital stays: 15 days</p> <p><b>74,3% had viral RNA clearance within 21 days after illness onset</b> (median: 15 days)</p> <p><u>Prolonged RNA shedding:</u></p> <ul style="list-style-type: none"> <li>- Male (p=0,009)</li> <li>- Old age (p=0,033)</li> <li>- Concomitant hypertension (p=0,009)</li> <li>- Invasive mechanical ventilation (p=0,006)</li> <li>- Use of corticosteroid (p=0,025)</li> <li>- Delay recovery on radiological image (p&lt;0,001)</li> </ul> <p>→ <u>Multivariate analysis:</u></p> <ul style="list-style-type: none"> <li>- Male (OR: 3,24)</li> <li>- Delay hospital admission (OR: 1,30)</li> <li>- Invasive mechanical support</li> </ul> <p><u>Limitations:</u></p> <ul style="list-style-type: none"> <li>- Viral RNA shedding ≠ viral shedding</li> <li>- Didn't not evaluated the effect of the treatment</li> </ul>
Obesity 9APR2020	<b>High prevalence of obesity in severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) requiring invasive mechanical ventilation</b>	Simonnet A and al, France <a href="https://doi.org/10.1002/oby.22831">https://doi.org/10.1002/oby.22831</a>	Clinic	<p><b>Retrospective study</b> – 124 patients in ICU</p> <p>Control group: non-SARS-CoV2 in ICU</p> <p>Median age: 60 years – 73% male -15% died</p> <p>68,6% required invasive mechanical ventilation (IMV)</p> <p><b>Obesity and severe obesity were significantly more frequent in SARS-CoV2 patients (p&lt;0,001)</b></p> <p>Median BMI in SARS-CoV2 patients higher than in non-SARS-CoV2: 29,6 vs 24,0 (p&lt;0,001)</p> <p><u>IMV vs non IMV:</u></p> <ul style="list-style-type: none"> <li>- BMI higher in IMV group: 31,1 vs 27,0 (p&lt;0,001)</li> </ul> <p><b>Need for IMV gradually increase with BMI category.</b></p> <p>Patients with obesity should take extra measure to avoid COVID19 contamination.</p>
F1000 Research 9APR2020	<b>In silico identification of vaccine targets for 2019-nCoV</b>	Chloe H. Lee and Hashem Koohy UK <a href="https://doi.org/10.12688/f1000research.22507.1">https://doi.org/10.12688/f1000research.22507.1</a>	Vaccine	<p>The authors define computationally identified immunogenic and/or cross-reactive peptides from 2019-nCoV, based on comparison with immunogenic peptides deposited in the Immune Epitope Database and Analysis Resource (IEDB). They found:</p> <p>i) 28 SARS-derived peptides having exact matches in 2019-nCoV proteome previously characterized to be immunogenic by in vitro T cell assays (high affinity to HLA-A class I and II and targeted by CD8+ and CD4+ T cells)</p> <p>ii) 22 nCoV peptides having a high sequence similarity with immunogenic peptides but with a greater predicted immunogenicity score</p> <p>iii) 44 nCoV peptides predicted to be immunogenic by the iPred algorithm and 1G4 TCR positional weight matrices respectively (<i>de novo in silico</i> search of immunogenic peptides against the 2019-nCoV proteome sequence)</p>
Psychotherapy and psychosomatics 9APR2020	<b>Mental Health and Psychosocial Problems of Medical Health Workers during the COVID-19 Epidemic in China</b>	Hong-xing Wang et al., China <a href="https://doi.org/10.1159/000507639">https://doi.org/10.1159/000507639</a>	Psy	<p>Method : <b>online survey (2182 participants from China)</b></p> <p><b>Result :</b></p> <ul style="list-style-type: none"> <li>• higher prevalence rates of psychological symptoms among medical health workers = insomnia, anxiety, depression, somatization, and obsessive-compulsive symptoms</li> <li>• risk factors : having organic disease, living in rural areas, being female, and being at risk of contact with COVID-19 patients</li> </ul> <p>Main reasons :</p> <ul style="list-style-type: none"> <li>• insufficient understanding of the virus initially</li> <li>• lack of prevention and control knowledge</li> <li>• long-term workload</li> <li>• high risk of exposure to patients with COVID-19</li> <li>• shortage of medical protective equipment,</li> <li>• lack of rest</li> <li>• exposure to critical life events, such as death.</li> </ul> <p>Need for :</p> <ul style="list-style-type: none"> <li>• health protection and adequate working conditions: lowering job demands and workload / increasing job control and reward, medical protective equipment, adequate rest...</li> <li>• recovery programs focused on resilience and psychological well-being</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Clinical infectious diseases 9APR2020	<b>Comparative replication and immune activation profiles of SARS-CoV-2 and SARS-CoV in human lungs: an ex vivo study with implications for the pathogenesis of COVID-19</b>	Chu, Hin; et al. China <a href="https://doi.org/10.1093/cid/ciaa410">https://doi.org/10.1093/cid/ciaa410</a>	Virology	<p>Ex vivo human lung tissues infected with SARS-CoV-2 compared to SARS-CoV :</p> <ul style="list-style-type: none"> <li>- SARS-CoV-2 infected and replicated in human lung tissues more efficiently, generating 3.20 folds more infectious virus particles within 48hrs.</li> <li>- Both viruses were similar in cell tropism: both targeting types I and II pneumocytes, and alveolar macrophages.</li> </ul> <p>-&gt; Despite a more efficient virus replication in the infected human lung tissue, SARS-CoV-2 did not significantly induce types I, II, or III interferons, and only upregulated 38 % (= IL6, MCP1, CXCL1, CXCL5, and CXCL10) of the 13 key inflammatory mediators tested (in contrast to 85 % for SARS-CoV).</p>
The Lancet. Global health 9APR2020	<b>COVID-19 and risks to the supply and quality of tests, drugs, and vaccines</b>	Paul N Newton et al., US <a href="https://doi.org/10.1016/S2214-109X(20)30136-4">https://doi.org/10.1016/S2214-109X(20)30136-4</a>	HSS/Politic	<p>Production /supply chains for COVID-19 candidate drugs / essential medical products impaired. Supply chains for vital drugs for other diseases disrupted by repurposing without adequate evidence.</p> <ul style="list-style-type: none"> <li>• Substandard drugs driven by cost reduction</li> <li>• Falsified agents thrive on shortages, especially when buyers depart from regulated supply chains (masks, diagnostic tests, false claim of treatments...).</li> </ul> <p>When proven efficacious treatment, robust policies need to ensure prompt affordable, access for all people in need + quality assured, not diverted from other treatments:</p> <ul style="list-style-type: none"> <li>• Coordinated information-sharing among medicine regulators on authorizations for clinical trials</li> <li>• Ensure global manufacture + investigational interventions for unregistered + off label use</li> <li>• Comprehensive/rapid reporting of shortages of active ingredients and finished products</li> <li>• Robust evaluation of diagnostic tests</li> <li>• Innovative regional mechanisms (e.g. African Vaccine Regulatory Forum) for nations without robust regulatory systems</li> </ul>
NEJM 9APR2020	<b>Disease Control, Civil Liberties, and Mass Testing — Calibrating Restrictions during the Covid-19 Pandemic</b>	David M. Studdert et al., US <a href="https://doi.org/10.1056/NEJMp2007637">https://doi.org/10.1056/NEJMp2007637</a>	HSS/Politic	<p>Civil liberties: courts insist coercive restrictions must be 1) necessary, 2) crafted as narrowly as possible, 3) not used to target ostracized groups.</p> <p>Clear criteria for quarantine for other diseases don't apply to social restrictions for COVID-19:</p> <ol style="list-style-type: none"> <li>1) Quarantine is community-wide and applies to government and private actors;</li> <li>2) Transmission dynamics make it difficult to identify / target risk groups.</li> <li>3) Stay-at-home restrictions unlikely to be a one-shot deal</li> </ol> <p>=&gt; Need for a graduated approach to restrictive measures.</p> <p>Curfew laws most adapted: courts give leeway to government to adapt + see COVID-19 restrictions as more of a public policy than a legal issue.</p> <p>Tailor restrictions using credible person-level information =&gt; identify people most likely to transmit infection through population wide program of testing and surveillance.</p> <p>Aggregate test results at community+state level to dial up or down.</p> <p>=&gt;Federal, state and local governments to finance &amp; oversee + rely on hospitals, pharmacies, private labs, mobile health services for implementation + civil organizations to foster compliance.</p>
Life Sciences 9APR2020	<b>In silico studies on therapeutic agents for COVID-19: Drug repurposing approach</b>	Shah, Bhumi et al, India <a href="https://doi.org/10.1016/j.jlfs.2020.117652">https://doi.org/10.1016/j.jlfs.2020.117652</a>	Therapeutic	<p>61 molecules that are already being used in clinics or under clinical scrutiny as antiviral agents are surveyed via <b>docking study</b>.</p> <p>37 molecules were found to interact with &gt;2 protein structures of COVID-19. Among them, <b>HIV protease inhibitors</b> and <b>RNA-dependent RNA polymerase inhibitors</b> showed promising features of binding to COVID-19 enzyme. Along with these, Methisazone an inhibitor of protein synthesis, <b>CGP42112A</b> an angiotensin AT2 receptor agonist and <b>ABT450</b> an inhibitor of the non-structural protein 3-4A might become convenient treatment option as well against COVID-19.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
EuroSur 9APR2020	<b>An alternative workflow for molecular detection of SARS-CoV-2 – escape from the NA extraction kit-shortage, Copenhagen, Denmark, March 2020</b>	Fomsgaard et al., Denmark <a href="https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.14.2000398">https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.14.2000398</a>	Diagnostics	<p>Development of an alternative simple and fast workflow for molecular detection of SARS-CoV-2 that does not require NA extraction and could serve as an alternative in diagnostic laboratories to overcome chemical-based kit-shortages.</p> <p>Approach consists of heating samples at 98°C for 5 min</p> <p>This simplified heat-approach should not be for general use but only if the gold standard approaches are not available. Simply heating the samples could serve as an easy, fast and inexpensive alternative to chemical extraction kits, which would detect 97.4% of the COVID-19-positive patients with no false positives; however, there might be a small risk of false negatives, which could be minimised by performing the assay in duplicates</p>
Clin Inf Dis 9APR2020	<b>Prediction for Progression Risk in Patients with COVID-19 Pneumonia: the CALL Score</b>	Ji, Dong and al., China <a href="https://academic.oup.com/cid/advance-article/doi/10.1093/cid/cia414/5818317">https://academic.oup.com/cid/advance-article/doi/10.1093/cid/cia414/5818317</a>	Diagnostics	<p>Overall, 208 patients were divided into stable group (n=168, 80.8%) and progressive group (n=40,19.2%) based on whether their conditions worsened during the hospitalization.</p> <p>Comorbidity, older age, lower lymphocyte and higher lactate dehydrogenase were shown to be independent high-risk factors for COVID-19 progression.</p> <p>By incorporating these 4 factors a novel scoring model, named as CALL, was established and tested.</p> <p>Conclusion : using the CALL score model can help the clinicians to improve the therapeutic effect and reduce the mortality of COVID-19 with more accurate and reasonable resolutions on medical resources.</p>
Clin Inf Dis 8APR2020	<b>PCR Assays Turned Positive in 25 Discharged COVID-19 Patient</b>	Yuan, Jing and al., China <a href="https://doi.org/10.1093/cid/cia398">https://doi.org/10.1093/cid/cia398</a>	Clinic	<p>172 COVID-19 infected patients discharged from Hospital:</p> <ol style="list-style-type: none"> <li>(1) Normal body temperature for more than 3 consecutive days.</li> <li>(2) Significant reduction of respiratory symptoms evaluated by following indicators: cough and expectoration disappeared, normal ranges for inflammatory markers IL-6 and CRP, as well as oxygenation index <math>\geq 350</math>.</li> <li>(3) Substantial improvement over conventional chest radiography detection.</li> <li>(4) At least two consecutively negative results of RT-PCR testing separated by at least 24-hour interval.</li> </ol> <p>All discharged patients were required another 14 days of self-segregating at home for further observation.</p> <p>-&gt; 25 discharged patients sent to hospital again because of the positive RT-PCR results. They experienced an average of <math>7.32 \pm 3.86</math> days from their last negative RT-PCR result to turning positive again.</p> <p><b>Some immunological parameters such as D-dimer and absolute lymphocyte count, and even antibody test should be combined with RT-PCR negative test as additional measures to assure that infected patients have completely recovered and can be released from quarantine.</b></p>
Clinical Immunology 8APR2020	<b>Epigenetic dysregulation of ACE2 and interferon-regulated genes might suggest increased COVID-19 susceptibility and severity in lupus patients</b>	Sawalha, Amr H. et al. USA-China <a href="https://doi.org/10.1016/j.clim.2020.108410">https://doi.org/10.1016/j.clim.2020.108410</a>	Virology	<p>Patients with <b>systemic lupus erythematosus</b> might be especially prone to severe COVID-19, independent of their immunosuppressed state.</p> <ul style="list-style-type: none"> <li>- <b>ACE2 is hypomethylated and overexpressed in lupus T cells</b> suggesting an increased susceptibility to SARS-CoV-2 infection.</li> <li>- increased oxidative stress induced by viral infection <b>exacerbates ACE2 demethylation defect</b> in lupus and <b>may enhance viremia</b>.</li> </ul> <p>&gt; Maintaining disease remission in lupus patients is critical to prevent DNA demethylation and increased oxidative stress, which may exacerbate susceptibility to SARS-CoV-2 infection and likelihood of cytokine storm.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA pediatric 8APR2020	<b>Screening and Severity of Coronavirus Disease 2019 (COVID-19) in Children in Madrid, Spain</b>	Tagarro A and al, Spain <a href="https://doi.org/10.1001/jamapediatrics.2020.1346">https://doi.org/10.1001/jamapediatrics.2020.1346</a>	Clinic.	365 screened children and 41 were positive = <b>11%</b> Median age = 1 year 34% had upper respiratory tract infection – 127 % fever without source - 5% viral like pneumonia 60% were hospitalized and <b>9,7% were admitted to PICU and needed respiratory support</b> <b>No one died</b> Limitations: probably more hospitalisation because of an increase awareness of COVID-19.
The Lancet 8APR2020	<b>First-wave COVID-19 transmissibility and severity in China outside Hubei after control measures, and second-wave scenario planning: a modelling impact assessment</b>	Leung et al., China <a href="https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30746-7/fulltext?utm_campaign=tlcoronavirus20&amp;utm_source=twitter&amp;utm_medium=social">https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30746-7/fulltext?utm_campaign=tlcoronavirus20&amp;utm_source=twitter&amp;utm_medium=social</a>	Public Health/Epidemiology	-> The first wave of COVID-19 outside of Hubei has abated because of <b>aggressive non-pharmaceutical interventions</b> .  -> the $R_t$ decreased substantially since Jan 23, when control measures were implemented, and have since remained below 1. -> <b>Relaxing the interventions (resulting in <math>R_t &gt; 1</math>)</b> when the epidemic size was still small would increase the cumulative case count <b>exponentially as a function of relaxation duration</b> , even if aggressive interventions could subsequently push disease prevalence back to the baseline level.  Given the substantial risk of viral reintroduction, particularly from overseas importation, close monitoring of $R_t$ and cCFR is needed to inform strategies against a potential second wave to achieve an optimal balance between health and economic protection.
The European respiratory journal, 8APR2020	<b>Clinical characteristics and outcomes of hospitalised patients with COVID-19 treated in Hubei (epicenter) and outside Hubei (non-epicenter): A Nationwide Analysis of China</b>	Wen-hua Liang et al, China <a href="https://doi.org/10.1183/13993003.00562-2020">https://doi.org/10.1183/13993003.00562-2020</a>	Clinic	<b>1590 cases from 575 hospitals in 31 provincial administrative regions</b> were collected (core cohort). <b>The overall rate of severe cases and mortality was 16.0% and 3.2%, respectively, but</b>  Potential risk factors analysed using proportional hazard (PH) Cox regression models <b>Patients in Hubei</b> [severe event rate 23.0% versus 11.1%, death rate 7.3% versus 0.3%, hazards ratio (HR) for critical illness 1.59, 95%CI 1.05–2.41] <b>have a poorer prognosis compared with patients outside of Hubei after adjusting for age and comorbidity</b> This might be <b>attributed to the prolonged duration of symptom onset to hospitalization in the epicenter</b> .
Journal of biomolecular structure & dynamics, 8APR2020	<b>In-silico homology assisted identification of inhibitor of RNA binding against 2019-nCoV N-protein (N terminal domain)</b>	Sarma, Phulen et al, India <a href="https://doi.org/10.1080/07391102.2020.1753580">https://doi.org/10.1080/07391102.2020.1753580</a>	Therapeutic	Two NTD structures of N proteins were selected (2OFZ and 1SSK, 92% homology) for virtual screening of 56,079 compounds from Asinex and Maybridge library to identify top 15 hits for each of the targets based on “docking score” <b>This study suggests two important class of compounds, theophylline and pyrimidone derivatives as possible inhibitors of RNA binding to the N terminal domain of N protein of coronavirus, thus opening new avenues for in vitro validations</b>
Pediatric Critical Care 7APR2020	<b>Coronavirus Disease 2019 in Critically Ill Children: A Narrative Review of the Literature</b>	Ong J and al, Singapore/Italy/Canada <a href="https://journals.lww.com/pccmjournal/Abstract/onlinefirst/Coronavirus_Disease_2019_in_Critically_Ill_98057.aspx">https://journals.lww.com/pccmjournal/Abstract/onlinefirst/Coronavirus_Disease_2019_in_Critically_Ill_98057.aspx</a>	Clinic	Children account for a few proportions of COVID19 diseases <b>Not severely ill:</b> asymptomatic++++ or mild Infant under 1 year appear to have an increased risk of severe disease. <u>Spared from severe disease:</u> - Less lymphopenia: 3,5% ≠ 70% in adults, - Appear to be less pro-inflammatory than adults, - Variation of expression of ACE2 and activity with age could protect against lung injury. <u>Management:</u> - Noninvasive ventilation or high-flow nasal cannula would be preferred, - Intubation should be performed by experienced practitioner with PPE and rapid sequence induction - Used cuffed tubes (minimized air leak), - Avoided disruption to the ventilator circuit - Creation of a family liaison contact



Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet Public Health 07APR2020	<b>The French response to COVID-19: intrinsic difficulties at the interface of science, public health, and policy</b>	Moatti, Jean.P et al., France <a href="https://doi.org/10.1016/S2468-2667(20)30087-6">https://doi.org/10.1016/S2468-2667(20)30087-6</a>	HSS/Politic	French authorities appointed an advisory board of 11 scientists to help manage the crisis =>evidence-based policy but: 1) France did not have logistic capacity to promote mass testing. Rather than setting goal of scaling up, authorities argued systematic testing was not needed; 2) Maintaining second round of elections # social science literature established disaster communication should avoid dissonant incentives/double binds. 3) Referring to clinical trials to prove efficacy of hydroxychloroquine without considering alternative evaluation methods for quicker evidence => reduced ability of authorities to mitigate effects or rumors and regulate prescription practices. Setting up second experts' committee: implicit recognition of the intrinsic difficulties of directly using science in political management of a health crisis.
J Mol Diag PRE-PROOF 7MAR2020	<b>Development of Reverse Transcription Loop-mediated Isothermal Amplification (RT-LAMP) Assays Targeting SARS-CoV-2</b>	Park et al., Republic of Korea <a href="https://www.biorxiv.org/content/10.1101/2020.03.09.983064v1">https://www.biorxiv.org/content/10.1101/2020.03.09.983064v1</a>	Diagnostic	-> Development of <b>highly specific RT-LAMP assays</b> for detection of SARS-CoV-2.  -> Results of these RT-LAMP assays can be detected within 30 minutes after amplification reaction begin. -> Optimization of reaction conditions where LCV colorimetric detection method is applied that can be used for point-of-care tests.
Clinical infectious diseases 7APR2020	<b>Towards Optimization of Hydroxychloroquine Dosing in Intensive Care Unit COVID-19 Patients</b>	Perinel, Sophie et al, France <a href="https://doi.org/10.1093/cid/ciaa394">https://doi.org/10.1093/cid/ciaa394</a>	Therapeutic	<b>Prospective pharmacokinetic study : 13 patients in intensive care unit received 200 mg x 3 of oral HCQ daily</b> , mean age 68 y. 46% obese, 31% with moderate or severe renal failure HCQ levels >1 mg/L and <2 mg/L were considered to be therapeutic. 161 blood levels recorded. Simulations performed based on data from patients with rheumatoid arthritis.  <b>PK studies are needed to define the optimal dosing regimen.</b> Based on simulations, <b>a loading dose of 800 mg once daily on day 1, followed by 200 mg twice daily for 7 days is proposed</b>
Science Translational Medicine 6APR2020	<b>An orally bioavailable broad-spectrum antiviral inhibits SARS-CoV-2 in human airway epithelial cell cultures and multiple coronaviruses in mice</b>	Sheahan, Timothy P. et al. USA <a href="https://doi.org/10.1126/scitranslmed.abb5883">https://doi.org/10.1126/scitranslmed.abb5883</a>	Therapeutic	In human airway epithelial cell cultures : - Orally bioavailable ribonucleoside analog (NHC, EIDD-1931) has broad spectrum antiviral activity against SARS-CoV-2, MERS-CoV, SARS-CoV, and related zoonotic group 2b or 2c Bat-CoVs, and a coronavirus bearing resistance mutations to remdesivir.  In mice: - Prophylactic and therapeutic administration improved pulmonary function, reduced virus titer and weight loss (mice infected with SARS-CoV or MERS-CoV)  -> potency of NHC/EIDD-2801 against multiple coronaviruses and oral bioavailability makes it a potential effective antiviral against SARS-CoV-2
Journal of Medical Virology 6APR2020	<b>Tocilizumab treatment in COVID-19: a single center experience</b>	Luo, Pan et al, China <a href="https://doi.org/10.1002/jmv.25801">https://doi.org/10.1002/jmv.25801</a>	Therapeutic	<b>15 COVID-19 patients</b> under <b>Tocilizumab (TCZ)</b> therapy were <b>retrospectively assessed</b> . - TCZ treatment ameliorated the increased CRP in all patients rapidly, - The 4 critically ill patients who received an only single dose of TCZ -> 3 died and the CRP level in the rest one patient failed to return to normal range with a clinical outcome of disease aggravation. - Serum IL-6 level tended to further spiked firstly and then decreased after TCZ therapy in 10 patients. - Persistent and dramatic increase of IL-6 was observed in these 4 patients who failed treatment.  <b>=&gt; A single dose of TCZ seems to fail to improve the disease activity in critically ill patients</b> although it was used in combination with glucocorticoid. However, <b>repeated doses of TCZ might improve the condition of critically ill patients.</b> <b>Limitations:</b> small number of cases reported ; use of laboratory parameters to define the disease activity is still challenging ; treatment duration observed may not be sufficient to make a final conclusion.



Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 6APR2020	<b>Baseline Characteristics and Outcomes of 1591 Patients Infected With SARS-CoV-2 Admitted to ICUs of the Lombardy Region, Italy</b>	Grasselli G et al, Italy <a href="https://doi.org/10.1001/jama.2020.5394">https://doi.org/10.1001/jama.2020.5394</a>	Clinic	<p>Retrospective - 1591 patients COVID-19 – multicentric</p> <p><u>Demographic:</u></p> <ul style="list-style-type: none"> <li>- <b>82% male</b> - median age: 63 years</li> <li>- 68% had at least 1 comorbidity (HTA+++)</li> </ul> <p><u>Clinical data</u></p> <ul style="list-style-type: none"> <li>- 1150 patients required mechanical ventilation (higher than reported for other ICU patients)</li> <li>- <b>Median PEEP: 14 cmH<sub>2</sub>O</b></li> <li>- Median P<sub>aO2</sub>/F<sub>IO2</sub> = 166 (IQR:114-220), higher in young patients (&lt; 63 years)</li> <li>- <b>Mortality: 26%</b>, higher in older patients (15% vs 36%, p&lt;0,001)</li> <li>- Median length of stay: 9 days in ICU</li> </ul> <p><u>Limitation:</u></p> <ul style="list-style-type: none"> <li>- Short follow up → mortality rate could change?</li> <li>- Missing data for some patients</li> </ul>
Clin Chem 4APR2020	<b>Potential false-negative nucleic acid testing results for Severe Acute Respiratory Syndrome Coronavirus 2 from thermal inactivation of samples with low viral loads</b>	Pan et al., China <a href="https://academic.oup.com/clinchem/advance-article/doi/10.1093/clinchem/hvaa091/5815979">https://academic.oup.com/clinchem/advance-article/doi/10.1093/clinchem/hvaa091/5815979</a>	Diagnostic	<p>-&gt; Ct values are increased (<b>higher threshold for detection</b>) in specimens from diagnosed COVID-19 patients in RT-PCR tests <b>after thermal incubation</b>.</p> <p>-&gt; About half of the weak-positive samples (7 of 15 samples, 46.7%) were RT-PCR negative after heat inactivation in at least one parallel testing</p> <p>Thermal inactivation adversely affected the efficiency of RT-PCR for SARS-CoV-2 detection. Given the limited applicability associated with chemical inactivators, other approaches to ensure the overall protection of laboratory personnel need consideration.</p>
Inter J Of Infect Dis 3APR2020	<b>A first Case of Meningitis/Encephalitis associated with SARS-Coronavirus-2</b>	Moriguchi, Takeshi et al, Japan <a href="https://doi.org/10.1016/j.ijid.2020.03.062">https://doi.org/10.1016/j.ijid.2020.03.062</a>	Clinic	<p><b>Case report : 23-year old male, with seizure accompanied by unconsciousness.</b></p> <p>The specific <b>SARS-CoV-2 RNA</b> was not detected in the nasopharyngeal swab but <b>was detected in a CSF brain MRI</b> : hyperintensity along the wall of right lateral ventricle and hyperintense signal changes in the right mesial temporal lobe and hippocampus, <b>suggesting the possibility of SARS-CoV-2 meningitis</b></p> <p>Chest CT <b>small ground glass opacities</b></p> <p>At D15 : still ventilated and with impaired consciousness</p> <p><b>This case warns the physicians of patients who have CNS symptoms.</b></p>
The Journal of infection 3APR2020	<b>Neutrophil-to-lymphocyte ratio as an independent risk factor for mortality in hospitalized patients with COVID-19</b>	Liu et al., China <a href="https://www.journalofinfection.com/article/S0163-4453(20)30208-5/pdf">https://www.journalofinfection.com/article/S0163-4453(20)30208-5/pdf</a>	Clinic	<p>245 COVID-19 patients :</p> <p>-&gt; Multivariate analysis demonstrated that there was 8% higher risk of in-hospital mortality for each unit increase in NLR.</p> <p>-&gt; Compared with patients in the lowest tertile, the NLR of patients in the highest tertile had a 15.04-fold higher risk of death after adjustment for potential confounders</p> <p>-&gt; Fully adjusted OR for mortality was 1.10 in males for each unit increase of NLR</p> <p><b>NLR is an independent risk factor of the in-hospital mortality for COVID-19 patients especially for male.</b></p>
The Journal of infection 3APR2020	<b>Arbidol Monotherapy is Superior to Lopinavir/ritonavir in Treating COVID-19</b>	Zhu, Zhen et al, China <a href="https://doi.org/10.1016/j.jinf.2020.03.060">https://doi.org/10.1016/j.jinf.2020.03.060</a>	Therapeutic	<p>50 patients into 2 groups</p> <ul style="list-style-type: none"> <li>- <b>lopinavir/ritonavir group (34 cases)</b></li> <li>- <b>arbidol group (16 cases).</b></li> </ul> <p>Data from these patients were <b>retrospectively analyzed</b>.</p> <p><b>At D14 post admission: no viral load was detected in arbidol group.</b></p> <p><b>44.1%</b> of patients in <b>lopinavir/ritonavir</b> group had <b>positive RNA test</b> on day 14.</p> <p>Patients in the arbidol group had a shorter duration of positive RNA test.</p> <p>No apparent side effects were found in both groups.</p> <p><b>=&gt; Arbidol monotherapy may be superior to lopinavir/ritonavir in treating COVID-19.</b></p> <p>The sample size is the major limitation of this study.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
CELL preproof	<b>Inhibition of SARS-CoV-2 infections in engineered human tissues using clinical-grade soluble human ACE2</b>	Monteil, Kwon et al,	Therapeutic	ACE2 has now also been identified as a key receptor for SARS-CoV-2 infections and it has been proposed that inhibiting this interaction might be used in treating patients with COVID-19. Clinical grade human recombinant soluble ACE2 ( <b>hrsACE2</b> ) <b>reduced SARS-CoV-2 recovery from Vero cells by a factor of 1,000-5,000</b> . An equivalent mouse rsACE2 had no effect. SARS-CoV-2 can also directly infect engineered human blood vessel organoids and human kidney organoids, which can be inhibited by hrsACE2. <b>These data demonstrate that hrsACE2 can significantly block early stages of SARS-CoV-2 infections.</b>
Nature Medicine 3APR2020	<b>Respiratory virus shedding in exhaled breath and efficacy of face masks</b>	Leung, Nancy H. L. et al., <a href="https://doi.org/10.1038/s41591-020-0843-2">https://doi.org/10.1038/s41591-020-0843-2</a> China - USA	Virology	Detection of <b>virus RNA shedding in exhaled breath and coughs</b> of children and adults with acute respiratory illness (influenza (n=23-28 infected individuals), coronavirus (n=10-11) and rhinoviruses (n=36-32)). 5 µm particle size fractionation.  <b><u>Without face mask:</u></b>  - Viral RNA identified in 30%, 26% and 28% of respiratory droplets and 40%, 35% and 56% of aerosols collected while not wearing a face mask (coronavirus, influenza virus and rhinovirus-infected participants, respectively)  Surgical face masks significantly reduced detection of viral RNA for: - Influenza virus in respiratory droplets, but not in aerosol. - Coronavirus in aerosols (≤5 µm particles), and non-significant reduction in respiratory droplets.  Limitation: infectivity of detected virus not tested.  <b>--&gt; Surgical face masks could prevent transmission of human coronaviruses and influenza viruses from symptomatic individuals</b>
Antiviral Research 3APR2020	<b>Remdesivir, lopinavir, emetine, and homoharringtonine inhibit SARS-CoV-2 replication in vitro</b>	Choy, Yin-Lam Wong et al, Hong Kong <a href="https://doi.org/10.1016/j.antiviral.2020.104786">https://doi.org/10.1016/j.antiviral.2020.104786</a>	Therapeutic	Evaluation of the in vitro antiviral effect of compounds that were previously reported to inhibit coronavirus replication and compounds that are currently under evaluation in clinical trials for SARS-CoV-2 patients.  <b>=&gt; Antiviral effect of remdesivir, lopinavir, homoharringtonine, and emetine against SARS-CoV-2 virus in Vero E6 cells with the estimated 50% effective concentration at 23.15 µM, 26.63 µM, 2.55 µM and 0.46 µM, respectively.</b> <b>=&gt; Ribavirin or favipiravir that are currently evaluated under clinical trials showed no inhibition at 100 µM.</b> <b>=&gt; Synergy between remdesivir and emetine was observed, and remdesivir at 6.25 µM in combination with emetine at 0.195 µM may achieve 64.9% inhibition in viral yield.</b>  <b>Combinational therapy may help to reduce the effective concentration of compounds below the therapeutic plasma concentrations and provide better clinical benefits.</b>
International Journal of Antimicrobial Agents 3APR2020	<b>Structural and molecular modeling studies reveal a new mechanism of action of chloroquine and hydroxychloroquine against SARS-CoV-2 infection</b>	Fantini, Di Scala et al, France <a href="https://doi.org/10.1016/j.ijantimicag.2020.105960">https://doi.org/10.1016/j.ijantimicag.2020.105960</a>	Therapeutic	Identification of a <b>new mechanism of action of CLQ and CLQ-OH</b> supporting the use of these repositioned drugs to cure SARS-CoV-2 infected patients. Using a combination of structural and molecular modeling approaches : <b>=&gt; chloroquine (CLQ) binds sialic acids and gangliosides with high affinity.</b> <b>=&gt; New type of ganglioside-binding domain at the tip of the N-terminal domain of the SARS-CoV-2 spike (S) protein identified.</b> This domain (aa 111-158), which is fully conserved among clinical isolates worldwide, may <b>improve the attachment of the virus to lipid rafts and facilitate the contact with the ACE-2 receptor.</b> <b>=&gt; In presence of CLQ (or of the more active derivative hydroxychloroquine, CLQ-OH), the viral spike is no longer able to bind gangliosides.</b>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Journal of Thrombosis and Thrombolysis 3APR2020	<b>Difference of coagulation features between severe pneumonia induced by SARS-CoV2 and non-SARS-CoV2</b>	Shiyu Y et al, China <a href="https://doi.org/10.1007/s11239-020-02105-8">https://doi.org/10.1007/s11239-020-02105-8</a>	Clinic	Retrospective – 2 groups 449 patients COVID-19 and 104 severe pneumonia non-COVID 28 days mortality → higher in COVID group (29,8% vs 15,4%, p<0,005) Heparin treatment: 22% in COVID and 21% non-COVID Platelet count: higher in COVID group significantly Elevated D-Dimer (sixfold of upper limit of normal) was associated with poor prognosis only in COVID group <b>Anticoagulant therapy may benefit to selected COVID patients (elevated D-Dimer)?</b> <u>Limits:</u> - Retrospective - Influence of others therapies?
Circulation 3APR2020	<b>The Variety of Cardiovascular Presentations of COVID-19</b>	Fried J et al, USA <a href="https://doi.org/10.1161/CIRCULATIONAHA.120.047164">https://doi.org/10.1161/CIRCULATIONAHA.120.047164</a>	Clinic	<b>4 cases reports</b> - SARS-CoV2 infection should be in the differential of typical cardiac syndrome during pandemic event without infection signs - Myocarditis like presentations with COVID-19 → <b>further study</b> - Direct cardiac injury = result of viral invasion OR cytokine storm induced by SARS-CoV2 → toxic effect on myocardium - COVID-19 can cause decompensation of heart failure → mixed shock
JAMA 3APR2020	<b>Personal Risk and Societal Obligation Amidst COVID-19</b>	Tsai et al., USA <a href="https://doi.org/10.1001/jama.2020.5450">https://doi.org/10.1001/jama.2020.5450</a>	HSS/Politic	Health workers with <b>pre-existing medical conditions</b> /in <b>older age groups</b> are at <b>greater risk</b> of severe illness and death if exposed to COVID-19. => Telemedicine  Issue : <b>guilt</b> – putting other colleagues at risk; what risk acceptable as a necessary part of a path in medicine? Assessment takes an inherently individual path: no single rule can guide a physician's involvement in high-risk scenarios  <b>Comforted by :</b> 1) High need for virtual-based care 2) Sense of purpose in the community  How much risk in the career of medicine should be acceptable to physicians? → <b>Need for medical profession to balance the obligations and duties of this profession with physicians' fundamentally human limitations and fears</b>
Physical and Engineering Sciences in Medicine, 3APR2020	<b>Covid-19: automatic detection from X-ray images utilizing transfer learning with convolutional neural networks</b>	Apostolopoulos et al., Greece <a href="https://link.springer.com/article/10.1007/s13246-020-00865-4">https://link.springer.com/article/10.1007/s13246-020-00865-4</a>	Diagnostic	Technology evaluated is called <b>Transfer Learning</b>  <u>Two sets of X-Ray images from patients were used as follow:</u> - 1st set: a collection of 1427 X-ray images including 224 images with confirmed Covid-19 disease, 700 images with confirmed common bacterial pneumonia, and 504 images of normal conditions. - 2nd set : 224 images with confirmed Covid-19 disease, 714 images with confirmed bacterial and viral pneumonia, and 504 images of normal conditions.  Deep Learning with X-ray imaging would extract significant biomarkers related to the Covid-19 disease, while the best accuracy, sensitivity, and specificity obtained is 96.78%, 98.66%, and 96.46% respectively. Further research are needed to confirm the efficiency of this type of technology for automatic detection of Covid-19 cases; moreover, it is necessary to develop models capable of distinguishing Covid-19 cases from other similar viral cases, but also from a greater variety of common pneumonia or even physiological X-rays
Antiviral Research, 3APR2020	<b>The FDA-approved Drug Ivermectin inhibits the replication of SARS-CoV-2 in vitro</b>	Caly, Leon et al, Australia <a href="https://doi.org/10.1016/j.antiviral.2020.104787">https://doi.org/10.1016/j.antiviral.2020.104787</a>	Therapeutic	<b>Ivermectin is an inhibitor of the COVID-19 causative virus (SARS-CoV-2)</b> on Vero/hSLAM cells. A single treatment able to effect ~5000-fold reduction in virus at 48h in cell culture compared to control sample.  <b>Ivermectin is FDA-approved for parasitic infections, Ivermectin is widely available, due to its inclusion on the WHO model list of essential medicines</b>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet Microbe 2APR2020	<b>Stability of SARS-CoV-2 in different environmental conditions</b>	Chin, Alex W. H. et al., China <a href="https://doi.org/10.1016/S2666-5247(20)30003-3">https://doi.org/10.1016/S2666-5247(20)30003-3</a>	Virology	<p><u>Infectious SARS-CoV-2 Stability at different temperatures (in virus transport medium) :</u></p> <ul style="list-style-type: none"> <li>- highly stable at 4°C (only ~ 0.7 log-unit reduction of infectious titre on day 14)</li> <li>- at 70°C, virus inactivation reduced to 5 mins</li> <li>- On a surgical mask, infectious virus detectable on day 7 (~0.1% of the original inoculum).</li> <li>- Except from hand soap, no infectious virus detected after 5-min incubation at RT (22°C) with virucidal disinfectants (Household bleach (1:49), Ethanol (70%), Povidone-iodine (7.5%), ...).</li> </ul> <p><b>--&gt; SARS-CoV-2 can be highly stable in a favourable environment, but also susceptible to standard disinfection methods.</b></p>
Liver Int 2APR2020	<b>Clinical characteristics of Non-ICU hospitalized patients with coronavirus disease 2019 and liver injury : A Retrospective study</b>	Xie et al., Chine, <a href="https://doi.org/10.1111/iv.14449">https://doi.org/10.1111/iv.14449</a>	Clinic	<p><b>Retrospective study of 79 patients</b>, median age 60 years and 55.7% male. <b>29 had liver injury</b> (elevated ALT, AST and/or bilirubin)</p> <p>Multivariate analysis suggested that <b>CT scores was an independent predictor for liver injury</b>. Patients with liver injury stayed longer in the hospital.</p>
American journal of nephrology 2APR2020	<b>Coronavirus Disease 19 Infection Does Not Result in Acute Kidney Injury: An Analysis of 116 Hospitalized Patients from Wuhan, China</b>	Wang Lu et al, China <a href="https://doi.org/10.1159/000507471">https://doi.org/10.1159/000507471</a>	Clinic	<p><b>116 patients</b> – Retrospective study</p> <p>Median age: 54y and 58% male 40% severe pneumonia and 9% were ARDS 4,3% had CKD with long-term hemodialysis</p> <p><b>None patient meet criteria for AKI</b> during or after treatment 12 patients without CKD showed mild increase of BUN or serum creatinine (≠ criteria of AKI). Temporary abnormal renal function → injury due to hypoxemia? SARS-CoV2 RNA in urine sediment was positive in 4 patients Mortality: 6%</p> <p><b>Results are similar with study on SARS-CoV infection in 2003</b> <b>Be careful because ACE2 expression is high in kidney.</b></p>
Liver Int 2APR2020	<b>Liver impairment in COVID-19 patients: a retrospective analysis of 115 cases from a single center in Wuhan city, China</b>	Zhang Y et al, China <a href="https://doi.org/10.1111/iv.14455">https://doi.org/10.1111/iv.14455</a>	Clinic	<p><b>2 groups: 115 COVID-19 and 114 controls (community acquired pneumonia)</b></p> <p>Controls significantly older – no other difference <b>No difference between group in the level of ALT or AST</b> Majority of COVID19 had mild abnormalities COVID-19 had reduction of albumin Liver is not the main target organ</p> <p><u>Relationship with the disease progression:</u></p> <ul style="list-style-type: none"> <li>- Higher level of ALT or AST in severe cases than mild cases,</li> <li>- Higher total bilirubin in severe cases,</li> <li>- Lower level of albumin in severe cases,</li> </ul> <p><b>Liver function did not show an independent association with severe COVID19</b></p>
CDC 1APR2020	<b>Presymptomatic Transmission of SARS-CoV-2 — Singapore, January 23–March 16, 2020</b>	Wei et al., China <a href="https://www.cdc.gov/mmwr/volumes/69/wr/mm6914e1.htm?s_cid=mm6914e1_w#contribAff">https://www.cdc.gov/mmwr/volumes/69/wr/mm6914e1.htm?s_cid=mm6914e1_w#contribAff</a>	Public Health/Epidemiology	<p>-&gt; Identification of <b>7 clusters</b> of COVID-19 in Singapore in which <b>presymptomatic transmission</b> likely occurred and which may explain the occurrence of secondary cases</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet Neurology 1APR2020	<b>Guillain-Barré syndrome associated with SARS-CoV-2 infection: causality or coincidence?</b>	Zhao, Hua; et al. China <a href="https://www.thelancet.com/journals/lanneur/article/PIIS1474-4422(20)30109-5/fulltext">https://www.thelancet.com/journals/lanneur/article/PIIS1474-4422(20)30109-5/fulltext</a>	Clinic	<p>Patient was diagnosed with Guillain-Barré syndrome (autoimmune disorder damaging the peripheral nervous system resulting in aggravating muscle weakness) with lymphocytopenia and thrombocytopenia upon hospitalization.</p> <p>Patient then developed symptoms of SARS-CoV-2 on day 8 and tested +ve by RT-PCR.</p> <p>Given the temporal overlap, authors speculate that SARS-CoV-2 infection might have been responsible for the development of Guillain-Barré syndrome with a possible parainfectious profile.</p> <p>Limitations: patient was not tested for SARS-CoV-2 upon admission, so causality is not clear.</p> <p><b>-&gt;This is the first, and single case report. It only suggests a possible association and more cases are necessary to support a causal relationship.</b></p>
Nature 1APR2020	<b>Virological assessment of hospitalized patients with COVID-2019</b>	Wolfel et al. Germany <a href="https://www.nature.com/articles/s41586-020-2196-x">https://www.nature.com/articles/s41586-020-2196-x</a>	Virology	<p>A detailed virological analysis of 9 cases, providing proof of active virus replication in upper respiratory tract tissues.</p> <p>-&gt; Pharyngeal virus shedding: very high during 1st week of symptoms.</p> <p>-&gt; Infectious virus was readily isolated from throat- and lung-derived samples, but not from stool samples (in spite of high virus RNA concentration).</p> <p>-&gt; Blood and urine never yielded virus.</p> <p>Active replication in the throat is confirmed by viral replicative RNA intermediates in throat samples.</p> <p>Sequence-distinct virus populations were consistently detected in throat and lung samples from the same patient, proving independent replication.</p> <p>Shedding of viral RNA from sputum outlasted the end of symptoms.</p> <p>Seroconversion occurred after 7 days in 50% of patients (14 days in all), but was not followed by a rapid decline in viral load.</p> <p>COVID-19 can present as a mild upper respiratory tract illness.</p> <p><b>Active virus replication in the upper respiratory tract puts the prospects of COVID-19 containment in perspective.</b></p>
NEJM 01APR2020	<b>Ten Weeks to Crush the Curve</b>	Fineberg, Harvey v. et al. USA <a href="https://doi.org/10.1056/NEJMe2007263">https://doi.org/10.1056/NEJMe2007263</a>	HSS/Politik	<p><b>1.</b> Appoint a commander who reports to the President with powers and authority of the President to mobilize all civil and military means (same at the level of the state) + redeploy limited national supplies where most needed.</p> <p><b>2.</b> Perform millions of diagnostic tests over the next 2 weeks Organize dedicated clinical trial sites, physically separate from other health centers.</p> <p><b>3.</b> Provide all health workers with personal protective equipment.</p> <p><b>4.</b> Act on the basis of symptoms, examinations, viral RNA detection tests and exposures to differentiate the population into 5 groups to be treated accordingly : 1 / infected; 2 / presumed infected; 3 / exposed; 4 / unknown exposure/infection; 5 / recovered &amp; sufficiently immune. Hospitalize severely affected or high-risk individuals+ create quarantine centers.</p> <p>Identify the fifth group by tests to enable economy to restart quickly and safely.</p> <p><b>5.</b> Mobilize the entire population. With PPE for all health workers, deliver surgical masks and hand sanitizer to every American household. If everyone is wearing a mask, no stigma.</p> <p><b>6.</b> Learn through real-time, fundamental research.</p> <p>Over the long-term: Reinvigorate the public health infrastructure for future threats.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet) 01APR2020	<b>Racism and discrimination in COVID-19 responses</b>	Devakumar, D. et al., UK <a href="https://doi.org/10.1016/S0140-6736(20)30792-2">https://doi.org/10.1016/S0140-6736(20)30792-2</a>	HSS/Politic	<p><b>COVID 19 engenders fear =&gt; social, political racism and xenophobia with racialised/ discriminatory responses to fear + disproportionately affecting marginalised groups</b></p> <p>Social dimension : COVID could have been an equalizer but disproportionately affects people of color + migrants</p> <ol style="list-style-type: none"> <li>1) Microaggression/Violence towards different ethnic groups (i.e.: Chinese)</li> <li>2) lower socio-economic groups (limited access to healthcare + precarious jobs)</li> <li>3) Ethnic minority groups at greater risk (comorbidities)</li> <li>4) Migrants avoid hospitals for fear of identification/reporting</li> </ol> <p>Political dimension : Misappropriation of Covid-19 crisis for political purpose (racial discrimination, conflating public health restrictions and border policies + trade policies).</p> <p>Health protection relies on a well functioning health system with universal coverage, + social inclusion, justice, and solidarity</p>
Virol Sin 31MAR2020	<b>Inefficiency of Sera from Mice Treated with Pseudotyped SARS-CoV to Neutralize 2019-nCoV Infection</b>	Zezhong Liu et al., China <a href="https://doi.org/10.1007/s12250-020-00214-5">https://doi.org/10.1007/s12250-020-00214-5</a>	Therapeutic	<p><b>S proteins:</b></p> <ul style="list-style-type: none"> <li>- 76% homology SARS CoV / SARS CoV-2</li> <li>- 29% homology SARS CoV / MERSCoV.</li> </ul> <p><b>RBS:</b></p> <ul style="list-style-type: none"> <li>- Significantly different, even if the bind to the same receptor (ACE2).</li> </ul> <p><b>Cross-reaction of sera ?</b></p> <p>SARS-CoV and MERCoV pseudovirus expression S protein: produced and injected into BALBc mice.</p> <ul style="list-style-type: none"> <li>-&gt; Sera tested on ACE2 expressing 293T cells.</li> <li>-&gt; Effective neutralization for SARS-PsV-treated mice but not MERS-PsV treated mice.</li> </ul> <p>When SARS-PsV-treated mice was exposed to a SARS-CoV-2 pseudovirus -&gt; no neutralization effect was evidenced.</p> <p><b>It may not be practical to treat SARS-CoV-2 patients.</b></p>
Emerging microbes & infections 31MAR2020	<b>Renin-angiotensin system inhibitors improve the clinical outcomes of COVID-19 patients with hypertension</b>	Meng, Juan et al, Chine, <a href="https://doi.org/10.1080/22221751.2020.1746200">https://doi.org/10.1080/22221751.2020.1746200</a>	Therapeutic	<p>Retrospective study of <b>42 patients with treated hypertension</b> admitted in hospitalization for COVID 19.</p> <p>Before hospitalization, <b>17 were on angiotensin-converting enzyme inhibitors</b> (ACEIs) or angiotensin II type 1 receptor blockers (ARBs), 25 were on other drugs.</p> <p><b>Results :</b> in patients from the ACEI/ARB group :</p> <ul style="list-style-type: none"> <li>- <b>Less severe cases</b></li> <li>- <b>trend toward lower IL-6 levels</b></li> <li>- <b>increased CD3 and CD8 T cell counts</b></li> <li>- <b>peak viral load during hospitalization significantly lower</b></li> </ul> <p>ACEI/ARB therapy may attenuate the inflammatory response, potentially through the inhibition of IL-6 levels</p> <p><b>Point of attention :</b> retrospective study, small sample.</p>
Annals of internal medicine 30MAR2020	<b>A Rush to Judgment? Rapid Reporting and Dissemination of Results and Its Consequences Regarding the Use of Hydroxychloroquine for COVID-19</b>	Kim, Alfred H.J et al., USA <a href="https://doi.org/10.7326/M20-1223">https://doi.org/10.7326/M20-1223</a>	HSS/Politic	<p><b>Urgency: certain limits of this study are acceptable (small sample size, use of an unvalidated surrogate end point, lack of randomization or blinding, ...). But methodological flaws that may affect the validity of the results :</b></p> <p><b>Conclusion:</b></p> <ul style="list-style-type: none"> <li>-&gt; Sufficient justification to continue investigation of the efficacy and safety of HCQ in patients hospitalized with COVID-19.</li> <li>-&gt; No data currently to recommend the use of HCQ as a prophylaxis for COVID-19.</li> <li>-&gt;No recommendation of its use outside of marketing authorization until it is justified and offer is reinforced.</li> <li>-&gt; Risk of penury to patients with rheumatic diseases who depend on HCQ for their survival.</li> <li>-&gt; HCQ shortage will limit availability to patients with COVID-19 if efficacy truly established.</li> </ul>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Medicine in Drug Discovery – pre-Proof 22MAR2020	<b>Novel decoy cellular vaccine strategy utilizing transgenic antigen-expressing cells as immune presenter and adjuvant in vaccine prototype against SARS-CoV-2 virus</b>	Henry Ji et al., China <a href="https://doi.org/10.1016/j.medidd.2020.100026">https://doi.org/10.1016/j.medidd.2020.100026</a>	Vaccine	<p>S1 SARS-CoV-2 protein is expressed on the surface of K562 human myelogenous leukemia cells (HLA negative - highly sensitive to NK mediating kills):</p> <p>-&gt; Provides a means of targeting and activating an innate driver of the host adaptive immune response. -&gt; Stable clones are selected and irradiated to be formulated as vaccine product and administered via intramuscular or subcutaneous injection. This kind of cell vaccine can drive the host cellular immune response toward Th1, generating both potent cytotoxic T cell immunity against the major determinant of SARS-CoV-2 cellular entry and pathogenesis.</p> <p><b>This approach has already being used for cancer vaccine treatments inducing robust cellular and humoral anti-tumor immune responses.</b></p>
Journal of infectious Disease 31MAR2020	<b>Characteristics of peripheral lymphocyte subset alteration in COVID-19 pneumonia</b>	Fan W et al, China <a href="https://doi.org/10.1093/infdis/jiaa150">https://doi.org/10.1093/infdis/jiaa150</a>	Immunology	<p><b>60 patients – monocentric – total lymphocytes in COVID-19 were compared to healthy controls (HC)</b> Median age 60 y 32% were serious illness <u>Compared to HCs, COVID-19 had a decrease in:</u></p> <ul style="list-style-type: none"> <li>- Total lymphocytes</li> <li>- CD4 + - CD8+ - NK cells and B cells</li> </ul> <p><u>Serious compared to mild patient:</u></p> <ul style="list-style-type: none"> <li>- Decrease total lymphocytes, CD4+, CD8+ and B cells in serious patients</li> </ul> <p><u>Post-treatment:</u></p> <ul style="list-style-type: none"> <li>- Total lymphocytes, CD8+ and B cells increased significantly in responders</li> <li>- No significant change in non responder's</li> </ul> <p><b>CD8+ cells potential predictor for disease severity and poor clinical efficacy</b></p>
The Lancet ID 30MAR2020	<b>Estimates of the severity of coronavirus disease 2019: a model-based analysis</b>	Verity et al., UK <a href="https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30243-7/fulltext#">https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30243-7/fulltext#</a>	Public Health/Epidemiology	<p><b><u>Using data on 24 deaths that occurred in mainland China and 165 recoveries outside of China:</u></b></p> <ul style="list-style-type: none"> <li>- Mean duration from onset of symptoms to death : 17,8 days</li> <li>- Mean duration from onset to hospital discharge: 24,7 days</li> <li>- Crude case fatality ratio: 3,67%</li> </ul> <p>After further adjusting for demography and under-ascertainment:</p> <ul style="list-style-type: none"> <li>- Case fatality ratio: 1,38% / &lt;60 y : 0,32% / &gt;60y: 6,4% / &gt;80y: 13,4%</li> </ul> <p>Estimates of case fatality ratio from international cases stratified by age were consistent with those from China (see paper for data)</p> <p>Estimated overall infection fatality ratio for China: 0-66%, with an increasing profile with age. Estimates of the proportion of infected individuals likely to be hospitalised increased with age up to a maximum of 18-4% in those aged 80 years or older.</p>
Journal of Gastroenterology and Hepatology 27MAR2020	<b>Covid-19 and the Digestive System</b>	Wong S et al, China <a href="https://doi.org/10.1111/jgh.15047">https://doi.org/10.1111/jgh.15047</a>	Clinic	<p><b>Diarrhoea</b> (2 to 10%) and <b>nausea/vomiting</b> (1 to 10%) are the most frequent gastrointestinal symptoms. Early in the disease course: earlier than pyrexia <b>Liver injury:</b> abnormal level of ALAT and ASAT in 15 to 53 % of patients – <b>mild and transient</b> → microvesicular steatosis and mild lobular activity → direct viral infection of hepatocytes (ACE2 receptor) or drug toxicity or immune-related injury</p> <p><b>Possible tropism of SARS-CoV-2 for gastrointestinal tract:</b> ACE2 receptor <b>Faecal source: viral transmission ?</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet ID 27MAR2020	<b>Clinical and virological data of the first cases of COVID-19 in Europe: a case series</b>	Lescure et al., France <a href="https://doi.org/10.1016/S1473-3099(20)30200-0">https://doi.org/10.1016/S1473-3099(20)30200-0</a>	Clinic	<p><b>5 Patients:</b> 3 men: aged 31 years, 48 years, and 80 years – 2 women: aged 30 years and 46 years</p> <p><b>3 different clinical evolutions:</b></p> <ul style="list-style-type: none"> <li>- 2 paucisymptomatic women diagnosed within a day of exhibiting symptoms, with high nasopharyngeal titres of SARS-CoV-2 within the first 24 h of the illness onset and viral RNA detection in stools</li> <li>- A two-step disease progression in 2 young men, with a secondary worsening around 10 days after disease onset despite a decreasing viral load in nasopharyngeal samples</li> <li>- an 80-year-old man with a rapid evolution towards multiple organ failure and a persistent high viral load in lower and upper respiratory tract with systemic virus dissemination and virus detection in plasma.</li> </ul> <p>The 80-year-old patient died on day 14 of illness. All other patients had recovered and been discharged by Feb 19, 2020.</p>
Clinical Infectious Disease 27MAR2020	<b>Clinical characteristics of refractory COVID-19 pneumonia in Wuhan, China</b>	Mo P et al, China <a href="https://doi.org/10.1093/cid/ciaa270">https://doi.org/10.1093/cid/ciaa270</a>	Clinic	<p><b>155 patients</b> with median age of 54 years – <b>85 refractory COVID-19:</b></p> <ul style="list-style-type: none"> <li>- Older and more male (<math>p &lt; 0,05</math>)</li> <li>- More comorbidities: diabetes, cardiovascular disease, cerebrovascular disease (<math>p &lt; 0,05</math>)</li> <li>- Higher incidence of breath shortness and anorexia (<math>p &lt; 0,05</math>)</li> <li>- Bilateral pneumonia</li> <li>- Higher CRP, LDH, ASAT and neutrophile</li> </ul> <p><b>Risk factors:</b></p> <ul style="list-style-type: none"> <li>- <b>Male (OR: 2,3 [1,0-4,8]) and anorexia admission (OR:3,9 [1,1-13,4])</b></li> </ul> <p>Received more oxygen (OR: 3,0), corticosteroid (OR:2,32) <b>Protective factor:</b> fever on admission (OR: 0,33 [0,1 – 0,9])</p>
JAMA 27MAR2020	<b>Treatment of 5 critically ill patients with COVID-19 with convalescent plasma</b>	Shen C et al, China <a href="https://jamanetwork-com.proxy.insermbiblio.iust.fr/journals/jama/fullarticle/2763983">https://jamanetwork-com.proxy.insermbiblio.iust.fr/journals/jama/fullarticle/2763983</a>	Therapeutic	<p><b>5 patients:</b> severe pneumonia + <math>P_{AO_2}/F_{IO_2} &lt; 300</math> mmHg + currently or has been supported by mechanical ventilation All received antiviral agents and steroids <b>Administered between 10 and 22 days after admission</b></p> <p><b>After transfusion:</b></p> <ul style="list-style-type: none"> <li>- Ct value and viral load declined</li> <li>- Value of inflammatory biomarkers decreased</li> <li>- Clinical improvement: improved <math>P_{AO_2}/F_{IO_2}</math>, reduced body temperature, improved chest imaging</li> <li>- No longer required respiratory support by 9 days after transfusion</li> </ul> <p><b>Limitations:</b></p> <ul style="list-style-type: none"> <li>- No control group and small cases</li> <li>- Improved without transfusion? / Improvement related to transfusion or other therapies?</li> <li>- Late administration of transfusion: different timing would be associated with different outcomes?</li> </ul>
The Lancet 27MAR2020	<b>Historical linkages: epidemic threat, economic risk, and xenophobia</b>	White, A. et al., USA <a href="https://doi.org/10.1016/S0140-6736(20)30737-6">https://doi.org/10.1016/S0140-6736(20)30737-6</a>	HSS/Politic	<p>Global management of pandemic disease threats and global commerce historically linked:</p> <ul style="list-style-type: none"> <li>- History of international infectious disease control shaped by a distinctly European/US perspective prioritizing epidemic threats from colonial/post-colonial sites potentially affecting trade (</li> <li>=&gt; aggressive control in sites of epidemic outbreak and aggressive scrutiny of those deemed responsible.</li> </ul> <p>- Importance of colonial trade from Asia led to bias against people of Asian descent. “Chinese virus” connected to a long legacy of associating epidemic disease threat and trade with movement of Asian peoples.</p> <p><b>Aggressive racist and xenophobic responses in the name of health controls.</b></p> <ul style="list-style-type: none"> <li>- Concern for trading relationships central to US economic growth pivotal for US Congress to endorse creation of WHO.</li> <li>- Nations have recently aligned infectious disease control policy alongside concerns for national security.</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
J. Med. Virol. 26MAR2020	<b>Stability Issues of RT-PCR Testing of SARS-CoV-2 for Hospitalized Patients Clinically Diagnosed with COVID-19</b>	Li et al., China <a href="https://doi.org/10.1002/jmv.25786">https://doi.org/10.1002/jmv.25786</a>	Diagnostic	<p><u>610 hospitalized patients from Wuhan</u></p> <p>-&gt; High false negative rate of RT-PCR testing</p> <p>-&gt; RT-PCR results from several tests at different points were variable from the same patients during the course of diagnosis and treatment of these patients</p> <p><b>Clinical indicators such as CT images should also be used not only for diagnosis and treatment but also for isolation, recovery/discharge and transferring for hospitalized patients clinically diagnosed with COVID-19 during the current epidemic.</b></p>
JAMA 26MAR2020	<b>Antibodies in Infants Born to Mothers With COVID-19 Pneumonia</b>	Zeng et al., China <a href="https://jamanetwork.com/journals/jama/fullarticle/2763854">https://jamanetwork.com/journals/jama/fullarticle/2763854</a>	Clinic	<p>A novel study on 6 pregnant women and their infants confirm no maternal-infant transmission of SARS-CoV-2 based on reverse transcriptase–polymerase chain reaction (RT-PCR) and reveals the presence of antibodies in all newborns :</p> <ul style="list-style-type: none"> <li>· 6 mothers had mild clinical manifestations and had cesarean deliveries in their third trimester</li> <li>· Neonatal throat swabs and blood samples are negative by RT-PCR test</li> <li>· All 6 infants had IgG and IgM virus-specific antibodies in their serum and their mothers also had elevated levels of IgG and IgM</li> <li>· Inflammatory cytokine IL-6 was significantly increased in all infants.</li> </ul> <p><b>Point of care/conclusion</b></p> <p>The detection of high level of IgM in 2 infants, is not usually. Whether the placentas of women were damaged and abnormal or whether IgM could have been produced by the infant if the virus crossed the placenta need to be confirm in a larger cohort.</p>
BMJ 26MARS2020	<b>The world's largest refugee camp prepares for covid-19</b>	Gaia Vince, UK <a href="https://doi.org/10.1136/bmj.m1205">https://doi.org/10.1136/bmj.m1205</a>	HSS/Politic	<p><b><u>Biggest camp in Cox's Bazar (Bengladesh):</u></b></p> <ul style="list-style-type: none"> <li>- Nearly 1 million people live in overcrowded conditions.</li> <li>- Particularly vulnerable (physical distancing impossible).</li> </ul> <p>United Nations Refugee Agency coordinate efforts to increase hand washing, using community leaders to inform (imams and women group leaders).</p> <p>Other initiative for preparedness : creasion of isolation unit in the camp. Aid workers are credible after experience of managing other crisis (malaria, dengue, cholera...) in the camp since 2 years and a half.</p>
The Lancet Public Health 25MAR2020	<b>The Italian health system and the COVID-19 challenge</b>	Armocida et al., Italy <a href="https://doi.org/10.1016/S2468-2667(20)30074-8">https://doi.org/10.1016/S2468-2667(20)30074-8</a>	HSS/Politic	<p>In Italy, National Healthcare Service is regionally based, with <b>local authorities responsible for the organisation</b> and delivery of health services. Due to progressive privatisation and finance cuts, system close to collapse. 4 lessons to be learned :</p> <ul style="list-style-type: none"> <li>- <b>Decentralisation and fragmentation of health services seems to have restricted timely interventions and effectiveness</b></li> <li>- Health-care systems capacity and financing need to be more flexible in case of emergencies</li> <li>- Solid partnerships between the private and public sector should be institutionalised</li> <li>- Recruitment of HR must be planned and financed with a long-term vision</li> </ul>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Lancet 25MAR2020	<b>Clinical features and obstetric and neonatal outcomes of pregnant patients with COVID-19 in Wuhan, China: a retrospective, single-centre, descriptive study</b>	Yu N. et al, China <a href="https://www.thelancet.com/pdfs/journals/laninf/PIIS1473-3099(20)30176-6.pdf">https://www.thelancet.com/pdfs/journals/laninf/PIIS1473-3099(20)30176-6.pdf</a>	Clinic	<p><b>Pregnant patients with COVID 19 – no ICU :</b> Mean age gestational: 39 + 1 - All caesarean section Fever (6) - Cough (1) - Shortness of breath (1) - Diarrhea (1) Laboratory tests: - Elevated CRP (7) - Lymphopenia (5) – thrombopenia (2) - Elevated IL-6 (4) <b>Chest CT:</b> all pneumonia → bilateral (6), unilateral (1) Treatment: oxygen + antiviral + antibiotic (single or combination) + traditional medicine. Methylprednisolone for 5 after caesarean section. <b>Neonatal:</b> 3 stays in observation and 1 was positive for SARS-CoV2 with mild shortness of breath. At 28 days after birth: all child was healthy <b>No arguments for vertical transmission</b></p>
SCIENCE 25MAR2020	<b>The effect of human mobility and control measures on the COVID-19 epidemic in China</b>	Kraemer et al., UK <a href="https://science.sciencemag.org/content/early/2020/03/25/science.abb4218">https://science.sciencemag.org/content/early/2020/03/25/science.abb4218</a>	Public Health/Epidemiology	<p><b>Use of real-time mobility data from Wuhan and detailed case data including travel history</b> -&gt; Early: spatial distribution of COVID-19 cases in China was explained well by human mobility data -&gt; After implementation of control measures: this correlation dropped and growth rates became negative in most locations</p> <p><b>Travel restrictions are particularly useful in the early stage of an outbreak when it is confined to a certain area that acts as a major source. However, travel restrictions may be less effective once the outbreak is more widespread.</b></p>
Inter J of Infectious Diseases 25MAR2020	<b>Epidemiological, clinical characteristics of cases of SARS-CoV-2 infection with abnormal imaging findings</b>	Zhang X et al, China <a href="https://www.ijidonline.com/article/S1201-9712(20)30172-7/fulltext">https://www.ijidonline.com/article/S1201-9712(20)30172-7/fulltext</a>	Clinic	<p><b>645 patients with 72 no-pneumonia and 573 pneumonia</b> Bilateral lung disease: 432 (67%) <u>Group with pneumonia:</u> - Older: 46,6 vs 34,9 years - One coexisting underlying: 28,8% vs 16% - Less exposure to Wuhan or confirmed patient - Time from onset to COVID-19 was longer: 5 days vs 2 days - Symptoms: fever and cough - Lower lymphocyte, albumin and NaCl - Higher LDH and CRP <u>Predictive factor of severe pneumonia:</u> - Lymphopenia and higher creatinine - Shortness of breath</p>
Disaster medicine and public health preparedness 24MAR2020	<b>Chronology of COVID-19 cases on the Diamond Princess cruise ship and ethical considerations: a report from Japan</b>	Nakazawa, et al. Japan <a href="http://www.ncbi.nlm.nih.gov/pubmed/32207674">http://www.ncbi.nlm.nih.gov/pubmed/32207674</a>	HSS/Politics	<p><b>Ship = virus incubator + "international miniature company"</b> -&gt; Difficulty in testing such a large number of people of various origins and faiths <b>Recommendations of the article:</b> <b>Politically:</b> -&gt; Alert political decision-makers to the impact of multiple, contradictory, false or unconfirmed information on the health of confined passengers -&gt; Mobilize collective intelligence / academic consensus by involving a large number of experts</p> <p><b>In terms of ethics and public health:</b> -&gt; When is the principle of confinement at sea justified: human rights dilemma (ensuring minimum well-being for passengers and crew) / health security (preventing the spread of the virus on land) -&gt; Two criteria for authorizing a ship to dock or not = "1) the nation's geopolitical status + 2) the nation's ability to provide adequate health care". -&gt; Optimizing the material and psychological conditions of confinement on a ship: access to medication; psychological support ; means of communication with the outside world ; transparency and consistency of media information + take into account cultural differences</p> <p><b>Legally:</b> -&gt; Design and implement international regulations because an epidemic on board a ship should not be dealt with solely with regard to local policies (territory in which the ship is at anchor) -&gt; Strengthen international cooperation.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet Global Health 24MAR2020	<b>Early in the epidemic: impact of preprints on global discourse about COVID-19 transmissibility</b> COMMENT	Maimuna et al., USA <a href="https://doi.org/10.1016/S2214-109X(20)30113-3">https://doi.org/10.1016/S2214-109X(20)30113-3</a>	HSS/Politic	<p><b><u>Novelty of SARS-CoV-2, so scientists rushed to fill epidemiological, virological, and clinical knowledge gap</u></b></p> <p>-&gt; 50 new studies about the virus between January 10 and January 30 alone.</p> <p>Use of a simple method to plot the ten R0 estimations posted as preprints before publication of the first peer-reviewed study on Jan 29. Result of the peer review R0 estimations are very similar to those in the peer-reviewed studies published on and after Jan 29.</p> <p><b><u>Conclusions :</u></b></p> <ul style="list-style-type: none"> <li>- Powerful role preprints can have during public health crises because of the timeliness with which they can disseminate new information.</li> <li>- Use of preprint does not jeopardise future peer-reviewed publication (first step : preprint and then peer reviewed)</li> <li>- Impact of preprints on discourse and decision making to the ongoing COVID-19 outbreak (even if in some cases the preprints have conveyed erroneous ideas)</li> </ul>
Emerging microbes and Infections 24MAR2020	<b>Establishment and validation of a pseudovirus neutralization assay for SARS-CoV-2</b>	Jianhui Nie et al., China <a href="http://www.ncbi.nlm.nih.gov/pubmed/32207377">http://www.ncbi.nlm.nih.gov/pubmed/32207377</a>	Therapeutic	<p>Necessity of <b>handling SARS-CoV-2 in BSL-3 facilities and accessibility to virus strains -&gt; barriers to develop candidate vaccines and therapeutics.</b></p> <p>-&gt; Hence, development of a SARS-CoV-2 pseudovirus based in neutralization assays using S viral genes cloned into pcDAN3.1 plasmids. -&gt; Expressed in a VSV pseudoviral platform. -&gt; Huh7 cells plated at 5x10<sup>4</sup>/well were identified as the best cell system for SARS-CoV2 pseudovirus infection (inocula of 650 TCID50/well).</p> <p><b>When tested against the SARS-CoV-2 pseudovirus, SARS-CoV-2 convalescent patient sera showed high neutralizing potency, which underscore its potential as therapeutics.</b></p> <ul style="list-style-type: none"> <li>• at home = stress can be eased</li> <li>• in local hospitals/ collective medical observation centers = separated from caregivers (children infected/suspected of being infected or caregivers infected/dead) Potential consequences: 1) grief and fear + 2) potential mood disorders and psychosis/death by suicide in adulthood</li> </ul> <p><b>30% = post-traumatic stress disorder</b></p> <p><b>Chinese gov. strategies to prevent risks :</b></p> <ol style="list-style-type: none"> <li>1. nurses 24 h per day</li> <li>2. guidance by nutritionists for children's diets</li> <li>3. communication with parents any time</li> <li>4. citizens volunteering as temporary mothers</li> <li>5. 24 h free psychological counselling hotlines</li> </ol> <p><b>Guidelines issued: more communication time with parents; access to disease information via comic books and videos; regular activity schedule ; night lights and gifts for children; referrals for psychiatrists</b></p> <p><b>Need for :</b></p> <ol style="list-style-type: none"> <li>1. formal training for paediatric health-care workers</li> <li>2. evidence-based guidelines</li> <li>3. national collaborative networks (psychiatrists, psychotherapists, researchers, community volunteers)</li> <li>4. post-pandemic surveillance of children</li> </ol>
The Lancet Child & Adolescent Health 24MAR2020	<b>Mental health considerations for children quarantined because of COVID-19</b>	Liu, Jia Jia; Bao, Yanping et al., China <a href="https://doi.org/10.1016/S2352-4642(20)30096-1">https://doi.org/10.1016/S2352-4642(20)30096-1</a>	HSS/Politic	<p><b>30% = post-traumatic stress disorder</b></p> <p><b>Chinese gov. strategies to prevent risks :</b></p> <ol style="list-style-type: none"> <li>1. nurses 24 h per day</li> <li>2. guidance by nutritionists for children's diets</li> <li>3. communication with parents any time</li> <li>4. citizens volunteering as temporary mothers</li> <li>5. 24 h free psychological counselling hotlines</li> </ol> <p><b>Guidelines issued: more communication time with parents; access to disease information via comic books and videos; regular activity schedule ; night lights and gifts for children; referrals for psychiatrists</b></p> <p><b>Need for :</b></p> <ol style="list-style-type: none"> <li>1. formal training for paediatric health-care workers</li> <li>2. evidence-based guidelines</li> <li>3. national collaborative networks (psychiatrists, psychotherapists, researchers, community volunteers)</li> <li>4. post-pandemic surveillance of children</li> </ol>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet 23MAR2020	<b>Temporal profiles of viral load in posterior oropharyngeal saliva samples and serum antibody responses during infection by SARS-CoV-2: an observational cohort study</b>	Kai-Wang et al., China <a href="https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30196-1/fulltext">https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30196-1/fulltext</a>	Virology	<p><b>23 persons were included</b></p> <ul style="list-style-type: none"> <li>- Median viral load in posterior oropharyngeal saliva or other respiratory specimens at presentation was 5.2 log<sub>10</sub> copies per mL</li> <li>- Salivary viral load: <b>highest during the first week after symptom onset</b> and subsequently declined with time</li> <li>- In one patient, viral RNA was detected <b>25 days after symptom onset</b>.</li> <li>- <b>Older age</b> was correlated with <b>higher viral load</b></li> <li>- For 16 patients with serum samples available 14 days or longer after symptom onset, <b>rates of seropositivity</b> were 94% for <b>anti-NP IgG</b>, 88% for <b>anti-NP IgM</b>, 100% for <b>anti-RBD IgG</b>, and 94% for <b>anti-RBD IgM</b>.</li> <li>- <b>Anti-SARS-CoV-2-NP</b> or <b>anti-SARS-CoV-2-RBD IgG</b> levels correlated with <b>virus neutralisation titre</b>.</li> </ul>
JAMA 23MAR2020	<b>Ethics Committee Reviews of Applications for Research Studies at 1 Hospital in China During the 2019 Novel Coronavirus Epidemic</b>	Zhang H et al.- China <a href="https://doi.org/10.1001/jama.2020.4362">https://doi.org/10.1001/jama.2020.4362</a>	HSS/Politic	<ul style="list-style-type: none"> <li>- <b>Henan hospital</b>: designated to provide care to COVID-19 patients.</li> <li>- Hospital ethics committee organized <b>4 emergency video conference in 35 days</b>.</li> <li>- Projects evaluated <b>within 2,13 days after submission: more quickly</b> than other previous boards organized in an outbreak context.</li> <li>- 41 applications were reviewed; 6 were approved; 4 rejected; and 31 referred for modification because of lack statistical basis for sample size calculation, deficiencies in inclusion/exclusion criteria or issues related to consent form. <b>Although the rush, review standards were not lowered during the outbreak.</b></li> </ul>
Open Forum Infect Dis 21MAR2020	<b>High-dose intravenous immunoglobulin as a therapeutic option for deteriorating patients with Coronavirus Disease 2019</b>	Wei Cao and al, China <a href="https://doi.org/10.1093/ofid/ofaa102">https://doi.org/10.1093/ofid/ofaa102</a>	Therapeutic	<p><b>3 adults (56, 34 and 35 y)</b></p> <ul style="list-style-type: none"> <li>- treated by <b>25 grams per day for five days</b> of <b>immunoglobulins</b> at the time of respiratory distress initiation + <b>antibiotic</b></li> <li>- <b>temperature back to normal in one to two days, and breathing difficulties alleviating in 3-5 days</b></li> </ul> <p>Point of attention: other treatments were given, antiviral for 2/3 patients, corticoid for 1.</p> <p>The <b>first few days of deterioration</b> may present a <b>critical point</b> when <b>potent suppression of inflammatory cascade could save the patients</b> from fatal immune-mediated injuries</p> <p><b>Hospitalized patients</b> : i) <b>age &gt;12 years</b> and ii) <b>PCR documented SARS-CoV-2 carriage in nasopharyngeal sample at admission</b></p>
International journal of antimicrobial agents 20MAR2020	<b>Hydroxychloroquine and azithromycin as a treatment of COVID-19: results of an open-label non-randomized clinical trial</b>	Gautret, and al, France <a href="https://www.sciencedirect.com/science/article/pii/S0924857920300996">https://www.sciencedirect.com/science/article/pii/S0924857920300996</a>	Therapeutic	<ul style="list-style-type: none"> <li>- Treatment: <b>oral hydroxychloroquine sulfate</b> 200 mg, 3/day during 10 days.</li> <li>- 26 treated among them, six patients received additional azithromycin.</li> <li>- Control group : 16 patients from another centre or refusal to participate</li> <li>- 6 patients treated were excluded from the analysis</li> <li><b>Primary endpoint</b> : <b>virological clearance at day-6 post-inclusion</b></li> </ul> <p><b>70% of hydroxychloroquine-treated patients (N=20) were virologically cured comparing with 12.5% in the control group (N=16) (p= 0.001)</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Journal Travel Medicine and Infectious Disease 20MAR2020	<b>COVID-19: Active measures to support community-dwelling older adults</b>	K, Kuwahara et al., Japan <a href="http://www.ncbi.nlm.nih.gov/pubmed/32205272">http://www.ncbi.nlm.nih.gov/pubmed/32205272</a>	HSS/Politic	<p>-&gt; With no proven drug and vaccine treatments, non-pharmaceutical measures, especially social distancing, are an essential to slow the spread of the epidemic.</p> <p>-&gt; Given the higher risk associated with older adults, practical information should be provided to community-dwelling adults to help maintain appropriate community activity levels.</p> <p>-&gt; Issue of social isolation: efforts are needed to mitigate the negative psychological impact. recent technological advances may help detect and provide care for groups at high risk of social isolation. plans and measures to maintain social ties should be prepared at the individual level (family, friends, neighborhood, etc.), organizational or community levels, and societal level.</p>
Travel Med Infect Dis 20MAR2020	<b>Rapid viral diagnosis and ambulatory management of suspected COVID-19 cases presenting at the infectious diseases referral hospital in Marseille, France, - January 31st to March 1st, 2020: A respiratory virus snapshot</b>	Amrane et al, France <a href="http://www.ncbi.nlm.nih.gov/pubmed/32205269">http://www.ncbi.nlm.nih.gov/pubmed/32205269</a>	Public Health/Epidemio	<p>Rapid viral detection performed on sputum and nasopharyngeal samples from the first 280 patients suspected to have COVID-19.</p> <p>No SARS-CoV-2 was detected. Other viral infections were identified in 49% of the patients, with most common pathogens being influenza A and B viruses, rhinovirus, metapneumovirus and common coronaviruses, notably HKU1 and NL63.</p>
Cell Mol Immunol 19MAR2020	<b>Characterization of the receptor-binding domain (RBD) of 2019 novel coronavirus: implication for development of RBD protein as a viral attachment inhibitor and vaccine</b>	Wanbo Tai et al., China <a href="https://www.nature.com/proxy.insermbiblio.inist.fr/articles/s41423-020-0400-4">https://www.nature.com/proxy.insermbiblio.inist.fr/articles/s41423-020-0400-4</a>	Vaccine	<p>-&gt; SARS-CoV-2 <b>receptor-binding domain (RBD)</b> protein could be used <b>as a therapeutic agent</b> against SARS-CoV-2 and SARS-CoV infection (from results <i>invitro</i>)</p> <p>-&gt; RBD in SARS-CoV-2 S protein was identified</p> <p>-&gt; RBD protein bound strongly to human and bat angiotensin-converting enzyme 2 (ACE2) receptors.</p> <p>- SARS-CoV RBD-specific antibodies could crossreact with SARS-CoV-2 RBD protein</p> <p>- SARS-CoV RBD-induced antisera could cross-neutralize SARS-CoV-2 -&gt; <b>potential to develop SARS-CoV RBD-based vaccines for prevention of SARS-CoV-2 and SARS-CoV infection.</b></p>
NEJM 19MAR2020	<b>A trial of liponavir-ritonavir in adults hospitalized with severe Covid-19</b>	Cao B et al, China <a href="https://www.nejm.org/doi/pdf/10.1056/NEJMoa2001282?articleTools=true">https://www.nejm.org/doi/pdf/10.1056/NEJMoa2001282?articleTools=true</a>	Therapeutic	<p>Randomized, controlled trial, open-label trial</p> <p>☑ <b>199 patients included:</b> 99 received lopinavir-ritonavir and 100 standard care alone:</p> <ul style="list-style-type: none"> <li><b>Lopinavir-ritonavir was not associated with clinical improvement or mortality:</b> median time to clinical improvement 16 days vs 16 days, HR = 1.31 [0.95 – 1.85]</li> </ul> <p>Others outcomes:</p> <ul style="list-style-type: none"> <li>28-days mortality lower in the lopinavir-ritonavir group: 19.2% vs 25%, difference -5.8 % [-17.3 – 5.7]</li> <li>Detectable viral RNA for SARS-CoV2 was similar between two groups: 40.7 % of the patients of lopinavir-ritonavir group at the end of trial (28d)</li> <li>Serious adverse events: 19 in the lopinavir-ritonavir group (4 serious gastrointestinal adverse events related to the trial medication) and 32 in the standard care alone.</li> <li>No difference on duration of oxygen therapy and duration hospitalization.</li> <li>Post hoc finding that early initiation of lopinavir-ritonavir might accelerate clinical recovery and reduced mortality</li> </ul> <p>Overall mortality at 22.1%</p> <p><b>No benefit was observed with lopinavir-ritonavir treatment</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
NEJM 19MAR2020	<b>SARS-CoV2 Infection in children</b>	Lu X. et al, China <a href="https://www.nejm.org/doi/pdf/10.1056/NEJMc2005073?articleTools=true">https://www.nejm.org/doi/pdf/10.1056/NEJMc2005073?articleTools=true</a>	Clinic	On the 1391 children tested at Wuhan Children's Hospital, <b>171 (12.3%) were positive for SARS-CoV2 infection.</b>  Median age: <b>6.7 years</b> - Male: 60.8 % Fever: <b>41.5 %</b> - Cough: 48.5 % Pneumonia: 64.9 % <b>3 patients</b> (with coexisting conditions) <b>require intensive care and 1 death</b> <b>Most children appear to be mild symptomatic.</b>
World Journal of Pediatrics 19MAR2020	<b>Clinical features of severe pediatric patients with coronavirus disease 2019 in Wuhan: a single center's observational study</b>	Sun D et al, China <a href="https://link.springer.com.proxy.insermbiblio.inist.fr/content/pdf/10.1007/s12519-020-00354-4.pdf">https://link.springer.com.proxy.insermbiblio.inist.fr/content/pdf/10.1007/s12519-020-00354-4.pdf</a>	Clinic	<b>8 children included:</b> 5 severely ill and 3 critically ill 2 months to 15 years <b>Symptoms:</b> - Polypnea 100% - Fever (6/8) - Cough (6/8) - Expectoration (4/8) <b>Abnormalities in chest scanning 100% patients:</b> - multiple patch-like shadows - ground glass opacity <b>Biological:</b> - increase CRP, PCT and LDH - elevated ALAT - increase IL6 (2/8), IL10 (5/8), IFN-γ (2/8) Level of IL6 and IL10 were significantly increase in 2 critically ill patient who remained in ICU for 20 days. <b>Specific laboratory abnormalities and excessive immune responses may lead to long-term lung damage and severe health complication</b>
Cell and Mol Biol 17MAR2020	<b>Elevated exhaustion levels and reduced functional diversity of T cells in peripheral blood may predict severe progression in COVID-19 patients</b>	Zheng et al., <a href="https://www.nature.com/articles/s41423-020-0401-3">https://www.nature.com/articles/s41423-020-0401-3</a>	Immunology	Immunological characteristics of peripheral blood leukocytes from 16 patients:  <b>Compared to healthy group (n=6):</b> - Frequency of multi-functional CD4+ T cells (positive for at least two cytokines) decreased significantly in the severe group - The proportion of non-functional (IFN-γ-TNF-α-IL-2-) subsets increased significantly. - No increase in neutrophils or decrease in lymphocytes. - No statistical differences in interleukin-6 (IL-6) and tumor necrosis factor-α (TNF-α) plasma concentrations were found among the three groups  - Levels of interferon-γ (IFN-γ) and TNF-α in CD4+ T cells were lower in the severe group than in the mild group, whereas the levels of granzyme B and perforin in CD8+ T cells were higher in the severe group than in the mild group.  - Frequency of multi-functional CD4+ T cells decreased significantly in the severe group and proportion of non-functional subsets increased significantly  <b>-&gt; Identification of potential immunological risk factors for COVID-19 pneumonia and provided clues for its clinical treatment.</b>
The NEJM 17MAR2020	<b>Aerosol and Surface Stability of SARS-CoV-2 as Compared with SARS-CoV-1</b>	Doremalen et al., USA <a href="https://www.nejm.org/doi/pdf/10.1056/NEJMc2004973?articleTools=true">https://www.nejm.org/doi/pdf/10.1056/NEJMc2004973?articleTools=true</a>	Virology	-> Stability of <b>SARS-CoV-2</b> was similar to that of <b>SARS-CoV-1</b> under the experimental circumstances tested.  -> Detectable in <b>aerosols</b> for up to <b>three hours</b> , up to <b>four hours</b> on <b>copper</b> , up to <b>24 hours</b> on <b>cardboard</b> and up to <b>two to three days</b> on <b>plastic</b> and <b>stainless steel</b> .  <b>Aerosol and fomite transmission of SARS-CoV-2 is plausible</b>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet 17MAR2020	<b>Prevention of SARS-CoV-2 infection in patients with decompensated cirrhosis</b>	Xiao et al., China <a href="https://www.thelancet.com/journals/langas/article/PIIS2468-1253(20)30080-7/fulltext">https://www.thelancet.com/journals/langas/article/PIIS2468-1253(20)30080-7/fulltext</a>	Clinic	<p><b>Previously known:</b> Patients with decompensated cirrhosis have a higher risk of, and mortality from, infection.</p> <p>-&gt; 111 patients with decompensated cirrhosis (were included) -&gt; <b>New precautionary procedures</b> were implemented (<a href="#">see paper</a>) -&gt; Incidence of COVID19 was lower than in other groups.</p> <p>The simple approach (<a href="#">see paper</a>) could be an effective means of preventing COVID-19 in patients with decompensated cirrhosis.</p>
International journal of infectious diseases 17MAR2020	<b>Transmission potential and severity of COVID-19 in South Korea</b>	Shim et al., Rep of Korea <a href="https://www.ijidonline.com/article/S1201-9712(20)30150-8/fulltext">https://www.ijidonline.com/article/S1201-9712(20)30150-8/fulltext</a>	Public Health/Epidemiology	<p>- COVID-19 caused 6,284 cases and 42 deaths in South Korea as of March 8, 2020.</p> <p>- The mean reproduction number <math>R_t</math> of COVID-19 in Korea was estimated at 1.5 (95% CI: 1.4-1.6)</p> <p>- The intrinsic growth rate was estimated at 0.6 (95% CI: 0.6, 0.7) and the scaling of growth parameter was estimated at 0.8 (95% CI: 0.7, 0.8), indicating sub-exponential growth dynamics of COVID-19</p> <p>- The crude case fatality rate is higher among males (1.1%) compared to females (0.4%) and increases with older age, from 0.1% among those 30-39 yrs to 6% among those &gt;= 80 yrs as of March 6, 2020.</p> <p>- Results indicate early sustained transmission of COVID-19 in South Korea and support the implementation of social distancing measures to rapidly control the outbreak.</p>
J Inf Dis 17MAR2020	<b>Clinical outcome of 55 asymptomatic cases at the time of hospital admission infected with SARS-Coronavirus-2 in Shenzhen, China.</b>	Wang et al., China <a href="https://academic.oup.com/ijid/advance-article/doi/10.1093/infdis/jiaa119/5807958">https://academic.oup.com/ijid/advance-article/doi/10.1093/infdis/jiaa119/5807958</a>	Clinic	<p><b>55 asymptomatic carriers</b></p> <p><u>Conclusions:</u> -&gt; Asymptomatic carriers occurred <b>more often in middle aged</b> people who had <b>close contact</b> with infected family members -&gt; Majority of the cases <b>developed to be mild and ordinary COVID-19</b> during hospital</p>
Am J Transplant. 17MAR2020	<b>Successful recovery of COVID-19 pneumonia in a renal transplant recipient with long-term immunosuppression.</b>	Zhu et al., China <a href="https://onlinelibrary.wiley.com/doi/abs/10.1111/ajt.15869">https://onlinelibrary.wiley.com/doi/abs/10.1111/ajt.15869</a>	Clinic	<p><b>52-year-old man</b> who received <b>kidney transplantation 12 years ago</b></p> <p>-&gt; Clinical characteristics (symptoms, laboratory examinations, and chest CT) were <b>similar to those of non-transplanted COVID-19</b> patients -&gt; Following a treatment regimen: reduced immunosuppressant use and low dose methylprednisolone-based therapy</p> <p><b>Effectively treated case has reference value for the future treatment of other transplant patients with COVID-19 pneumonia. Analysis of additional cases is necessary to determine if this remains true.</b></p>
J Med Virol 17MAR2020	<b>Platelet-to-lymphocyte ratio is associated with prognosis in patients with Corona Virus Disease-19.</b>	Qu et al., China <a href="https://onlinelibrary.wiley.com/doi/abs/10.1002/jmv.25767">https://onlinelibrary.wiley.com/doi/abs/10.1002/jmv.25767</a>	Clinic	<p>-Retrospective analysis of <b>30 hospitalized patients</b> -&gt; Patients with platelet peaks during treatment: longer hospitalization. -&gt; Patients with platelet peaks were <b>older</b> -&gt; <b>Higher PLT</b> (platelet to lymphocyte ratio): <b>longer hospitalisation</b>. It may be related to <b>cytokine storm</b>.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet 17MAR2020	<b>Prisons and custodial settings are part of a comprehensive response to COVID-19</b>	Kinnet et al., Australia <a href="https://www.thelancet.com/journals/lanpub/article/PIIS2468-2667(20)30058-X/fulltext">https://www.thelancet.com/journals/lanpub/article/PIIS2468-2667(20)30058-X/fulltext</a>	Public Health/Epidemiology	<p><b>Prisons are epicentres for infectious diseases:</b></p> <ul style="list-style-type: none"> <li>- higher background prevalence of infection</li> <li>- higher levels of risk factors for infection</li> <li>- unavoidable close contact in often overcrowded, poorly ventilated, and unsanitary facilities,</li> <li>- poor access to health-care services relative to that in community settings</li> </ul> <p>-&gt; The public health importance of prison responses to influenza outbreaks has been recognised in the USA, where the Centers for Disease Control and Prevention have developed a checklist for pandemic influenza preparedness in correctional settings. WHO has also issued prison-specific guidance for responding to COVID-19.</p>
Arch Pathol Lab Med. 17MAR2020	<b>An Analysis of 38 Pregnant Women with COVID-19, Their Newborn Infants, and Maternal-Fetal Transmission of SARS-CoV-2: Maternal Coronavirus Infections and Pregnancy Outcomes</b>	Schwartz et al., USA <a href="https://www.archivesofpathology.org/doi/pdf/10.5858/arpa.2020-0901-SA">https://www.archivesofpathology.org/doi/pdf/10.5858/arpa.2020-0901-SA</a>	Clinic	<p>Analyzing literature describing 38 pregnant women with COVID-19 and their newborns in China</p> <p>-&gt; Unlike coronavirus infections of pregnant women caused by SARS and MERS, COVID-19 did not lead to maternal deaths</p> <p>-&gt; Similar to pregnancies with SARS and MERS: no confirmed cases of intrauterine transmission of SARS-CoV-2</p> <p><b>There is no evidence that SARS-CoV-2 undergoes intrauterine or transplacental transmission from infected pregnant women to their fetuses.</b></p>
Gynecologie, obstetrique, fertilité & senologie 16 MAR2020	<b>Infection with SARS-CoV-2 in pregnancy. Information and proposed care. CNGOF</b>	Peyronnet et al., France <a href="https://www.sciencedirect.com/science/article/pii/S2468718920301100?via=ihub">https://www.sciencedirect.com/science/article/pii/S2468718920301100?via=ihub</a>	Clinic	<p>Few pregnant women have been described</p> <p>Same symptoms as rest of adult's patients</p> <p>Some cases of ARDS or pneumonia</p> <p><b>2 pregnant women with invasive ventilation have been described</b></p> <p>Risk: cesarian and prematurity</p> <p>No miscarriage described</p> <p><b>Neonatal:</b></p> <ul style="list-style-type: none"> <li>- no case of vertical transmission</li> <li>- milder symptomatic</li> <li>- symptoms probably due to maternal hypoxemia</li> </ul>
Nat Med 16MAR2020	<b>Breadth of concomitant immune responses prior to patient recovery: a case report of non-severe COVID-19</b>	Thevarajan et al., Australia <a href="https://www.nature.com/articles/s41591-020-0819-2">https://www.nature.com/articles/s41591-020-0819-2</a>	Immunology	<p>-&gt; Kinetics of immune responses in relation to clinical and virological features of a patient with mild-to-moderate coronavirus disease 2019 (COVID-19) that required hospitalization.</p> <p><b>The emergence and rapid increase in activated CD38+HLA-DR+ T cells, especially CD8+ T cells, at days 7–9 preceded the resolution of symptoms:</b></p> <p>-&gt; ASCs appeared in the blood at the time of viral clearance (day 7; 1.48%) and peaked on day 8 (6.91%).</p> <p>-&gt; Emergence of cTFH cells in blood at day 7 (1.98%), increasing on day 8 (3.25%) and day 9 (4.46%)</p> <p>-&gt; The frequency of co-expression of CD38 and HLA-DR on CD8+ T cells increased in this patient from day 7 (3.57%) to day 8 (5.32%) and day 9 (11.8%) as well as the frequency of co-expression of CD38 and HLA-DR on CD4+ T cells between day 7 (0.55%) and day 9 (3.33%) although at lower levels than that of CD8+ T cells.</p> <p>-&gt; CD38+HLA-DR+ CD8+ T cells, produced larger amounts of granzymes A and B and perforin (~34–54% higher) than did their parent cells (CD8+or CD4+ populations).</p> <p>-&gt; Interestingly, minimal pro-inflammatory cytokines and chemokines were found in this patient with COVID-19, even at days 7–9.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
SCIENCE 16MAR2020	<b>Substantial undocumented infection facilitates the rapid dissemination of novel coronavirus(SARS-CoV2).</b>	Li et al, UK <a href="https://science.sciencemag.org/content/early/2020/03/13/science.abb3221.fulltext">https://science.sciencemag.org/content/early/2020/03/13/science.abb3221.fulltext</a>	Public Health/Epidemiology	<p>From observations of reported infection within China + mobility data + a networked dynamic metapopulation model and Bayesian inference</p> <p>-&gt; <b>86%</b> of all infections were undocumented (95% CI: [82%–90%]) prior to 23 January 2020 travel restrictions.</p> <p>-&gt; <b>Undocumented infections</b> were the infection <b>source for 79%</b> of documented cases</p> <p><b>It explain the rapid geographic spread of SARS-CoV2 and indicate containment of this virus will be particularly challenging</b></p>
The Lancet 16MAR2020	<b>Preparedness is essential for malaria-endemic regions during the COVID-19 pandemic</b>	Wang et al., China <a href="https://www.thelancet.com/journals/lanres/article/PIIS0140-6736(20)30561-4/fulltext">https://www.thelancet.com/journals/lanres/article/PIIS0140-6736(20)30561-4/fulltext</a>	Public Health/Epidemiology	<p>-&gt; Relevant lessons from the 2014–16 outbreak of Ebola virus disease in west Africa</p> <p>-&gt; Much like Ebola, the early symptoms of COVID-19, including fever, myalgia, and fatigue, might be confused with malaria and lead to challenges in early clinical diagnosis</p>
The Lancet 16MAR2020	<b>Screening of faecal microbiota transplant donors during the COVID-19 outbreak: suggestions for urgent updates from an international expert panel</b>	Ianiro et al., Italy <a href="https://www.thelancet.com/journals/lanres/article/PIIS0140-6736(20)30082-0/fulltext">https://www.thelancet.com/journals/lanres/article/PIIS0140-6736(20)30082-0/fulltext</a>	Public Health/Epidemiology	<p>-&gt; Before each donation, physicians should screen for two main items: the presence of typical COVID-19 symptoms</p> <p>-&gt; In endemic countries, the RT-PCR assay should be considered in all donors</p> <p>-&gt; Stool banks should retrospectively check the health status of the donor before using frozen faeces, according to local epidemiology, to avoid further potential spreading of SARS-CoV-2</p>
JAMA 13MAR2020	<b>Risk Factors Associated With Acute Respiratory Distress Syndrome and Death in Patients With Coronavirus Disease 2019 Pneumonia in Wuhan, China</b>	Wu et al., China <a href="https://jamanetwork.com/journals/jamainternalmedicine/fullarticle/2763184?resultClick=1">https://jamanetwork.com/journals/jamainternalmedicine/fullarticle/2763184?resultClick=1</a>	Clinic	<p>-&gt; <b>201 patients</b> included in the study</p> <p><b>Risk factors to develop ARDS:</b></p> <ul style="list-style-type: none"> <li>Older age, neutrophilia, and organ and coagulation dysfunction (eg, higher LDH and D-dimer)</li> <li>Associated with ARDS but not death: Comorbidities, lymphocyte counts, CD3 and CD4 T-cell counts, AST, prealbumin, creatinine, glucose, low-density lipoprotein, serum ferritin, PT</li> <li>Although <b>high fever</b> was <b>positively associated</b> with development of <b>ARDS</b>, it was <b>negatively related to death</b></li> <li><b>Higher CD3 and CD4 T-cell counts</b> might <b>protect</b> patients from developing ARDS</li> <li><b>Persistent and gradual increases in lymphocyte responses</b> might be required for effective immunity against SARS-CoV-2 infection.</li> </ul>
Euro Surveill 12MAR2020	<b>Retrospective analysis of the possibility of predicting the COVID-19 outbreak from Internet searches and social media data, China, 2020</b>	Li et al., China <a href="https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.10.2000199">https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.10.2000199</a>	Public Health/Epidemiology	<p><b>To predict the development of this outbreak as early and as reliably as possible</b></p> <p>-&gt; Data obtained from Google Trends, Baidu Index and Sina Weibo Index on searches for the keywords ‘coronavirus’ and ‘pneumonia’ correlated with the published NHC data on daily incidence of laboratory-confirmed and suspected cases of COVID-19, with the maximum <math>r &gt; 0.89</math>.</p> <p>-&gt; Peak interest for these keywords in Internet search engines and social media data was <b>10–14 days earlier than the incidence peak of COVID-19 published by the NHC</b>.</p> <p>-&gt; The lag correlation showed a maximum correlation at 8–12 days for laboratory-confirmed cases and 6–8 days for suspected cases</p>
The Lancet 12MAR2020	<b>SARS-CoV-2 RNA more readily detected in induced sputum than in throat swabs of convalescent COVID-19 patients</b>	Han et al., China <a href="https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30174-2/fulltext">https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30174-2/fulltext</a>	Diagnostic	<p>-&gt; 2 cases in <b>convalescence</b></p> <p>-&gt; Both <b>negative</b> with throat swab and anal swabs</p> <p>-&gt; <b>Positive</b> in <b>induced sputum</b></p> <p>To reduce the risk of disease spread, <b>viral RNA tests of induced sputum, not throat swabs</b>, should be assessed as a criterion for releasing COVID-19 patients.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet 12MAR2020	<b>Real estimates of mortality following COVID-19 infection</b>	Baud et al., Switzerland <a href="https://www.thelancet.com/action/showPdf?pii=S1473-3099%2820%2930195-X">https://www.thelancet.com/action/showPdf?pii=S1473-3099%2820%2930195-X</a>	Public Health/Epidemiology	<p>Mortality rate estimates are based on the number of deaths relative to number of confirmed cases of infection -&gt; <b>not representative of actual death rate.</b></p> <p><b>Real rates:</b></p> <ul style="list-style-type: none"> <li>- 5-6% for China</li> <li>- 15-2% outside China</li> </ul> <p><b>Current figures might underestimate the potential threat of COVID-19 in symptomatic patients</b></p>
The Lancet 11MAR2020	<b>Are patients with hypertension and diabetes mellitus at increased risk for COVID-19 infection?</b>	Fang et al., Switzerland <a href="https://www.thelancet.com/journals/lanres/article/PIIS2213-2600(20)30116-8/fulltext">https://www.thelancet.com/journals/lanres/article/PIIS2213-2600(20)30116-8/fulltext</a>	Clinic	<p>Patients with <b>cardiac diseases, hypertension, or diabetes</b>, who are treated with <b>ACE2-increasing drugs</b>, may be at <b>higher risk</b> for severe COVID-19 infection</p> <p>-&gt; They <b>should be monitored for ACE2-modulating medications</b>, such as ACE inhibitors or ARBs.</p> <p>-&gt; No evidence to suggest that <b>antihypertensive calcium channel blockers increased ACE2 expression or activity</b>: these could be a <b>suitable alternative treatment</b> in these patients.</p>
The Lancet 11MAR2020	<b>Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: a retrospective cohort study.</b>	Zhou et al., China <a href="https://www.thelancet.com/journals/lanres/article/PIIS2213-2600(20)30566-3/fulltext">https://www.thelancet.com/journals/lanres/article/PIIS2213-2600(20)30566-3/fulltext</a>	Clinic	<p>-&gt; 191 patients: 137 discharged and 54 died</p> <p>1- Comorbidity: 48%, with hypertension (30%), diabetes (19%), coronary heart disease (8%).</p> <p>2- Death associated with older age, higher SOFA score, d-dimer greater than 1 µg/mL on admission.</p> <p>3- Viral shedding: median 20 days in survivors, otherwise until death. Longest viral shedding: 37 days</p>
The Lancet 11MAR2020	<b>Early dynamics of transmission and control of COVID-19: a mathematical modelling study</b>	Kucharski et al., UK <a href="https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30144-4/fulltext">https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30144-4/fulltext</a>	Public Health/Epidemiology	<p>Calculation the <b>probability that newly introduced cases might generate outbreaks in other areas.</b></p> <p>-&gt; Estimations: The median daily reproduction number (<math>R_t</math>) in Wuhan <b>declined from 2.35</b> (95% CI 1.15–4.77) 1 week before travel restrictions were introduced on Jan 23, 2020, to <b>1.05</b> (0.41–2.39) 1 week after.</p> <p>-&gt; In locations with similar transmission potential to Wuhan in early January, <b>once there are at least four independently introduced cases</b>, there is a <b>more than 50% chance the infection will establish within that population.</b></p>
JAMA 11MAR2020	<b>Detection of SARS-CoV-2 in Different Types of Clinical Specimens</b>	Wang et al., China <a href="https://jamanetwork.com/journals/jama/fullarticle/2762997">https://jamanetwork.com/journals/jama/fullarticle/2762997</a>	Diagnostic	<p>-&gt; <b>1070 specimens collected from 205 patients</b></p> <p><b>POSITIVITY</b> by RT-PCR:</p> <ul style="list-style-type: none"> <li>Bronchoalveolar lavage fluid (<b>93%</b>)</li> <li>Sputum (<b>72%</b>)</li> <li>Nasal Swabs (<b>63%</b>)</li> <li>Fibrobronchoscope brush biopsy (<b>46%</b>)</li> <li>Pharyngeal swabs (<b>32%</b>)</li> <li>Feces (<b>29%</b>)</li> <li>Blood (<b>1%</b>)</li> <li>Urine (<b>0%</b>)</li> </ul>
Sci Rep 11MAR2020	<b>A high ATP concentration enhances the cooperative translocation of the SARS coronavirus helicase nsP13 in the unwinding of duplex RNA</b>	Jang et al., Republic of Korea <a href="https://www.nature.com/articles/s41598-020-61432-1">https://www.nature.com/articles/s41598-020-61432-1</a>	Fundamental Research	<p><b>To know: RNA Helicase nsP13 is essential for the viral RNA replication of the SARS coronavirus</b></p> <p><b>Here:</b></p> <p>-&gt;RNA helicase nsP13 would have higher binding affinity to RNA than to DNA, at same ATP concentrations.</p> <p>-&gt; The open state of nsP13 binding with a higher affinity to RNA than to DNA, is a considerably energy-consuming reaction</p> <p>-&gt;Unwinding of duplex RNA by nsP13 is a considerably energy-consuming reaction</p> <p><b>SARS coronavirus nsP13 may require more ATPs to promote stable helicase translocation necessary for delicate RNA replication.</b></p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Emerge Inf Dis 09MAR2020	<b>Detection of Novel Coronavirus by RT-PCR in Stool Specimen from Asymptomatic Child, China</b>	Tang et al., China <a href="https://wwwnc.cdc.gov/eid/article/26/6/20-0301_article">https://wwwnc.cdc.gov/eid/article/26/6/20-0301_article</a>	Public Health/Epidemio	<p>-&gt; <b>Asymptomatic child positive</b> for COVID-19 by RT-PCR in stool, <b>17 days after the last virus exposure</b></p> <p>-&gt; Still positive 9 days after that (in stool)</p> <p>-&gt; <b>Never positive in respiratory tracts specimens</b></p> <p>-&gt; no data on urine and blood</p> <p>-&gt; The child might have transmitted the virus to numerous persons. <b>Stool from COVID-19 patients might serve as another vehicle for virus transmission</b></p>
Clin Inf Dis 09MAR2020	<b>In Vitro Antiviral Activity and Projection of Optimized Dosing Design of Hydroxychloroquine for the Treatment of Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2)</b>	Yao et al., China <a href="https://academic.oup.com/cid/advance-article/doi/10.1093/cid/ciaa237/5801998">https://academic.oup.com/cid/advance-article/doi/10.1093/cid/ciaa237/5801998</a>	Therapeutic	<p>-&gt; Vero cells were treated by <b>Choloroquine and Hydroxychloroquine</b> before (prophylaxy) and after (anti-viral) infection by SARS-CoV-2.</p> <p>-&gt; <b>EC50</b> are calculated</p> <p>-&gt; <b>Hydroxychloroquine has superior antiviral and prophylactic activity than chloroquine</b></p> <p>-&gt; Physiologically-based pharmacokinetic (PBBPK) -&gt; to <b>predict</b> (in silico) <b>drug concentrations</b> in lung, plasma and blood.</p> <p>-PBBK model has acceptable prediction accuracy.</p> <p>-Kinetics were simulated with different scenari of dose regimens</p> <p>-Dose regiment was optimized (<b>recommendations</b>).</p>
Science 06MAR2020	<b>The effect of travel restrictions on the spread of the 2019 novel coronavirus (COVID-19) outbreak.</b>	Chinazzi et al., USA <a href="https://science.sciencemag.org/content/early/2020/03/05/science.aba9757.long">https://science.sciencemag.org/content/early/2020/03/05/science.aba9757.long</a>	Public Health/Epidemio	<p>-&gt; <b>Global metapopulation disease transmission model</b> to project the impact of travel limitations on the national and international spread of the epidemic.</p> <p>-&gt; <b>Travel quarantine of Wuhan</b> delayed the overall epidemic progression by <b>only 3 to 5 days in Mainland China</b></p> <p>-&gt; More marked effect <u>at the international scale</u>, where case importations were <b>reduced by nearly 80%</b> until mid February</p> <p>-&gt; Sustained 90% travel restrictions to and from Mainland China <b>only modestly affect</b> the epidemic trajectory <b>unless combined with a 50% or higher reduction of transmission in the community</b></p> <p>-&gt; Potential uses for the <b>definition of optimized containment schemes and mitigation policies</b> that includes <b>the local and international dimension</b> of the COVID-19 epidemic</p>
EuroSurveillance 05MAR2020	Evaluation of a quantitative RT-PCR assay for the detection of the emerging coronavirus SARS-CoV-2 using a high throughput system	Pfefferle et al. Germany <a href="https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7068162">https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7068162</a>	Diagnostic	<p>Assessment of a molecular assay for the detection of SARS-CoV-2 on a high-throughput platform, the cobas 6800, using the 'open channel' for integration of a laboratory-developed assay.</p> <p>Evaluated samples are swab samples.</p> <p>Good analytical performance in clinical specimens. The fully automated workflow enables high-throughput testing with minimal hands-on time, while offering fast and reliable results.</p> <p>Special notes : by its nature as a screening test targeting only a single viral gene, positive results should always be confirmed with an independent PCR as recommended]. Importance of closely coordinating with local reference centres and public health authorities for determining clinical indications for testing</p>
Cell 04MAR2020	<b>SARS-CoV-2 Cell Entry Depends on ACE2 and TMPRSS2 and Is Blocked by a Clinically Proven Protease Inhibitor</b>	Hoffman et al., Germany <a href="https://www.cell.com/cell/fulltext/S0092-8674(20)30229-4?_returnURL=https%3A%2F%2Flinkinghub.elsevier.com%2Fretrieve%2Fpii%2FS0092867420302294%3Fshoal%3Dtrue">https://www.cell.com/cell/fulltext/S0092-8674(20)30229-4?_returnURL=https%3A%2F%2Flinkinghub.elsevier.com%2Fretrieve%2Fpii%2FS0092867420302294%3Fshoal%3Dtrue</a>	Therapeutic	<p>-&gt; <b>Priming of S proteins</b> by host <b>cell proteases (TMPRSS2)</b> is <b>essential for viral entry</b> into cells.</p> <p>-&gt; <b>ACE 2 can be blocked</b> by a clinically proven <b>inhibitor of TMPRSS2</b></p> <p>-&gt; <b>The study suggests that antibody responses raised against SARS-CoV could at least partially protect against SARS- CoV-2 infection</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
Science 04MAR2020	<b>Structural basis for the recognition of the SARS-CoV-2 by full-length human ACE2</b>	Yan et al., China <a href="https://science.sciencemag.org/content/early/2020/03/03/science.abb2762/ta-b-pdf">https://science.sciencemag.org/content/early/2020/03/03/science.abb2762/ta-b-pdf</a>	Fundamental Research	<p>-&gt; <b>Cryo-EM structures of human ACE2</b>, in the presence of a neutral amino acid transporter B0AT1, with or without the receptor binding domain (RBD) of the surface spike glycoprotein (S protein) of SARS-CoV-2</p> <p>-&gt; <b>ACE2 may be a homodimer even in the absence of B0AT1</b></p> <p>-&gt; <b>A dimeric ACE2 can accommodate two S protein trimers</b>, each through a monomer of ACE2</p> <p>-&gt; Structure-based rational <b>design of binders with enhanced affinities to either ACE2 or the S protein of the coronaviruses</b> may facilitate development of <b>decoy ligands or neutralizing antibodies</b> for suppression of viral infection.</p>
J Clin Microbiol 04MAR2020	<b>Multicenter Evaluation of the QIAstat-Dx Respiratory Panel for the Detection of Viruses and Bacteria in Nasopharyngeal Swab Specimens</b>	Leber et al., USA <a href="https://jcm.asm.org/content/early/2020/02/28/JCM.00155-20.long">https://jcm.asm.org/content/early/2020/02/28/JCM.00155-20.long</a>	Diagnostic	<p>-&gt; <b>Multiplex <i>in vitro</i> diagnostic test</b> for the <b>qualitative</b> detection of <b>20 pathogens</b> directly from <b>nasopharyngeal swab specimens</b>.</p> <p>-&gt; Results available in approximately <b>69 minutes</b></p> <p>-&gt; <b>Pathogens identified:</b> adenovirus, coronavirus 229E, coronavirus HKU1, coronavirus NL63, coronavirus OC43, human metapneumovirus A+B, influenza A, influenza A H1, influenza A H3, influenza A H1N1/2009, influenza B, parainfluenza virus 1, parainfluenza virus 2, parainfluenza virus 3, parainfluenza virus 4, rhinovirus/enterovirus, respiratory syncytial virus A+B, Bordetella pertussis, Chlamydia pneumoniae and Mycoplasma pneumoniae</p> <p>-&gt; Compared to the BioFire FilmArray Respiratory Panel version 1.7: <b>percent agreement: 99,5% . negative percent agreement of ≥ 97.9%</b></p> <p><b>Robust and accurate assay for rapid, comprehensive testing for respiratory pathogens.</b></p>
Sci. China Life Sci. 04MAR2020	<b>Clinical characteristics of 24 asymptomatic infections with COVID-19 screened among close contacts in Nanjing, China</b>	<a href="https://link.springer.com/article/10.1007%2Fs11427-020-1661-4">https://link.springer.com/article/10.1007%2Fs11427-020-1661-4</a>	Clinic	<p>-&gt; Laboratory-confirmed positive for the COVID-19 (pharyngeal swab)</p> <p>-&gt; No obvious symptoms <b>at time of screening</b> (all of them)</p> <p>-&gt; <b>20.8%</b> developed symptoms (fever, cough, fatigue, etc.)</p> <p>-&gt; 50.0% cases showed typical CT images of ground-glass chest</p> <p>-&gt; 20.8% presented stripe shadowing in the lungs</p> <p>-&gt; 29.2% cases showed normal CT image and had no symptoms during hospitalization (<b>these cases were younger</b>)</p> <p>-&gt; <b>Epidemiological investigation revealed asymptomatic transmission</b></p>
JAMA 04MAR2020	<b>Air, Surface Environmental, and Personal Protective Equipment Contamination by Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) From a Symptomatic Patient</b>	Ong et al., Singapore <a href="https://jamanetwork.com/journals/jama/fullarticle/2762692?resultClick=1">https://jamanetwork.com/journals/jama/fullarticle/2762692?resultClick=1</a>	Public Health/Epidemiology	<p>Extensive <b>environmental contamination</b> by 1 SARS-CoV-2 patient with mild upper respiratory tract involvement</p> <p>-&gt; Toilet bowl and sink samples were positive</p> <p>-&gt; Swabs taken from the air exhaust outlets tested positive</p> <p>-&gt; Air samples were negative</p> <p>-&gt; Risk of transmission from contaminated footwear is likely low: negative results in the anteroom and clean corridor</p> <p><b>Limit of the study:</b> viral culture was not done to demonstrate viability</p>
Nat Sci Rev 03MAR2020	<b>On the origin and continuing evolution of SARS-CoV-2</b>	Tang et al., China <a href="https://academic.oup.com/nsr/advance-article/doi/10.1093/nsr/nwaa036/5775463?searchres=ult=1">https://academic.oup.com/nsr/advance-article/doi/10.1093/nsr/nwaa036/5775463?searchres=ult=1</a>	Genomic	<p>-&gt; Assessment of the <b>molecular phylogeny</b> and the divergence between <b>SARS-CoV-2</b> and <b>related coronaviruses</b>.</p> <p>-&gt; Population genetic analyses of 103 genomes of SARS-CoV-2 indicate that there are <b>two major types of viruses</b> (designated <b>L</b> and <b>S</b>) currently circulating between humans.</p> <p>-&gt; The <b>L type is predominant (70%)</b> against 30% for S type).</p> <p>-&gt; <b>This article suggests that the L type is more aggressive.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
JAMA 03MAR2020	<b>Epidemiologic Features and Clinical Course of Patients Infected With SARS-CoV-2 in Singapore</b>	Young et al., Singapore <a href="https://jamanetwork.com/journals/jama/fullarticle/2762688">https://jamanetwork.com/journals/jama/fullarticle/2762688</a>	Clinic	<p>-&gt; <b>18patients</b> diagnosed with SARS-CoV-2 infection in Singapore between January 23 and February 3, 2020</p> <p>-&gt; Respiratory tract infection with <b>prolonged viral shedding from the nasopharynx of 7 days or longer</b> in 15 patients (83%)</p> <p>-&gt; Supplemental oxygen was required in 6 patients (33%), <b>5 of whom</b> were treated with <b>lopinavir-ritonavir</b>, with <b>variable clinical outcomes</b> following treatment.</p>
Int J Infect Dis 02MAR2020	<b>Recurrence of positive SARS-CoV-2 RNA in COVID-19: A case report</b>	Chen et al., China <a href="https://www.ijidonline.com/article/S1201-9712(20)30122-3/pdf">https://www.ijidonline.com/article/S1201-9712(20)30122-3/pdf</a>	Virology	<p>- 46-year-old woman with multiple patchy ground glass opacities in bilateral subpleural areas by CT</p> <p>- <b>Oropharyngeal swab test was positive by RT-PCR.</b></p> <p>-&gt; Received <b>symptomatic treatment</b> and <b>antimicrobial therapy</b> including oseltamivir, arbidol, Lopinavir/ritonavir and moxifloxacin</p> <p>-&gt; 6 testing from 28 Jan to 17FEB, all negative <b>but one the 2FEB</b>. Discharged on 9FEB and testing remained negative during follow-up.</p> <p><b>SARS-CoV-2 RNA of respiratory tract specimen may be persistent or recurrent positive during the course.</b></p>
Jour of Infect 29FEB2020	<b>Identification of the hyper-variable genomic hotspot for the novel coronavirus SARS-CoV-2</b>	Wen et al., China <a href="https://www.journalofinfection.com/article/S0163-4453(20)30108-0/pdf">https://www.journalofinfection.com/article/S0163-4453(20)30108-0/pdf</a>	Genomic	<p>-&gt; Confirmation of the relationship of SARS-CoV-2 with other beta coronaviruses on the amino acid level.</p> <p>-&gt; Hyper-variable genomic hotspot established in SARS-CoV-2 <b>population at the nucleotide but not the amino acid level</b> -&gt; means <b>no beneficial mutations</b>.</p> <p>-&gt; <b>Mutations in nsp1, nsp3, nsp15, and gene S would be associated with the SARS-CoV-2 epidemic (compared with RaTG13) / required for human adaptation?</b></p>
J Med Virol 28FEB2020	<b>Development of Epitope-Based Peptide Vaccine Against Novel Coronavirus 2019 (SARS-COV-2): Immunoinformatics Approach</b>	Bhattacharya et al., India <a href="https://onlinelibrary.wiley.com/doi/abs/10.1002/jmv.25736">https://onlinelibrary.wiley.com/doi/abs/10.1002/jmv.25736</a>	Vaccine	<p>1-&gt; Characterization of the <b>spike glycoprotein</b> to obtain immunogenic epitopes</p> <p>2-&gt; Immunoinformatic analysis of 13 MHC I and 3 MHC II epitopes which <b>have antigenic properties</b></p> <p>3-&gt; These identified epitopes are candidate to formulate a <b>multi-epitopic peptide vaccine</b>.</p> <p><b>Need for <i>in vitro</i> and <i>in vivo</i> validations</b></p>
The NEJM 28FEB2020	<b>Clinical Characteristics of Coronavirus Disease 2019 in China</b>	Ni et al., China <a href="https://www.nejm.org/doi/pdf/10.1056/NEJMoa2002032?articleTools=true&amp;downloadfile=showPdf&amp;articleTools=true&amp;doi=10.1056/NEJMoa2002032">https://www.nejm.org/doi/pdf/10.1056/NEJMoa2002032?articleTools=true&amp;downloadfile=showPdf&amp;articleTools=true&amp;doi=10.1056/NEJMoa2002032</a>	Clinic	<p>Median age : <b>47 years</b> / Female: 41.9%</p> <p><b>Primary composite end point</b> (admission in ICU, use of mechanical ventilation and death) in <b>6.1%</b>, with <b>5.0% in ICU</b>, <b>2.3% with invasive mechanical ventilation</b>, and <b>1.4% who died</b>.</p> <p>History of direct contact with <b>wildlife: 1.9%</b></p> <p>Among nonresidents of Wuhan, <b>72.3% had contact with residents of Wuhan</b>, including 31.3% who had visited the city.</p> <p>Most common symptoms: fever (43.8% on admission and 88.7% during hospitalization) and cough (67.8%). Diarrhea was uncommon (3.8%).</p> <p>Median <b>incubation period: 4 days</b> (interquartile range, 2 to 7).</p> <p><b>CT:</b> ground-glass opacity was the most common radiologic: 56.4%.</p> <p><b>No radiographic or CT abnormality:</b> 17.9% with nonsevere disease and 2.9% with severe disease.</p> <p><b>Lymphocytopenia: 83.2%</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
EuroSurv 27FEB2020	<b>Early transmission patterns of coronavirus disease 2019 (COVID-19) in travellers from Wuhan to Thailand, January 2020</b>	Okada et al., Thailand <a href="https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.8.2000097">https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.8.2000097</a>	Public Health/Epidemiology	<p>-&gt; 2 women arriving in Thailand at different times (8 and 13 January)</p> <p>-&gt; The <b>two viral genomes are identical</b> to four sequences from Wuhan, while no direct link to the Huanan Seafood Market.</p> <p>-&gt; <b>Identical genomes of up to 30 kb are rare and a strong sign of recent transmission linkage</b></p> <p>-&gt; Data suggest that <b>transmission within Wuhan beyond the Huanan Seafood Market is likely to have occurred in the first week of January or earlier.</b></p>
J Clin Med 27FEB2020	<b>Epidemiological Identification of A Novel Pathogen in Real Time: Analysis of the Atypical Pneumonia Outbreak in Wuhan, China, 2019—2020</b>	Jung et al., Japan <a href="https://www.mdpi.com/2077-0383/9/3/637">https://www.mdpi.com/2077-0383/9/3/637</a>	Public Health/Epidemiology	<p>-&gt; <b>Non-virological descriptive characteristics</b> could have determined that the outbreak is caused by a novel pathogen in <b>advance of virological testing.</b></p> <p>-&gt; Characteristics of the outbreak <b>were collected in real time and compared with characteristics of eleven pathogens</b> that have previously caused cases of atypical pneumonia.</p> <p>-&gt; The <b>probability that a new virus was driving</b> the outbreak was assessed as <b>over 29%</b> on 31 December 2019, <b>one week before virus identification.</b></p>
The Lancet 27FEB2020	<b>Secondary attack rate and superspreading events for SARS-CoV-2</b>	Liu et al., UK <a href="https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30462-1/fulltext">https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30462-1/fulltext</a>	Public Health/Epidemiology	<p><b>The Ro value</b> only captures the average dynamics of transmission.</p> <p><b>The secondary attack rate (SAR)</b> is the probability that an infection occurs among susceptible people <b>within a specific group.</b></p> <p><b>SAR among close contacts would be of 35%</b> (95% CI 27–44).</p> <p>-&gt; An infection with a high household SAR but a modest R0 suggests transmission is driven by a relatively small number of high-risk contacts.</p> <p>-&gt; A large household SAR further suggests that between-household transmission risk is lower; otherwise the observed R0 would be larger.</p> <p><b>More data are needed.</b></p>
The Lancet 27FEB2020	<b>COVID-19: combining antiviral and anti-inflammatory treatments</b>  <b>COMMENT</b>	Stebbing et al., UK <a href="https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30132-8/fulltext">https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30132-8/fulltext</a>	Therapeutic	<p>-&gt; COVID-19 characterised by an overexuberant inflammatory response</p> <p>SARS -&gt; viral load is not correlated with the worsening of symptoms</p> <p>-&gt; Inhibition of <b>numb-associated kinase (NAK)</b> family would reduce viral infection in vitro (inhibit clathrin-mediated endocytosis and thereby inhibit viral infection of cells)</p> <p>-&gt; <b>JAK-STAT</b> signalling inhibitors, could be effective against the consequences of the elevated levels of cytokines (including interferon) typically observed in people with COVID-19</p> <p>-&gt; Baricitinib is a NAK inhibitor (<b>anti-viral</b>)</p> <p>-&gt; Baricitinib, fedratinib, and ruxolitinib are JAK inhibitors (<b>anti-inflammatory</b>)</p> <p><b>-&gt; Baricitinib is the best of the group</b></p>
The Lancet 27FEB2020	<b>Positive RT-PCR Test Results in Patients Recovered From COVID-19</b>	Lan et al., China <a href="https://jamanetwork.com/journals/jama/fullarticle/2762452">https://jamanetwork.com/journals/jama/fullarticle/2762452</a>	Public Health/Epidemiology	<p><b>Little attention</b> has been paid to the <b>follow-up of recovered</b> patients so far.</p> <p><b>4 patients</b> with COVID-19 who met criteria for hospital discharge or discontinuation of quarantine in China (absence of clinical symptoms and radiological abnormalities and 2 negative RT-PCR test results) had <b>positive RT-PCR test results 5 to 13 days later</b>, while they were still <b>asymptomatic.</b></p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
The Lancet 27FEB2020	<b>Convalescent plasma as a potential therapy for COVID-19</b>  <b>COMMENT</b>	Chen et al., China <a href="https://www.thelancet.com/pdfs/journals/laninf/PIIS1473-3099(20)30141-9.pdf">https://www.thelancet.com/pdfs/journals/laninf/PIIS1473-3099(20)30141-9.pdf</a>	Therapeutic	-> In 2014, the use of convalescent plasma collected from patients who had recovered from <b>Ebola virus disease</b> was <b>recommended by WHO</b> as an empirical treatment during outbreaks. -> A protocol for the use of convalescent plasma in the treatment of MERS coronavirus was established in 2015. -> <b>H1N1</b> : significant reduction of relative risk of mortality / <b>no adverse event</b> . -> and other studies <b>Antibodies from convalescent plasma might suppress viraemia</b>
Emerg Microb Infects 26FEB2020	<b>Detectable 2019-nCoV viral RNA in blood is a strong indicator for the further clinical severity</b>	Chen et al., China <a href="https://www.tandfonline.com/doi/full/10.1080/22221751.2020.1732837">https://www.tandfonline.com/doi/full/10.1080/22221751.2020.1732837</a>	Clinic	-> All patients (n=6 / 57) with <b>detectable viral RNA in the blood</b> progressed to severe symptom stage, indicating a strong <b>correlation of serum viral RNA with the disease severity</b> (p-value = 0.0001). -> 8 of the 11 patients with <b>annal swab virus-positive</b> was in <b>severe clinical stage</b> . -> Concentration of viral RNA in the <b>anal swab was higher than in the blood: virus might replicate in the digestive tract</b>
The Lancet, 26FEB2020	<b>The psychological impact of quarantine and how to reduce it: rapid review of the evidence</b>	Brooks et al., UK <a href="https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30460-8/fulltext">https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30460-8/fulltext</a>	HSS/Politic	-> Information is key; people who are quarantined need to understand the situation -> The quarantine period should be short and the duration <b>should not be changed</b> unless in extreme circumstances -> Most of the adverse effects come from the imposition of a restriction of liberty; voluntary quarantine is associated with less distress and fewer long-term complications -> <b>Public health officials should emphasise the altruistic choice of self-isolating</b>
Viruses 25FEB2020	<b>Preliminary Identification of Potential Vaccine Targets for the COVID-19 Coronavirus (SARS-CoV-2) Based on SARS-CoV Immunological Studies</b>	Ahmed et al., China <a href="https://www.mdpi.com/1999-4915/12/3/254">https://www.mdpi.com/1999-4915/12/3/254</a>	Vaccine	-> <b>High genetic similarity</b> between <b>SARS-CoV-2 and SARS-Co</b> . -> Identification of a set of <b>B cell and T cell epitopes</b> derived from the spike (S) and nucleocapsid (N) proteins that <b>map identically</b> to SARS-CoV-2 proteins. -> <b>No mutation</b> has been observed in these epitopes (as of 21 February 2020). -> <b>Immune targeting of these epitopes</b> may offer protection against this novel virus
EuroSurv 25FEV2020	<b>Differential diagnosis of illness in patients under investigation for the novel coronavirus (SARS-CoV-2), Italy, February 2020.</b>	Bordi et al., Italy <a href="https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.8.2000170">https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.8.2000170</a>	Public Health/Epidemio	-> <b>Similarity of symptoms</b> shared with more common respiratory infections. -> <b>Broad screening</b> requested. -> <b>Influenza virus</b> infections: <b>28.5% of all suspected cases</b> of SARS-CoV-2 infection. -> <b>Alternative diagnoses may clarify an individual patient's risk and may allow adjusting public health containment measures.</b>
The Lancet 25FEB2020	<b>Potential association between COVID-19 mortality and health-care resource availability</b>	Ji et al., China <a href="https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30068-1/fulltext">https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30068-1/fulltext</a>	Public Health/Epidemio	Plotting mortality against the incidence of COVID-19 (cumulative number of confirmed cases since the start of the outbreak, per 10 000 population) showed a significant positive correlation, suggesting that <b>mortality is correlated with health-care burden</b>
The Lancet 24FEB2020	<b>COVID-19 control in China during mass population movements at New Year</b>	Chen et al., China <a href="https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30421-9/fulltext">https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30421-9/fulltext</a>	Public Health/Epidemio	Several lessons that can be drawn from China's extension of the Lunar New Year holiday: 1-> Countries should consider <b>periods of recommended or mandatory closure of non-essential workplaces and public institutions</b> — to slow the rate of transmission. 2-> To tailor the design of these actions according to specific epidemic characteristics (incubation period and transmission routes). 3-> This is to prevent people with asymptomatic infections from spreading the disease.  As such, <b>governments should use the closure period for information and education campaigns, community screening, active contact tracing, and isolation and quarantine</b> to maximise impact.

Journal and date	Title	Authors and link	Field of expertise	Key facts
J Clin Med 24FEB2020	<b>Assessing the Impact of Reduced Travel on Exportation Dynamics of Novel Coronavirus Infection (COVID-19)</b>	Anzai et al., Japan <a href="https://www.mdpi.com/2077-0383/9/2/601">https://www.mdpi.com/2077-0383/9/2/601</a>	Public Health/Epidemiology	<p>-&gt; From <b>28 January to 7 February 2020</b>, around 226 exported cases were prevented (=70.4% reduction in incidence)</p> <p>-&gt; Reduced probability of a major epidemic in Japan: from 7% to 20% (=median time delay: of 2 days)</p> <p>-&gt; Depending on the scenario, the estimated delay may be less than one day. As the <b>delay is small</b>, the decision to control travel volume through restrictions on freedom of movement should be <b>balanced</b> between the <b>resulting estimated epidemiological impact</b> and <b>predicted economic fallout</b>.</p>
Cell Discov 24FEB2020	<b>Comparative genetic analysis of the novel coronavirus (2019-nCoV/SARS-CoV-2) receptor ACE2 in different populations</b>	Cao et al., China <a href="https://jcm.asm.org/content/early/2020/02/28/jcm.00155-20.long">https://jcm.asm.org/content/early/2020/02/28/jcm.00155-20.long</a>	Fundamental Research	<p>-&gt; Previous studies demonstrated the <b>positive correlation of ACE2 expression and the infection of SARS-CoV in vitro</b></p> <p>-&gt; <b>Here:</b> Systematic analysis of coding-region variants in ACE2 and the eQTL variants (may affect the expression of ACE2) among different populations (GTEx database)/</p> <p>-&gt; The <b>East Asian</b> populations have <b>much higher AFs</b> in the <b>eQTL variants</b> associated with <b>higher ACE2 expression</b> in tissues which may suggest <b>different susceptibility or response</b> to 2019-nCoV/SARS-CoV-2 <b>from different populations</b> under the similar conditions.</p> <p>-&gt; No direct evidence supporting the existence of <b>coronavirus S-protein binding-resistant ACE2 mutants</b> in different populations.</p>
The Lancet 24FEB2020	<b>Clinical course and outcomes of critically ill patients with SARS-CoV-2 pneumonia in Wuhan, China: a single-centered, retrospective, observational study</b>	Xiaobo Yang et al., China <a href="https://www.thelancet.com/journals/lanres/article/PIIS2213-2600(20)30079-5/fulltext">https://www.thelancet.com/journals/lanres/article/PIIS2213-2600(20)30079-5/fulltext</a>	Clinic	<p>- <b>Mortality is high</b>. The survival term of the non-survivors is likely to be within 1–2 weeks after ICU admission.</p> <p>- Older patients (&gt;65 years) with comorbidities and ARDS are at increased risk of death.</p>
The Lancet 24FEB2020	<b>Viral load of SARS-CoV-2 in clinical samples</b>	Pan et al., China <a href="https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30113-4/fulltext">https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30113-4/fulltext</a>	Virology	<p>- The <b>viral loads</b> in throat swab and sputum samples peaked at around <b>5–6 days after symptom onset</b>, ranging from around 104 to 107 copies per mL during this time</p> <p>- <b>Sputum samples</b> generally showed higher viral loads than throat swab samples.</p>
The Lancet 24FEB2020	<b>COVID-19 pneumonia: what has CT taught us?</b>	Lee et al., China <a href="https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30134-1/fulltext">https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30134-1/fulltext</a>	Diagnostic	<p>- The predominant CT findings included <b>ground-glass opacification, consolidation, bilateral involvement, and peripheral and diffuse distribution</b>.</p> <p>- More research is needed to correlate of CT findings with clinical severity and progression, the predictive value of baseline CT or temporal changes for disease outcome, and the sequelae of acute lung injury induced by COVID-19.</p>
J Med Virol 21FEB2020	<b>COVID-2019: the role of the nsp2 and nsp3 in its pathogenesis.</b>	Angeletti et al., Rome, Italy <a href="https://onlinelibrary.wiley.com/doi/abs/10.1002/jmv.25719">https://onlinelibrary.wiley.com/doi/abs/10.1002/jmv.25719</a>	Fundamental Research	<p>The Open Reading Frame 1ab (ORF1ab) of COVID-2019 has been analyzed to evidence the <b>presence of mutation caused by selective pressure</b> on the virus.</p> <p>Which are the probably most common sites undergoing to an aminoacidic change ?</p> <p>-&gt; Insight of some important proteins of the COVID-2019 that are involved in the mechanism of viral entry and viral replication</p> <p>Results: Both nsp2 and nsp3 are under selective pressure. <b>nsp2</b>-&gt; could explain why this virus is more contagious than SARS <b>nsp 3</b>-&gt; could suggest a potential mechanism differentiating COVID-2019 from SARS</p>



Journal and date	Title	Authors and link	Field of expertise	Key facts
Radiology 20 FEB 2020	<b>Chest CT Findings in Coronavirus Disease-19 (COVID-19): Relationship to Duration of Infection.</b>	Bernheim et al., <a href="https://pubs.rsna.org/doi/10.1148/radiol.2020200463">https://pubs.rsna.org/doi/10.1148/radiol.2020200463</a>	Diagnostic	Frequency of CT findings is related to infection time course.
The Lancet, 20 FEB 2020	<b>Preparedness and vulnerability of African countries against importations of COVID-19: a modelling study.</b>	Gilbert et al., Vittoria's team <a href="https://www.thelancet.com/journals/laninf/article/PIIS0140-6736(20)30411-6/fulltext">https://www.thelancet.com/journals/laninf/article/PIIS0140-6736(20)30411-6/fulltext</a>	Public Health/Epidemiology	<ul style="list-style-type: none"> <li>- Highest importation risk: Egypt, Algeria, and South Africa -&gt; moderate to high capacity to respond to outbreaks</li> <li>- Moderate risk: Nigeria, Ethiopia, Sudan, Angola, Tanzania, Ghana, and Kenya -&gt; variable capacity and high vulnerability</li> </ul>
The Lancet 19FEB2020	<b>Asymptomatic cases in a family cluster with SARS-CoV-2 infection</b>	Pan et al., China <a href="https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30114-6/fulltext">https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(20)30114-6/fulltext</a>	Public Health/Epidemiology	<ul style="list-style-type: none"> <li>- In this family cluster, although <b>all individuals tested positive</b> for SARS-CoV-2 infection on qRT-PCR, only patient 1 showed clinical symptoms, decreased lymphocyte count, and abnormal chest CT images.</li> <li>- However, <b>any of the three individuals could have been the first one to become infected</b> and thus transmitted the virus to the other two family members.</li> </ul>
The Lancet 19FEB2020	<b>Enteric involvement of coronaviruses: is faecal–oral transmission of SARS-CoV-2 possible?</b>	Yeo et al., Singapore <a href="https://www.thelancet.com/journals/langas/article/PIIS2468-1253(20)30048-0/fulltext">https://www.thelancet.com/journals/langas/article/PIIS2468-1253(20)30048-0/fulltext</a>	Virology	<ul style="list-style-type: none"> <li>- Considering the evidence of faecal excretion for both SARS-CoV and MERS-CoV, and their ability to remain viable in conditions that could facilitate faecal–oral transmission, it is possible that SARS-CoV-2 could also be transmitted via this route.</li> <li>-&gt;When <b>SARS-CoV</b> was seeded into sewage water obtained from the hospitals in a separate experiment, the virus was found to remain infectious for 14 days at 4°C, but for only 2 days at 20°C. SARS-CoV can survive for up to 2 weeks after drying, remaining viable for up to 5 days at temperatures of 22–25°C and 40–50% relative humidity, with a gradual decline in virus infectivity thereafter. Viability of the SARS-CoV virus decreased after 24 h at 38°C and 80–90% relative humidity.</li> <li>-&gt; <b>MERS-CoV</b> is viable in low temperature, low humidity conditions. The virus was viable on different surfaces for 48 h at 20°C and 40% relative humidity, although viability decreased to 8 h at 30°C and 80% relative humidity conditions.</li> </ul>
THE NEJM, 19FEB2020	<b>SARS-CoV-2 Viral Load in Upper Respiratory Specimens of Infected Patients</b>	Zou et al., China <a href="https://www.nejm.org/doi/full/10.1056/NEJMc2001737">https://www.nejm.org/doi/full/10.1056/NEJMc2001737</a>	Virology	<ul style="list-style-type: none"> <li>- <b>The higher viral loads were detected soon after symptom onset.</b></li> <li>- <b>Higher viral loads detected in the nose</b> than in the throat.</li> <li>- Our analysis <b>suggests</b> that the viral nucleic acid shedding pattern of patients infected with SARS-CoV-2 resembles that of patients with influenza and appears different from that seen in patients infected with SARS-CoV.</li> <li>- The viral load that was detected in the <b>asymptomatic patient</b> was <b>similar</b> to that in the <b>symptomatic patients</b>, which <b>suggests the transmission potential of asymptomatic or minimally symptomatic patients.</b></li> </ul>
Biosci Trends, 19FEB2020	<b>Breakthrough: Chloroquine phosphate has shown apparent efficacy in treatment of COVID-19 associated pneumonia in clinical studies.</b>	Gao et al., <a href="https://www.ncbi.nlm.nih.gov/pubmed/32074550">https://www.ncbi.nlm.nih.gov/pubmed/32074550</a>	Therapeutic	<p><b>Chloroquine phosphate</b>, an old drug for treatment of malaria, is shown to have apparent efficacy and acceptable safety against COVID-19 associated pneumonia in multicenter clinical trials conducted in China. <b>(DATA NOT SHOWN !)</b>.</p> <p>The drug is recommended to be included in the next version of the Guidelines for the Prevention, Diagnosis, and Treatment of Pneumonia Caused by COVID-19 issued by the National Health Commission of the People's Republic of China for treatment of COVID-19 infection in larger populations in the future.</p>

Journal and date	Title	Authors and link	Field of expertise	Key facts
J Infect Dis. 18FEB2020	<b>A familial cluster of infection associated with the 2019 novel coronavirus indicating potential person-to-person transmission during the incubation period.</b>	Yu et al., China <a href="https://academic.oup.com/jid/advance-article/doi/10.1093/infdis/jiaa077/5739751">https://academic.oup.com/jid/advance-article/doi/10.1093/infdis/jiaa077/5739751</a>	Public Health/Epidemiology	Familial cluster of four patients in Shanghai. One was 88 years old man with moving difficulties and was only exposed to his asymptomatic family members who <b>developed symptoms later</b> .  The epidemiological evidence has shown a potential transmission of the 2019-nCoV during the incubation period.
The Lancet 18FEB2020	<b>Tracking online heroisation and blame in epidemics</b>  <b>COMMENT</b>	Atlani Duault et al., France <a href="https://www.thelancet.com/action/showPdf?pii=S2468-2667%2820%2930033-5">https://www.thelancet.com/action/showPdf?pii=S2468-2667%2820%2930033-5</a>	HSS/Politics	-> Gathering online <b>data on local perceptions</b> has the potential to help public authorities mount more robust responses and better targeted health communications -> It is important to track the evolving dynamics of blame in <b>real time</b> , both <b>to correct inaccurate</b> information and <b>to respond to online scapegoating</b> . -> <b>Trust is a crucial</b> support to public health systems. <b>Public health authorities need to be aware of « complex geographies of hope and blame » while planning responses to the epidemic.</b>
Biochem Biophys Res Comm 17 FEB 2020	<b>Structure analysis of the receptor binding of 2019-nCoV</b>	Chen et al., China and USA <a href="https://www.sciencedirect.com/science/article/pii/S006291X20303399">https://www.sciencedirect.com/science/article/pii/S006291X20303399</a>	Fundamental Research	Structural analysis of the receptor binding domain (RBD) -> 72% identity with SARS CoV / Higher affinity with ACE 2.  ACE2 is widely expressed with conserved primary structures <u>throughout</u> the animal kingdom ( <b>possible hosts ?</b> ) Since ACE2 is predominantly expressed in intestines, testis, and kidney, <b>fecal-oral</b> and <b>other routes</b> of transmission are also <b>possible</b> .  Finally, antibodies and small molecular inhibitors that can block the interaction of ACE2 with RBD should be developed to combat the virus.
J Clin Med 17 FEB 2020	<b>Incubation Period and Other Epidemiological Characteristics of 2019 Novel Coronavirus Infections with Right Truncation: A Statistical Analysis of Publicly Available Case Data.</b>	Linton et al., Japan <a href="https://www.mdpi.com/2077-0383/9/2/538">https://www.mdpi.com/2077-0383/9/2/538</a>	Public Health/Epidemiology	Incubation period falls within the range of 2–14 days with 95% confidence and has a mean of around 5 days.  The mean time from illness onset to hospital admission (for treatment and/or isolation) was estimated at 3–4 days without truncation and at 5–9 days.
PNAS, 13FEB2020	<b>Prophylactic and therapeutic remdesivir (GS-5734) treatment in the rhesus macaque model of MERS-CoV infection</b>	De Wit et al., USA <a href="https://www.pnas.org/content/early/2020/02/12/1922083117">https://www.pnas.org/content/early/2020/02/12/1922083117</a>	Therapeutic	- 24 h prior to inoculation -> <b>completely prevented MERS-CoV-induced clinical disease</b> , strongly inhibited MERS-CoV replication in respiratory tissues, and prevented the formation of lung lesions. - 12 h postinoculation -> <b>clear clinical benefit</b> , with a reduction in clinical signs, reduced virus replication in the lungs, and decreased presence and severity of lung lesions.  <b>- Remdesivir may be considered for SARS-CoV -2</b>
The Lancet 12 FEB 2020	<b>What are the risks of COVID-19 infection in pregnant women?</b>	Qiao et al., China <a href="https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30365-2/fulltext">https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30365-2/fulltext</a>	Clinic	The clinical characteristics reported in pregnant women with confirmed COVID-19 infection are <b>similar to those reported for non-pregnant</b> adults with confirmed COVID-19 infection in the general population and are indicative of a relatively optimistic clinical course and outcomes for COVID-19 infection compared with SARS-CoV-1 infection.
The Lancet 12FEB2020	<b>Clinical characteristics and intrauterine vertical transmission potential of COVID-19 infection in nine pregnant women: a retrospective review of medical records</b>	Chen et al., China <a href="https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30360-3/fulltext">https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)30360-3/fulltext</a>	Clinic	Evidence of intrauterine vertical transmission was assessed by testing for the presence of SARS-CoV-2 in amniotic fluid, cord blood, and neonatal throat swab samples. <b>All samples tested negative</b> <b>None of the 9 patients developed severe COVID-19 pneumonia or died.</b>
Cell Res 4FEB2020	<b>Remdesivir and chloroquine effectively inhibit the recently emerged novel coronavirus (2019-nCoV) in vitro</b>	Wang et al., China <a href="https://www.nature.com/articles/s41422-020-0282-0">https://www.nature.com/articles/s41422-020-0282-0</a>	Therapeutic	Remdesivir and chloroquine <b>are highly effective</b> in the control of 2019-nCoV infection <b>in vitro</b> . These compounds have been used in human patients with a safety track record and shown to be effective against various ailments. They should be assessed in human patients suffering from the <b>novel coronavirus disease</b> .

Journal and date	Title	Authors and link	Field of expertise	Key facts
Euro Surveill 6FEB2020	<b>Effectiveness of airport screening at detecting travellers infected with novel coronavirus (2019-nCoV).</b>	Quilty et al., UK <a href="https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.5.2000080">https://www.eurosurveillance.org/content/10.2807/1560-7917.ES.2020.25.5.2000080</a>	Public Health/Epidemiology	<p>Estimation: <b>46%</b> of infected travellers would <b>not be detected</b>, depending on incubation period, sensitivity of exit and entry screening, and proportion of asymptomatic cases.</p> <p>-&gt; Airport screening is unlikely to detect a sufficient proportion of 2019-nCoV infected travellers to avoid entry of infected travellers.</p>
The Lancet 03FEB2020	<b>Baricitinib as potential treatment for 2019-nCoV acute respiratory disease</b>	Richardson et al., UK <a href="https://www.thelancet.com/pdfs/journals/lancet/Pii/S0140-6736(20)30304-4.pdf">https://www.thelancet.com/pdfs/journals/lancet/Pii/S0140-6736(20)30304-4.pdf</a>	Therapeutic	<p>The receptor that 2019-nCoV uses to infect lung cells might be ACE2, a cell-surface protein on cells in the kidney, blood vessels, heart, and, importantly, lung AT2 alveolar epithelial cells. One of the known regulators of endocytosis is the AP2-associated protein kinase 1 (AAK1). The plasma concentration of Baricitinib on therapeutic dosing (either as 2 mg or 4 mg once daily) is sufficient to inhibit AAK1, we suggest it could be trialled.</p>
Emerging Microbes Infect 03FEB2020	<b>Potent binding of 2019 novel coronavirus spike protein by a SARS coronavirus-specific human monoclonal antibody.</b>	Tian et al., China <a href="https://www.biorxiv.org/content/10.1101/2020.01.28.923011v1">https://www.biorxiv.org/content/10.1101/2020.01.28.923011v1</a>	Fundamental Research	<p>A SARS-CoV-specific human monoclonal antibody, CR3022, could bind potently with 2019-nCoV RBD.</p> <p>-&gt; Potential to be developed as candidate therapeutics ?</p> <p>Some of the most potent SARS-CoV-specific neutralizing antibodies that target the ACE2 binding site of SARS-CoV failed to bind 2019-nCoV spike protein. -&gt; It is still <b>necessary to develop novel monoclonal antibodies</b> that could bind specifically to 2019-nCoV RBD.</p>