**Recent Efforts and Advances for the Management of COVID-19**

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**Abstract**

A novel type beta strain of coronavirus, the severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2), caused the latest COVID-19 pandemic disease that emerged in Wuhan City, Hubei Province, China, in early December 2019. It is an enveloped positive-stranded RNA virus recognized for exhibiting interactions predominantly with lung epithelial cells. The WHO recommended avoiding gathering in public places, maintaining physical distance, using alcohol-based sanitizer, and washing hands thoroughly with soap as precautionary steps. No standard treatment and medicine are currently available to fight against COVID-19, challenging research communities and pharmaceutical industries to revolutionize drug discovery and vaccine development. Meanwhile, efforts to boost immunity power appear to be the need of the hour to combat COVID-19 illness. This review has been structured to provide detailed information on the causative agents of COVID-19, possible diagnostic and therapeutic approaches, and potential drugs and their impact on health. In addition, we briefly highlight stem cell-based therapy as another approach to disease management and control. Immune responses elicited in the human body against SARS-CoV-2 are also discussed. The recent emergence of the SARS-CoV-2 strain called OMICRON has sparked worldwide concern. This review also covers the identification and global spread of OMICRON, which has expanded to 77 countries, resulting in numerous speculations concerning its origin and degree of infectivity. Identifying mutations in the spike protein's RBD (receptor binding domain) region is a cause for concern because it goes beyond vaccination immunity. The following will discuss its transmission potential, infectivity, and impact of COVID-19 vaccinations.

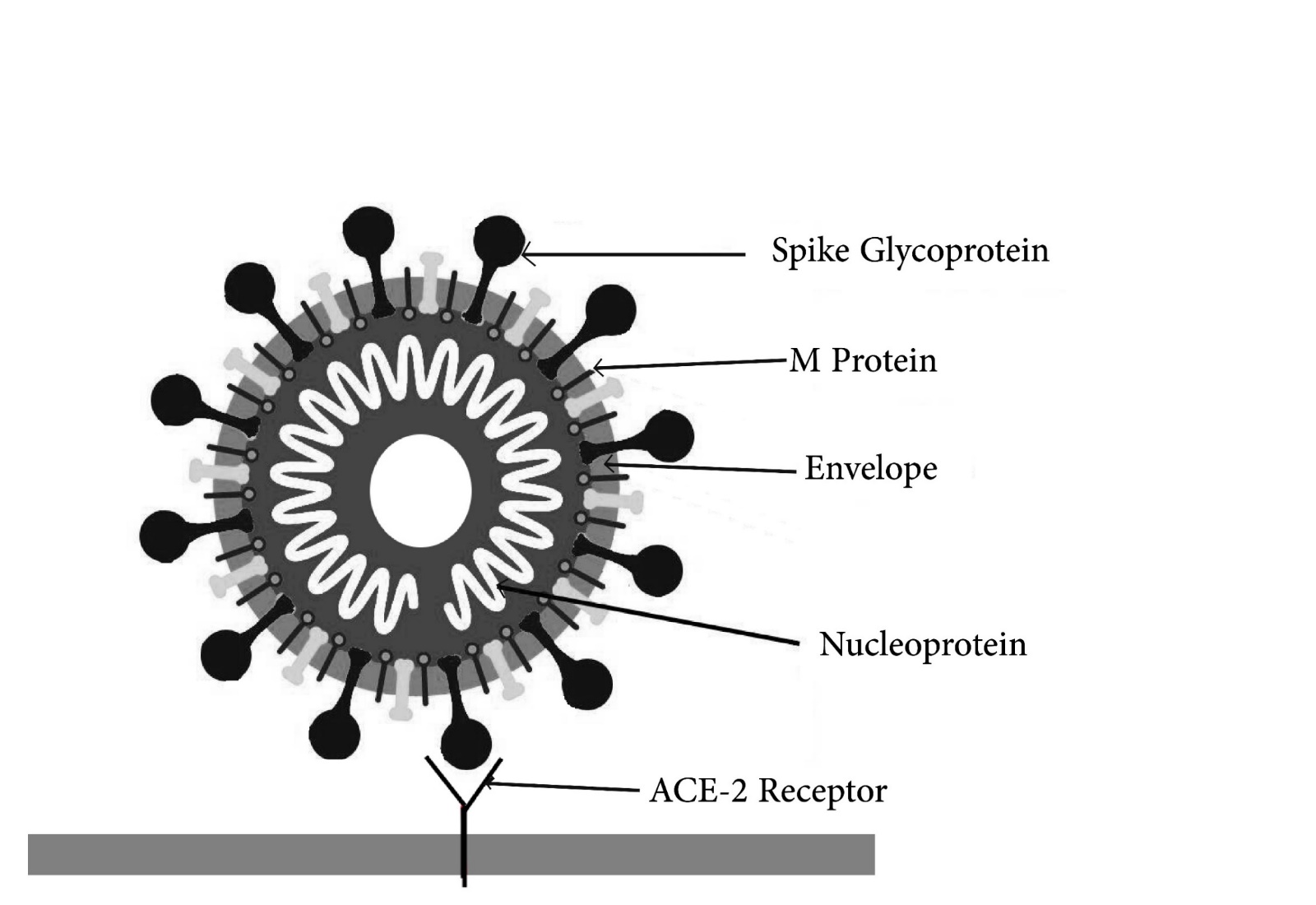
### **Keywords**

COVID-19, SARS-CoV-2, MERS, Drug Repurposing, Stem Cells, Immune Response.

1. **Introduction**

Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) is a newly evolving infectious disease and an epidemic reported worldwide caused by this novel coronavirus. Coronaviruses belong to the family of enveloped RNA viruses that can cause disease in humans and animals. Numerous viruses in this family are Middle East respiratory syndrome (MERS) and severe acute respiratory syndrome (SARS), which lead to respiratory infections in humans with symptoms such as common colds and more fatal infections 1. Patients were found to have pneumonia-like symptoms of unknown cause in December 2019, and most of them were found to have some connection to the wholesale seafood market in Wuhan. Subsequently, SARS-CoV-2, the Ortho Coronaviridae subfamily of the Coronaviridae family, was discovered as the primary leading agent for this new form of disease 2. SARS-CoV-2 is now declared a pandemic impacting several countries around the world 1. As the virus poses a significant risk to human health and the economy, there is a persevered worldwide for the immediate production of effective therapies and prophylactics for its control and prevention. As of December 14, 2020, a total of 72,893,236+ confirmed COVID-19 cases, including 1,623,425+ deaths, 51,121,100+ recovered, and 20,148,711+ active cases were reported worldwide with the infection fatality rate of 3 percent 3. The case-fatality rate (CFR) of SARS-CoV-2 has been estimated by the World Health Organization (WHO) to be between 0.3 and 1 percent, comparatively influenza A with CFR 0.1 percent lower than that of COVID-19 4. Some observational studies conducted in countries adopting COVID-19 preventive strategies have suggested that 80% of COVID-19 patients have had symptomless or moderate disease 4–6, 14% of patients had serious illnesses and 6% were in severe situations 4,6.

The outer surface of the “corona’’ appears to have crown-like spikes, thus referred to as coronavirus. Coronavirus has a small size which is ranging from 65-125 nm in diameter, and the nucleic material is single-stranded positive-sense RNA with a length of 26 to 32 kb. Coronavirus family is classified into four subgroups, i) Alpha Coronavirus: Human coronavirus HCoV-229E and HCoV-NL63, ii) Beta Coronavirus: this includes HCoV-OC43 and SARS-CoV, HCoV-HKU1 and MERS-CoV iii) Gamma Coronavirus: representing viruses of birds and whales and iv) Delta Coronavirus: a category of viruses isolated from birds and pigs 7–9. Viruses such as SARS coronavirus, avian influenza, swine flu 2009, and MERS coronavirus have similar symptoms like lung infection and breathing difficulties that ultimately lead to lung damage and deaths. The genetic material of all coronaviruses is generally organized in a manner wherein the 5′ end encodes the enzyme locus replicase, and the structural proteins are encoded by ORFs 10, 11 on 1/3rd of the genome within the 3' terminus. Coronavirus genes have four structural proteins structured in the 5'-3' sequence as spike (S), envelope (E), membrane (M) & nucleocapsid (N) (Figure 1) 10. Other coronaviruses, such as some beta coronaviruses, have specific structural and additional proteins, such as hemagglutinin esterase (HE) 10. Enclosed nucleic acid capsid protein connected with the RNA genetic material forming an icosahedral shape within the outer membrane. The envelope spike protein forms the peplomers and provides the crown-like structure to coronaviruses 11. 5' end of the coronavirus genome encodes the replicase enzyme, which comprises two open reading frames. Two large replicase polyproteins are translated by the replicase enzymes that are cleaved co-translationally into 16 proteins, which include 2 to 3 proteases, several modification enzymes, polymerases, and helicases 11.

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**Figure 1.** Schematic representation of coronavirus structure and viral receptor, i.e., angiotensin-converting enzyme 2 (ACE2) on the host cell surface.

SARS-CoV-2, with the help of its class I fusion spike (S) glycoprotein, recognizes the angiotensin I converting enzyme 2 (ACE2) as a binding site on the host cell, particularly epithelial cells and consecutively fuses with the host cell. The entry of the virus into target cells and further its transmission is allowed by the serine protease transmembrane protease, serine 2 (TMPRSS2) 4, which helps in priming of the virus spike protein and involves the cleavage of the viral spike protein at the S1/S2 site. The S1 subunit facilitates the binding of the spike protein via its receptor-binding domain. Upon attaching to the receptor, conformational changes in the spike protein occur 4,12. The S2 subunit helps the membrane to integrate and internalize the virus. ACE2 and TMPRSS2 receptors are usually found in alveolar type II cells and capillary endothelial cells in the lungs 4,13. As a result, SARS-CoV-2 infection demonstrates extreme respiratory disease as among the main symptoms. Importantly, infection with SARS-CoV-2 triggers the overproduction of cytokines and increases the pro-inflammatory response in the lungs. Excess levels of cytokines cause oedema, difficulty in breathing, acute respiratory failure and several other infections that can even lethe ad to the death of patients 4.

Delta and Omicron are two recent variations of concern. India was the first nation to identify the Delta variant in late 2020. The Delta variety was discovered on May 3, November 22, 2021, and spread to 179 countries. The World Health Organization (WHO) predicted in June 2021 that the Delta form would be among the most stressful worldwide. It appears to have feature mutations T478K, P,681R and, L452R in the gene encoding the SARS-CoV-2 spike protein, which has been linked to changes in the virus's infectiousness and how it can be neutralized with antigens against recently circulating COVID-19 virus subtypes. It is thought to be one of the most contagious severe illnesses ever fou. The variation is suspected of being involved in the devastating second wave of the epidemic, which began in India in February 2021. As a result, the third wave hit Fiji, the United Kingdom, and South Africa. By late July, it had also contributed to an increase in attacks in Asia, the United States, Australia, and New Zealand 19–25.

To date, more than 430 million people worldwide have been diagnosed with COVID-19, resulting in 5.9 million fatalities (According to WHO dashboards information March 1, 2022). Since its discovery, Omicron has caused an increase in infections in several countries and at several locations. According to WHO news, the number of verified COVID-19 cases in the United States surpassed one million in a single day in early January 2022. This rapid increase is consistent with the Omicron variant outbreak in the United States.

The Omicron variant has a constellation of over 50 mutations, roughly 30 of which are in the spike protein, which is the most concerning aspect of the variant. The 15 altered locations in the receptor-binding domain (RBD), which interacts with human cells before cell entry and may increase transmissibility 26, are the most concerning matter. Since the discovery of this variety in South Africa, scientists have been looking for evidence of its origin and the likely path of the unknown pandemic. Based on the examination of the Omicron and other SARS-CoV-2 variant sequences, researchers hypothesized that Omicron evolved in parallel and most likely diverged during early or possibly in mid-2020 from other strains 27. RBD (receptor-binding domain) analysis revealed the existence of two Omicron sub-clades. Furthermore, phylogenetic analysis of 3590 SARS-CoV-2 sequences revealed the development of Omicron from the B.1.1.519 family (clade 20B) 28. The presence of most mutations in the Omicron variant in the body of an immunocompromised HIV-infected individual shows the possibility of adaptations in COVID-19 patients who are persistently infected. A contributing cause to Omicron may be South Africa's high (19%) HIV prevalence rate 26,29.

#### **Origin of SARS-CoV-2 is linked with SARS-CoV and MERS-CoV**

The source of origin and transmission of the infection must be identified for the establishment of therapeutic strategies to control the disease. The previously confirmed SARS disease case was identified in Foshan, China in November 2002. Cases involving 300 health workers were reported in China in February 2003. The SARS outbreak in China later spread to other countries through infected individuals. The SARS epidemic 2003 occurred due to a new strain of human coronavirus as it was discovered. Some countries which came in contact with this virus were Hong Kong, Vietnam, Toronto, Taiwan, Canada, and the United States 30–32. Also, WHO issued a worldwide notice on March 12, 2003, warning of the spread of atypical pneumonia. After three days, WHO coined the name SARS for the spreading virus and provided an emergency alert along with caution and travel advisory. WHO observed air transport as one of the primary reasons for its spreading throughout the world. By April 04, 2003, the WHO had recorded 2353 cases. Approximately 4 percent of SARS patients died, as per the reports 33. In literature, it was revealed that the natural reservoir of SARS-CoV was horseshoe bats and civets and the animals sold in the China wet market that incredibly contributed to the SARS transmission from animals to humans 34. Talking or breathing, coughing, and sneezing served as common modes of transmission. SERS and MERS showed an incubation period of ~5 days and the illness was reported by more than 95% of patients approximately 13 days after exposure. Severe symptoms included fever, chills, coughing, and headache mild symptoms covered diarrhea, vomiting, and nausea in the first or second week of the illness. Patients who required intensive care infected with SARS-CoV were approximately 20-3,%; while in the case of MERS, 90-100% of ported required intensive care 30–32.

According to a report from Pro MED (Program for Monitoring Emerging Diseases) submitted on September 20, 2012, a new virus called MERS- was discovered from sputum taken from a man belonging to Arabian Peninsula, who died due to pneumonia and renal failure 3 months earlier 30,35,36. Acute res The Acute respiratory disease outbreak happened in 2012 in the Zarqa, Jordan, public hospital. The cause of the epidemic was an episode at the time, and laboratory studies were conducted shortly after the outbreak 36. The earliest case of severe respiratory illness, dated in April 2012 in Jordan Hospital, was subsequently diagnosed as MERS. Later, in September 2012, three MERS patients were found in the UK as well. Since 2012, MERS cases have been reported in 27 nations throughout the world, but a maximum of 80% of cases befell in Arabian Peninsula 30. MERS-CoV possesses a range of similarities with zoonotic viruses, as evidenced by the instances of its transmission from animals to humans. WHO reported that dromedary camels worked as a source of infection either directly or indirectly 30. By April 26, 2016, it was reported in 27 nations with 1,728 active MERS cases, 624 deaths, and 34.3% mortality rate 30,36. The primary symptoms of MERS included fever, cough, and breathlessness.

SARS-CoV-2 shows a great extent of similarities to SARS-CoV in terms of their mode of transmission and symptoms, and therefore, it is known to have emerged as a new modified pathogen that has caused COVID-19 disease. WHO reported the outbreak of unknown causes on January 05, 2020, and the organization called for an emergency health situation around the end of January 2020 37. On February 11, 2020 the organization announced COVID-19 as a pandemic 1. Like SARS and MERS, COVID-19 shows symptoms such as coughing, sneezing droplets, and breathing problems (Table 1). Also, the incubation time of SARS-CoV-2 is found to be 5-6 days after transmission 1. Although these three viruses show a great range of similarities, they exhibit unique features regarding the severity of its infection. Notably, SARS and MERS so far showed higher mortality rates in comparison to COVID-19 37. SARS-CoV-2 is, however, observed to be more infectious than MERS-CoV and SARS-CoV because SARS-CoV-2 infected asymptomatic individuals can transfer the virus potentially similar to symptomatic individuals. Initial analyses indicate that 79% genome of SARS-CoV-2 is similar to that of SARS-CoV but not identical at the nucleotide level. However, such changes are significantly noticed at the level of gene expression, as SARS-CoV-2 is found to possess only 72% similarity in its spike protein which is a crucial surface glycoprotein that interacts with receptors on host cells 38.

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| **Symptoms** | **Covid-19** | **Common Cold** | **Flu** | **Allergies** |
| **Fever** | Common | Rare | Common | Sometimes |
| **Dry Cough** | Common | Mild | Common | Sometimes |
| **Shortness Of Breath** | Common | No | No | Common |
| **Headache** | Sometimes | Rare | Common | Sometimes |
| **Aches And Pains** | Sometimes | Common | Common | No |
| **Sore Throat** | Sometimes | Common | Common | No |
| **Fatigue** | Sometimes | Sometimes | Common | Sometimes |
| **Diarrhea** | Rare | No | Sometimes | No |
| **Runny Nose** | Rare | Common | Sometimes | Common |
| **Sneezing** | No | Common | No | Common |

**Table 1:** Symptoms of COVID-19 in comparison with similar contagious diseases.

A study of Genome sequencing of SARS-CoV-2 and SARS-CoV revealed that the virus is similar to a bat coronavirus (BtCoV) RaTG13 originated from Yunnan Province in Rhinolophus with genome sequence similarity of 96.2% 39. In addition, the study reported that no recombination events occurred in the genome of SARS-CoV-2 from other bat’s originated viruses. Thus, literature so far published suggested that bats might be the natural host of SARS-CoV-2 39,40. However, more intensive research is needed to identify the possible intermediate host responsible for the transmission of this virus to humans 40. Studies reported so far reveal that SARS-CoV-2 causes excessive release of GCSF, IP10, MCPI, MIPIA, IL-2, IL-6, IL-7, and TNFα, which further lead to pulmonary edema, dysfunction of air-exchange, acute respiratory syndrome, acute cardiac injury and lastly even death. Thus, SARS-CoV-2 pathogenesis causes overproduction of immune cells and cytokine release in the lung 4.

In the middle of November 202141, the new OMICRON strain was discovered for the first time in the Gauteng area of South Africa. The advanced genome sequencing infrastructure in South Africa makes it easier to find novel variants early. According to WHO, the new SARS-CoV-2 strain, e.g., Omicron, has already been detected as of 15 December 2021 in 77 different nations, in most instances coming from the United Kingdom, South Africa, and the United States 42. India has also seen Omicron-positive cases. According to BBC News, the new COVID-19 variant's first death was recorded in the United Kingdom.

#### **Symptoms, Risk factor and Transmission Rate of COVID-19**

COVID-19 usually displays symptoms around 2 to 14 days after infection, informed by the Centers for Disease Control and Prevention (CDC) and WHO 43,44. According to WHO, patients diagnosed with COVID-19 show the most common symptoms such as dry cough, nausea, exhaustion, and headache; other common side effects include headaches, pharyngitis, sickness, eye infections, fever, ageusia, exanthem rash and discoloration of the skin (Table 1). The severe signs include breathing difficulty, chest discomfort, and loss of voice 44. The transmission rate of the virus is quite severe and depends a lot on the demography and quarantine level of the infected population 43. Infectivity is characterized as a pathogen's ability to develop an infection, denoted by the disease's reproductive number (R0, pronounced "R naught") 45. SARS-CoV-2 is predicted to have a R0 of 2 to 2.5 1; indicating that in a population, 2 to 2.5 individuals are susceptible to infection by each infected person.

COVID-19 spreads rapidly, with a primary reproductive number of 2.2 to 2.5 defined in Wuhan 46,47. However, this value estimation can vary on the basis of geography, population density, and size 48. Considering all the reports available in the literature, we have observed that the incubation period (detection time) in general for the SARS-CoV-2 is 5 days before any initial symptoms; however, it is likely that the same person can infect anyone after 2-3 days of the infection 49. Although an asymptomatic infection is recommended to be quarantined for at least 14 days, further studies are required to determine the infectivity duration in asymptomatic cases 50. The study recorded that during this phase, 44 percent of viral infections occurred 49,51. As per the CDC, a significant proportion of sick people with no symptoms are infectious 51,52.Based on research data, WHO suggested that kids, pregnant women, cancer patients, and people who are above 65 years of age are more prone in comparison to other age brackets and may unfold the sickness and transmission. Further added that, since they are at higher risk of getting exposed and showing symptoms, they should make sure to self-quarantine and avoid contact with others to prevent further transmission 1,52. Studies also indicate the susceptibility of an individual towards infection of this virus based on the blood group. In a study, individuals with A blood group are found to be more vulnerable to getting infected with SARS-CoV-2 53.

According to the World Health Organization (WHO), no research has shown that the Omicron version is more severe than other VOCs (Variants of Concern). Concerns about high transmissibility, virulence, increased risk of reinfection, and decreased efficacy of existing diagnostics, vaccines, and therapies remain unsolved. Chen et al. 54 used an artificial intelligence (AI) model (TopNetmAb) to predict the effect of 15 RBD mutations on the Omicron infectivity and efficacy of current vaccinations. The analysis found that mutations at the N440K, T478K, and N501Y locations may confer ten times and two times greater infectivity to Omicron, respectively, than the original SARS-CoV-2 and Delta variants. A study of 35,670 reinfections among 2.8 million positive cases in South Africa found significant population-level evidence for immune evasion from past infection. This suggests that the Omicron variation is involved in infections in people who have recovered 55. In support of this, the Omicron (pseudotyped) construct demonstrated an ED50 of 66 when evaluated against a panel of human sera collected from convalescent COVID-19 patients, representing an 8.4-fold reduction in neutralization 56.

The Omicron version shows a 13-fold increase in infectiousness, making it approximately 2.8 times more infectious than the Delta variant 54. The Omicron variant's primary reproductive number (R0), or the average number of additional cases created by a single infected individual, is similarly increased. The original SARSCoV2 strain has an R0 of 2-3, the Delta variation has an R0 of 5-8, and Omicron's R0 is believed to be as high as 10 57–59.

### **Diagnostic approaches for COVID-19**

COVID-19 diagnostic testing is critical to the monitoring of the virus, knowledge of its epidemiology, the reporting of case monitoring, and preventing the spread of the virus. For understanding drug repurposing, management of cases, and control transmission, it is essential to diagnose and trace the virus 60. On January 17, 2020, WHO released guidelines for laboratory testing of SARS-CoV-2 suspected individuals, acknowledging that the worldwide unfold of COVID-19 has surprisingly redoubled the quantity of suspected cases in the geographic regions wherever testing is increased and enforced properly. Rigorous tracing and testing of COVID-19 suspects have also led to a shortfall of testing reagents worldwide and alternative molecular medicines or repurposed drugs. In addition, there are constraints on treatment capability in several regions, particularly in low and medium-earning countries 61. Presently, there are multiple ways that are being utilized for testing COVID-19 suspects that are as follows-

#### **Real-time RT-PCR**

In laboratories that have validated a wide range of coronaviruses using real-time reverse transcription polymerase chain reaction (RT-PCR) assays, it is strongly recommended that the primers be checked against the published SARS-CoV-2 genomic sequence, as well as be tested if the primers overlap and have the capability to identify the presence of SARS-CoV-2. To determine the exact virus detected (e.g., on an amplicon of an unconserved region), sequencing should be performed on positive tests. On January 17, 2020, the WHO suggested visualizing the primers against the revealed SARS-CoV-2 genomic sequence and observing if primers complement and have the capability to find the SARS-CoV-2 61. It is a nucleic acid-derived method for the detection of the genetic material of infectious agents, including viruses, bacteria, etc. This equipment is one amongst the foremost commonly used for the diagnosis of the COVID-19 genome. In the past, for the identification of different diseases, such as those caused by the Ebola virus and some other viruses, several nations have used real-time RT–PCR assays. Therefore, several countries require this equipment as top-priority for identifying the presence of SARS-CoV-2 and also envision making strategies to expand further their national virus examination facilities 62.

Quantitative RT-PCR is a fluorescent-based technique that helps to examine small quantities of nucleic acid in a vast range of samples. It is a promising technology used in different fields, including life sciences, agriculture, biotechnology, and medical sciences, which gives better results at the molecular level. It is simple to operate in conjunction with speed, sensitivity, and specificity in a homogeneous assay 63. Coronavirus possesses a positive sense RNA strand and RNA-dependent RNA polymerase enzyme, which enables the virus to translate its own RNA copy by utilizing host machinery and replicate a large number of complementary DNA (cDNA) as an intermediate template that helps the synthesis of multiple sub-genomic mRNAs 64.

##### **Benefits of real-time RT-PCR**

Due to its sensitivity, specificity, high throughput, and reliable instrumentation, real-time RT-PCR is used frequently for the detection of COVID-19 virus. It takes approximately 3 h to obtain reliable results, although the laboratory takes between six and eight hours on average. Real-time RT-PCR is considerably quicker and encompasses fewer possible errors because the entire method runs within a single tube. It is proven as a reliable technique and the best available methodology for testing SARS-CoV-2 64,65.

##### **Drawbacks:**

Detection of past infections in real-time RT-PCR cannot be performed, which is necessary for the analysis of the virus’s emergence and transmission, as pathogens are known to be available in the body for a prolonged period of time. Alternate methods to diagnose, monitor and analyze previous infections, especially those that may have transmitted the virus without signs. RT-PCR is not a blood-based test, requires too much time to test, and involves collecting the sample from the throat and nasal swabs 64.

#### **Serological test**

Serological testing is very useful for confirming immunological response from a particular viral group, for example, coronaviruses. The best serological test results include the collection of paired serum samples (in acute and convalescent phases) from the recovered cases 61. CDC looks at antibody test data to determine the overall amount of persons tainted with the virus causing SARS-CoV-2 infection in the USA 66. CDC also uses antibody testing to understand more about how the immune system in the body reacts to the virus and how the virus spreads among people who are exposed to the virus 67. The information that CDC is investigating comes from several sources, including blood donors of people with symptoms and COVID-19 diagnoses.

**Types of serological assays**

##### **Rapid diagnostic test (RDT)**

RDT is generally a compact, movable, qualitative lateral flow test that could be used in healthcare. Such procedures require fingertip blood samples, drool samples, or nasal swab fluids.

This test is somewhat familiar to pregnancy kit tests because the test displays colored lines for users showing either positive or negative outcomes. Such tests most often recognize patients' antibodies (IgG and IgM) or viral antigens in the sense of COVID-19. For certain instances, it may be helpful in estimating the starting point (before infection) of IgG and IgM titers 68.

Antibodies are proteins that help to combat infections and typically provide protection against the recurrence of the disease (immunity). Antibody tests analyze the blood for antibodies that determine if you have had a prior infection with the COVID-19 virus 67. Antibodies are typically disease-specific; e.g., Measles antibodies can protect a person who is once again exposed to measles but has no effect if he is exposed to mumps 67. CDC is evaluating commercial antibody tests, as CDC works with other government agencies to assess the performance of commercially produced antibody tests 66.

##### **ELISA (Enzyme-Linked Immunosorbent Assay)**

ELISA is used as a qualitative or quantitative test depending on the test types required considering sensitivity and specificity. It is also known as rapid fast tests, which are only used for surveillance purposes. In this technique, whole blood, plasma, or serum of infected patients are required as samples. The test depends on a virus protein like spike protein or inactivated form of the virus coated on a multi-well plate. The sample is incubated in multi-well plates, and eventually, an antibody-protein complex is supposed to be formed if antibodies are already present in the sample against the viral protein coated on the substrate. For the estimation, a fluorescent-labeled secondary antibody is used which produces a color change when bound with an antibody-protein complex, thus detecting antibodies against the virus. This technique is utilized for detection of antibodies that are developed in response to SAR-CoV-2 infection 68. In particular, an immunoglobulin G (IgG) based ELISA test is preferred and developed by various research groups across the world, which has the advantage of analyzing a large number of samples in one go within 2-3 h. This test, in general, is characterized by 100% specificity and 98.7% sensitivity, suggesting that it can give reliable data that will indirectly confirm the exposure of a person to the coronavirus 69.

##### **Neutralization assay**

Neutralizing assay is a cell culture-based method that requires the growth of SARS-CoV-2 culturing cells (e.g., Vero E6 cells). This test is primarily based on observing the potency of antibodies of the patient to stop viral infection of cells in a suitable culture environment. If the patient has active and effective antibodies against coronavirus, this assay can recognize the viruses easily. In this test, samples such as whole blood, plasma, or serum are required from the infected patient. Neutralizing assay generally visualizes and quantifies the extent of blocking of virus replication through antibodies present in the patient’s serum. This blocking phenomenon reveals that antibodies specifically bind to a surface moiety of the virus meant for facilitating cell entry and thus help neutralize them 68.

##### **Chemiluminescent immunoassay (CLIA)**

This immunoassay is a quantitative biochemical technique requiring whole blood plasma or serum as patients’ samples. This test is also known as a chemiluminescent microparticle immunoassay due to magnetic protein-coated microparticles. In this type of test, several reagents such as buffer solution, antigen-antibody complex formed by specific primary antibodies, and enzyme-linked secondary antibodies were added sequentially to the patient sample. Their binding generates a light-emitting chemical reaction. The amount of light (radiance) emitted from each sample is then utilized to evaluate the number of antibodies found in the patient sample. This test can look for antibodies, namely IgM, IgG, and IgA. However, the drawback of this type of testing is that it cannot predict whether the antibodies detected can block the growth of the viruses 68,70.

#### **TrueNat/CBNAAT-based testing for COVID-19**

Following initial validation of TrueNat test machines for screening of SARS-CoV-2 infected patients, the Indian Council of Medical Research (ICMR) released revised guidelines on May 19, 2020, calling the test a 'comprehensive method for the screening and confirmation of COVID-19 cases 71. The TrueNat system, a cartridge-based nucleic acid amplification test (CBNAAT) indigenously manufactured in India, was originally designed for tuberculosis diagnosis 71,72.

**Working principle of TrueNat tests**

TrueNat is the Rt-PCR series, a battery-powered and chip-based machine that uses a two-step SARS-CoV-2 detection method. The first stage is detecting the presence of E-gene in the COVID-19 virus. E-gene allows the virus to create a spherical envelope around it. Step two recognizes the RNA-dependent RNA polymerase (RdRp) gene in the viral RNA and confirms the presence of the virus. For confirmation of the existence of E-gene in the sample, RT-PCR studies can be performed. However, new systems are designed to search for the RNA-dependent RNA polymerase (RdRp) enzyme in the RNA virus. Consequently, a two-step test can confirm all the samples tested positive for E-gene in the same laboratory. The ICMR has determined that such test results could be seen as evidence of a new coronavirus 73.

The sample of the throat and nasal swab is extracted and placed in a viral transmission medium where it is neutralized. It is then transported to another solvent, the viral lysis buffer, where the cells are broken down, and the contaminants are excluded. A portion of a solvent is then transported to a cartridge that appears like a disc. The process extracts the RNA in about 20 minutes. This isolated RNA is then moved to another system where the solvent is transferred into a minor well-connected to an electronic chip that is not more significant than that of the human thumb.

The RNA is activated by the reagent in the miniature well; it is the chip loaded with all viral load data to help determine whether or not the person carries the virus. Compared to traditional PCR tests, the reagents may not demand high heat in this method 74, with a smaller amount of swabs required for diagnostics. The main difference seems to be that the system is portable and costs much less than the conventional RT-PCR tests 72.

### **Impact of SARS-CoV-2 on the economy**

The coronavirus outbreak has shown a tremendous downfall in the global economy since its emergence. The lockdown was applied across the globe, which halted the economic movement. Stock markets took a worldwide hit, and the human race started finding high financial difficulty. Investors fear that the spread of the coronavirus will destroy economic growth, and it has already curbed the investments funds for businesses. In response, central banks in many countries across the globe have slashed interest rates. Global markets recovered some ground when governments worldwide started to drop some decent funds in the economy for its boost. The IMF (International Monetary Fund) defined this as similar to the great depression of the 1930s. While coronavirus is said to have plunged the world into a worst-hit epidemic, global growth is projected to increase to 5.8% next year if the pandemic declines in late 2020 75. Technology usage across the globe saw a surge as the government urged companies to work from home. Zoom (a video call)/meetup platform saw a massive uplift in share pricing during this period. Also, Netflix and Amazon saw a surge in their shares. Heavy lockdown triggered drastic changes in economic conditions and significantly declined the pollution level. Countries across the globe got their nature back in shape, with pollution levels dropping noticeably across the globe 75. Now easing lockdowns in various countries are helping the economy to move better than the March to May period, and even pollution levels in different countries are returning to normal at a gradual pace 75.

### **Mortality Rate of COVID-19**

The mortality rate of COVID-19 appears to be higher than influenza. Better understanding the actual death rate of COVID-19-infected people will take some time; the data sh that the crude mortality (the number of confirmed deaths divided by recorded cases) is between 3-4%. The infection fatality rate (the number of confirmed deaths divided by the number of infections would be lower. Mortality for seasonal influenza is typically well below 0.1 percent. Mortality, however, is largely determined by healthcare access and quality available in a given country 76.

As of January 1st, 2022, India had reported over 35 million cases of SARS-CoV-2, the virus that causes COVID-19. This placed the country as the second-highest in the world for confirmed cases, behind only the United States 77. The official cumulative COVID death count in India at the time was 0.48 million, translating to a death rate of approximately 345 per million population, which was one-seventh of the death rate in the United States 78.

However, it was widely believed that India's reported COVID death totals were underreported due to incomplete certification of COVID deaths and misattribution to other causes, as well as the fact that most deaths occurred in rural areas without medical attention 79,80. The United Nations Population Division (UNPD) estimated that India had up to 10 million deaths in 2020, with over 3 million of those deaths going unregistered and over 8 million not undergoing medical certification. As of November 9th, 2022, the United States has reported 1,070,947 COVID-19-related deaths, 81. During the first two years of the pandemic, COVID-19 emerged as one of the country's top three leading causes of death, ranked behind only heart disease and cancer 82,83.

The COVID-19 pandemic in 2020 and 2021 was marked by the widespread transmission of the SARS-CoV-2 virus within households, including among multiple generations, with a high prevalence of antibodies being detected. 84. The elevated death rate from COVID-19 in India in 2021 compared to the rate observed in 2020 requires further investigation. The dissemination of the virus to rural areas is a potential contributing factor. Still, variations in the pathogenicity between the Wuhan strain of the virus in 2020 and the Alpha and Delta variants that dominated the 2021 wave 85, as well as changes in biological markers predictive of severe infections, may also play a role. The continuous monitoring of COVID-19 death rates is crucial in assessing the impact of the ongoing Omicron wave in India and any future viral variants that may emerge.

### **Precaution, Prevention, and Treatment**

Medical companies and researchers across the globe are in a race to develop medication and vaccines for COVID-19 disease. Many companies are in the clinical trial phase, trying to expedite the steps of the clinical trial phases. Nevertheless, for remedial and preventive measure purposes, many governments in different countries have authorized various treatment procedures as valid steps for preventive measures till the vaccine or medicine is developed. WHO recommended washing hands properly with soap, using a 70% alcohol-based hand sanitizer, which helps kill the viruses. When anyone goes outside, it is suggested to use proper masks and gloves that can help avoid getting an infection and transmitting the infection the others. Also, maintaining at least a 1-meter physical distance is indicated as one of the significant precautions 86.

### **Ayurveda is an as natural treatment**

Ayurveda is a gift of mother nature and an ancient traditional method practiced in treatian diseases and other health disorders. Various doctors and government agencies in India are recommending Ayurvedic medicines to manage pre-symptomatic and asymptomatic COVID-19 patients. Basically, the Ayurvedic concept suggests ta hat boost in immunity can lower the risk of severe infection and help the fast recovery of the patients 87. Therefore, Ayurveda suggests consuming immunomodulatory herbs, which can enhance immunity. A few suggested ways are -

1. Consume decoction once or twice a day made from Ocimum basilicum (basil), cinnamon, Piper nigrum (black pepper), Zingiber officinale (dry ginger), and raisin (kishmish).
2. Golden Milk is also useful for enhancing immunity power. For its preparation, turmeric is added to hot milk, which is recommended to consume once or twice a day.
3. Practise drinking lukewarm water and use common spices such as turmeric, cumin, coriander, and garlic in daily food 87.

### **Homeopathy**

Homeopathy is an alternative medication easy to consume with no side effects. It is believed that homeopathic medicines are able to boost immunity. The Ministry of AYUSH, Government of India, released a health guideline and suggested that drugs from homeopathy and Unani could be beneficial in controlling SARS-CoV-2 infections 88. Arsenicum album 30 of homeopathic medicine has been meant to be used empty stomach daily as a prophylactic treatment against the infection for three days. The same dose must be repeated every month following the same schedule 88. However, there is no evidence that Arsenicum album 30 holds any preventive properties for COVID-19.

1. **Allopathy**

Allopathy refers to science-based conventional medication which targets the infectious agent or biological moieties to suppress symptoms. For the prevention of COVID-19, many pharmaceutical firms have shown their promptness to come forward for manufacturing testing kits, vaccines, drug development, and assessing the effectiveness of drug repurposing. We describe below some of the medicines undergoing phase I, II, and III trials, and a few are even being used as treatment alternatives alone or in combination.

#### **10.1. Chloroquine and hydroxychloroquine**

The Food and Drug Administration (FDA), USA, has only approved the use of hydroxychloroquine in emergency circumstances in people diagnosed with SARS-CoV-2 who are hospitalized or clinically enrolled. It is advised to stay under a doctor’s supervision while using hydroxychloroquine for COVID-19 89. The medications presently being studied for repurposing to treat COVID-19 appear to fall into two categories: those that control the viral replication process and those that seek to manage the disease-specific symptoms. Chloroquine and hydroxychloroquine aminoquinolines are generally taken as antimalarial medications. They are helpful in malaria to inhibit heme polymerase, which causes the parasite to accumulate toxic heme that may lead to death. In COVID-19, it is envisaged that the drugs can help avoid the interaction of the virus with the host cells by blocking host receptor glycosylation and breaking down viral protein production by inhibiting endosomal acidification. Although initial studies appeared promising, the research groups noticed various experimental flaws. In a randomized study of thirty patients in China who were found positive for COVID-19 at the same time, it was observed that they had no benefit over the standard treatment and that hydroxychloroquine administration could not help decrease death among hospitalized COVID-19 patients 90.



**10.2. Dexamethasone**

It is a low-cost steroid that is widely used. Although this matter is under discussion and further examination, researchers confirmed from the latest recovery study that the death rate of COVID-19 patients noticeably decreased following dexamethasone administration. This is an anti-inflammatory medicine widely used to treat disorders wherein the body's immunity is not working usually and triggers damage. Dexamethasone significantly lowers the level of inflammation and decreases the activation of the immune function by influencing the functioning of white blood cells. Dexamethasone falls under the corticosteroids category that closely matches the cortisol hormone released by human adrenal glands. This is widely used to treat chronic rheumatological disorders such as swelling of the muscles, irritation inside the blood vessels, persistent joint pain, and immune disorders. This is generally prescribed for cardiovascular disease, dysfunction of the kidney,s and skin irritations, and also to minimize swelling of the brain and spinal tumor. Corticosteroid treatment was used earlier during the SARS epidemic 2003 to help alleviate inflammatory lung injury 91. As a result, several countries are studying the effectiveness of corticosteroid treatment in COVID-19 acute respiratory infection patients. The WHO has listed it as a priority for the review of its clinical trials with a view to determining safety and effectiveness.



The WHO has advised the management of viral pneumonia "under the regimen of systemic corticosteroid" in an interim recommendation on COVID-19 therapy published on 27 May 2020 92. However, a recent systematic examination and analysis of the effects of dexamethasone treatment on people infected with coronavirus found that dexamethasone did not lower the mortality rate, did not decrease the time of hospitalization, did not decrease the ICU admission rate and oxygen treatment, many harmful consequences 93. In comparison, a recent study conducted by Oxford researchers, in which 2104 admitted patients received dexamethasone 6 mg for ten days, revealed that the drug significantly reduced deaths in patients on respirators by one-third and patients on oxygen therapy by one-fifth. The drug was shown to reduce the mortality rate by 17 percent after 28 days, with an "extremely important" pattern indicating "greatest gain" in patients needing ventilation 94. However, the experiment did not research patients beyond the clinical setting.

#### **10.3. Corticosteroids**

In parallel, scientists are investigating molecules to counteract the possible 'cytokine wind' in some patients, which results in lung damage which causes respiratory issues. Although immunomodulators are a promising candidate for this purpose, they may have adverse downstream effects, such as raising the likelihood of certain infections. Corticosteroids are the principal of these immunomodulatory drugs being studied for COVID-19 treatment. These drugs are well studied but are also known to be one of the most-blunt tools for immune system mutation. In addition, cardiovascular disease and the loss of bone density are associated with its long-term use. A previous meta-analysis showed that using corticosteroids in people with flu pneumonia correlated with a high death rate 95. Another retrospective study in China found that its use was associated with reduced death among those who developed ARDS. Clinical studies of these drugs, such as methylprednisolone and glucocorticoids, are still underway 96,97. The experts also advise thata higher glucocorticoid dosage could prolong coronavirus elimination due to immunosuppressive results. On 25 May 2020, the Lancet released a paper stating that "incorrect use of systemic corticosteroids can raise the risk of femoral head osteonecrosis (ONFH)” 91,98. Osteonecrosis leads to bone tissue mortality due to a loss of oxygen flow. Even the WHO suggests that regular corticosteroids must be avoided "given the lack of effectiveness and potential damage" unless they are indicated for some cause. While advising patients about their medications, it is strongly recommended to consider other health-related factors, like upper respiratory infection or chronic obstructive pulmonary disease (COPD), organ failure, and specific patient peril/satisfaction assessments 92.

**10.4. Lopinavir and Ritonavir**

In vitro experiments have found that lopinavir and ritonavir suppress 3-chymotrypsin-like protease activity; that is why this drug is also known as human immunodeficiency virus protease inhibitor. This drug was reported to be effective against previously emerged coronaviruses; however, there is no evidence that it can work against COVID-19 as well. In China, a randomized study was performed on 200 hospitalized patients but did not produce an effective outcome for the support of this drug combination in comparison to standard care. According to the Journal of the American Medical Association, this drug shows some side effects such as increased nausea, diarrhea, and risk of liver damage similar to the symptoms of SARS-Cov-2. New England Journal of Medicine published a randomized controlled study, which reveals that the drug is not beneficial to treat COVID-19 patients and, therefore, not helpful in the recovery of SARS-CoV-2 infected patients. Another trial on mild COVID-19 patients revealed that those who were treated with lopinavir, ritonavir, IFN-B, and ribavirin recovered fast, stayed for a short time in the hospital, and minimized symptoms noticeably, as compared to those receiving lopinavir and ritonavir alone 99.



#### **10.5. Nafamostat and Camostat**

Nafamostat and camostat are approved for use against human pancreatitis in Japan. Both are inhibitors of serine protease. In an in vitro study, camostat was found to work as an antagonist to the TMPRSS2 serine protease and to block SARS-CoV from reaching the host cells 12. Researchers claimed that both compounds may have inhibiting effects on SARS-CoV-2 as well. Recent in vitro tests have displayed that both can block COVID-19 entry into cells, while a preprint study stated that nafamostat blocks the virus entry inside cells 15-fold more efficiently than camostat 100. In the USA and Japan, these drugs undergo phase II 101 and phase II/III 102 clinical trials to assess their effectiveness against SARS-CoV-2.



**10.6. Famotidine**

Famotidine is an H2 receptor blocker 103,104 and is thought to be a potential candidate for treating patients infected with SARS-CoV-2. In China, Michael Callahan and colleagues reported that patients taking famotidine heartburn medication seemed less affected or less likely to need intubation during severe COVID-19. No such peer-reviewed results were released as a preprint 103. A randomized phase III clinical trial is being performed in New York, in which COVID-19 patients who are in critical condition were received intravenous famotidine with hydroxychloroquine. However, the mode of action of the drug is unknown 105. A Virginia-based biodefense consultant, Robert Malone, who is working on the famotidine, proposed a hypothesis that famotidine binds a papain-like protease encoded by the COVID-19 virus genome and is recognised as a critical component of viral entry into cells; however, none of the assay findings experimentally support this hypothesis 104.



#### **10.7. Umifenovir**

Umifenovir is approved as a prophylaxis for the prevention of influenza viruses A and B only in Russia and China, and several research groups have presumed that it has large-spectrum antiviral properties; however, there is no data available yet to assist its effectiveness against SARS-CoV-2. Umifenovir is a small indole derivative hydrophobic molecule that interacts with lipids and proteins. It targets viral lipid membranes and inhibits contact between the virus and host cell, thus, preventing virus entry into target cells 106,107. A randomized phase III clinical trial is currently being conducted to examine the efficacy, safety, and tolerability of an anti-viral umifenovir drug.



**10.8. Nitazoxanide**

Nitazoxanide is an antiparasitic and antiviral thiazolidine drug used as a medicine for the treatment of parasitic, bacterial, and viral infections. This drug acts by blocking the maturation of viral capsid N protein, which helps the formation of viral particles. In clinical trials, it is being tested with other anti-parasitic drugs like ivermectin and also against hydroxychloroquine 108.



**10.9. Ivermectin**

Ivermectin is a lipophilic macrolide medication used to treat many parasitic infections and is usually used as an anti-parasitic drug. It acts upon parasitic cell membranes, wherein glutamate-gated chloride ion channels are present, increasing the permeability of the cell membrane and resulting in paralysis or death of the parasite. Scientists at Monash University in Melbourne, Australia, have observed the efficacy of this drug against COVID-19 in an in vitro experiment 109. In addition, clinical studies ought to be performed to support its effectiveness in humans with COVID-19.



**10.10. Tocilizumab and sarilumab**

In clinical trials, drugs such as tocilizumab and sarilumab that block cytokines are currently being tested. These drugs are monoclonal antibodies that serve as IL-6 receptor inhibitors and are commonly utilized in treating chronic inflammatory conditions that badly affect joints and damage various body systems. An unarranged controlled trial showed very positive results 110. Tocilizumab is the first IL-6 inhibiting antibody approved and has demonstrated its protection and efficacy in rheumatoid arthritis therapy 111. Another RA-approved IL-6 receptor blocker, Sarilumab, has been studied in a multicenter, double-blind, phase 2/3 trial in hospitalized severe COVID-19 patients 112. From the affiliate Hospital of the University of Science and Technology of China and Anhui Fuyang Second People's Hospital, twenty-one sufferers identified with serious COVID-19 were selected and administered tocilizumab therapy to check whether the targeted interleukin-6 (IL-6) could be an efficient and helpful way to reduce COVID-19 fatalities. The outcomes of tocilizumab therapy are promising. The temperature of all patients stabilized very rapidly, and the respiratory function and all other conditions improved considerably. Of these 21 patients, 20 were stabilized and released within 2 weeks of treatment. No adverse drug reactions were reported during treatment with tocilizumab 111,113–115.

#### **10.11. Bevacizumab**

Bevacizumab is a monoclonal antibody typically used in various types of cancer against the signaling protein vascular endothelial growth factor (VEGF). In China and Italy, researchers are conducting clinical trials of Bevacizumab. This drug helps in suppressing tumors by inhibiting the growth of blood vessels that are supplied to the tumor. This drug is also helpful in the reduction of vascular permeability and thereby lowering the volume of fluid that enters the lungs 116.

#### **10.12. Fluvoxamine**

Fluvoxamine is an FDA-licenced antidepressant medicine utilized to cure obsessive-compulsive disorder and is a better source of immunomodulation. According to an animal study, fluvoxamine in cells shuts down the inflammatory cascade from the endoplasmic reticulum when it binds to sigma-1 receptors 116. A clinical trial is still underway at the Washington University School of Medicine in St Louis, Missouri, USA, to investigate the significance of this medicine for SARS-Cov-2 therapy 117.



#### **10.13. Favipiravir**

Favipiravir is mainly produced by Toyama Chemical Co. Ltd. in Japan. Favipiravir is an antiviral medication, which is an analog of modified pyrazine used to treat influenza in Japan. Favipiravir is a pro-drug converted by a host enzyme into its active form ribofuranosyl-5’-triphosphate, and inhibits RNA virus through selective inhibition of viral RNA-replicase. RNA-replicase helps RNA virus’s replication and transcription to make multiple copies of their genome. In 2014, Favipiravir was licensed for avian influenza storage in Japan and is now an alternative option for influenza strains resistant to neuraminidase inhibitors. Favipiravir has been approved to treat Ebola, Lassa viruses and is being used for curing SARS-CoV-2 118–120.



On June 20, 2020, Glenmark, India, issued a press release. It announced that the company, with the help of the Research and Development team, successfully developed a FabiFlu, an active pharmaceutical ingredient, and formulation, and claimed that favipiravir shows 88% clinical improvement in COVID-19 and lowers the viral load within 4 days 121. In India, Glenmark conducted a randomized multi-center drug efficacy and safety study on mild to severe Covid-19 Indian patients with a standard of mixed health care vs. standard of care alone. The study included one hundred fifty patients. The Chinese National Medical Products Administration licensed favipiravir as the very first anti-SARS-Cov-2 medicine in China, as the clinical research indicated effectiveness with minimal adverse effects. A retrospective, randomised, controlled, open-label, multicenter study conducted by Chen et al. included adult patients with SARS-CoV-2 122.

### **Stem cell-based therapy for the treatment of SARS-CoV-2**

The emergence of severe acute respiratory syndrome coronavirus-2 in the first half of the 21st century has shaken the world and scientists. This pandemic has enforced many researchers to find solutions to overcome this situation and ensure the life of others. Many researchers and scientists from different life sciences or engineering fields are working continuously for alternative solutions and developing therapeutic approaches to bring everyday life back on track. In this direction, efforts related to stem cell-based therapy have also been found to be very positive as an efficient approach for treating COVID-19 patients (Figure 2). The mesenchymal stem cells ( MSCs) could be used as one of the potential strategies to prevent and cure SARS-CoV-2 infection recently announced by the International Society for Stem Cell Research (ISSCR) 4,123.

Cell-based experiments have currently proved to be one of the potential methods that provide treatment options for several diseases that were previously considered to be incurable. Because MSCs are safe from legal and ethical issues and have a great potential to make a copy of themselves within a short time, and have less invasiveness, MSC therapy is preferred over alternative treatments. MSCs can be derived from a wide variety of adult tissues, including bone marrow, adipose tissues, dental pulp, neonatal birth-associated tissues, placenta (PL), umbilical cord (UC), Wharton jelly (WJ), amniotic fluid (AF), cord blood (CB), fetal liver and Bichat fat pads, etc. Mesenchymal stem cells are known to be multipotent in nature and generally stored for future possible therapeutic purposes. Also, no unfavorable reaction toward allogeneic MSCs has been seen in the clinical trials of MSCs therapy 4.

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**Figure 2.** Stem cell-based therapy as an efficient approach for the treatment of COVID-19 patients.

MSCs are opposed to viral contamination due to the existence of enhanced quality of particular cytokines. These features are present in the inner niche of the MSCs before the separation process. Therefore, even if grafted to a recipient with verified SARS-CoV-2, MSCs may be expected to survive. The concept of MSC therapy in patients with COVID-19 is thus promising (Table 2). MSC therapy is inclined to prevent the inflammatory processes by the body's defense system and facilitate intrinsic reconstruction by the regenerative qualities of the stem cells 4. After injecting directly into the vein, some of the MSC population is considered to be trapped in the lungs, sometimes referred to as a restriction of systemic infusion. Nevertheless, these MSCs can restore the lung microenvironment, shield epithelial cells, stop lung damage, and treat pulmonary impairment and SARS-CoV-2 infection 4. The fundamental limitation of this therapy is the availability of therapeutic-grade MSCs. As a result, MSCs could be the ideal candidate for research studies or beneficial strategies for curing patients infected with coronavirus-2 4,13.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Clinical trial number** | **Diseases** | **Cell type as a treatment** | **Phase** | **Reference** |
| **NCT04336254** | COVID-19 | Allogeneic human dental pulp stem cells | Phase I/II | 124 |
| **NCT04366323** | SARS-CoV-2 | Allogenic and expanded adipose tissue-derived mesenchymal stem cells | Phase I/II | 125 |
| **NCT04366063** | SARS-CoV-2 | Mesenchymal Stem Cell | Phase II/III | 126 |
| **NCT04392778** | COVID-19  Pneumonia  Multiple Organ Failure  Corona Virus Infection | Mesenchymal Stem Cell | Phase I/II | 127 |
| **NCT04355728** | Corona Virus Infection  ARDS  Human Acute Respiratory Distress Syndrome  COVID-19 | Umbilical Cord Mesenchymal Stem Cells | Phase I/II | 128 |
| **NCT03042143** | Acute Respiratory Distress Syndrome (ARDS) | Human umbilical cord-derived CD362 enriched MSCs | Phase I/II | 129 |
| **NCT04288102** | Corona Virus Disease 2019 (COVID-19) | Human Umbilical Cord-derived Mesenchymal Stem Cells | Phase II | 130 |

**Table 2:** List some clinical trials using stem cells to treat COVID-19 patients.

Clinical trials have recently started in the United States, China, Jordan, Iran, and several other nations, and some reports have also been released. In particular, mesenchymal stem cell (MSC) therapy has been widely used to cure diabetes mellitus, autoimmune disorders, nerve damage, and many different illnesses 4. The immunomodulatory properties and differentiating ability of MSC are therefore intended to stop the dying of pulmonary tissue by combating cytokine storms and helping to restore and restructure damaged tissues. Clinical treatment of avian influenza with similar consequences on the lungs by using these cells has recently been shown in the study 131. Similarly, a recent case study of a woman who suffered from acute SARS-CoV-2 infection in China has shown that her laboratory tests and CT images have improved significantly after 21 days of curing with funiculus umbilicalis MSCs 4,132. In one other study, the patient was medicated with lopinavir/ritonavir drugs which stop the growth of the virus, and also with moxifloxacin, xuebijing, and methylprednisolone injected directly into veins, which resulted in an 87% rise in neutrophil and a decrease of 9.8% in lymphocyte. The infected person was also exposed to non-invasive mechanical ventilation due to insufficient oxygenation to improve respiration and alleviate muscle fatigue. The infected person was treated with an MSC cord and 5×107 thymosin α1 cells every 3 times as vital symptoms worsened. The test results showed that blood albumin, C-reactive protein, and aspartate transaminase/alanine transaminase decreased gradually after the second dose, and in parallel, this improved life-sustaining signs. Subsequently, the infected person was discharged from the life support machine and found to be able to move. And, the amount of WBCs and neutrophils in the infected person lowered to usual, whereas the amount of immune cells continued to increase to average. More significantly, the number of immune cells increased dramatically. In addition, qualitative studies obtained from CT scans and pictures during the second and third intravenous administration of umbilical cord stem cells have shown that respiratory disease could have been somewhat relaxed. Subsequently, after the third injection, the patient was released from the intensive care unit, and most of the essential standards, like pulse rate, heartbeat, etc., were fine. The findings indicate that umbilical cord mesenchymal stem cell therapy can be an option for SARS-CoV-2 infected persons, separately or with other immune modulators 4,132. Mesenchymal stem cells have the potential to alter the behavior of immune cells 133. In another recent research conducted in China in alliance with the USA, from January 23 to February 16, 2020, in Beijing Youan Hospital, seven patients with COVID-19 pneumonia who experienced mesenchymal stem cell transplantation improved their health outcomes and normalization in the immunological profile 13. Furthermore, genetic analysis of MSCs suggests that they are ACE2- and TMPRSS2-;, revealing that MSCs cannot directly interact with SARS-CoV-2 13. Studies indicate that mesenchymal stem cells could be secure and beneficial in treating infected persons with SARS-CoV-2 pneumonia, particularly people with severe illnesses.

### **Some of the potential vaccines which are under research, testing, and trial**

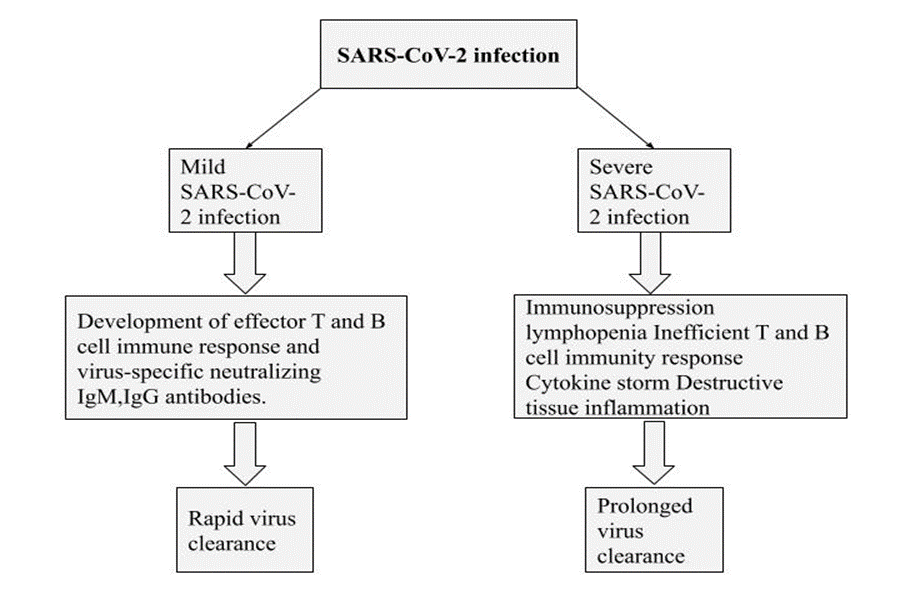
Many companies are developing vaccines against coronavirus alone and some in collaboration. The vaccines which are currently under test and trials are as follows:

1. The National Institute of Allergy and Infectious Diseases (NIAID), USA, conducted a trial and review of preliminary phase 1 results for the mRNA-1273 vaccine called mRNA-1273, and Moderna company announced and published satisfactory positive interim phase 1 results for mRNA-1273 against COVID-19. 134. On July 27, 2020 phase III trial began in collaboration with NIH (National Institute of Health) and BARDA (Biomedical Advanced Research and Development Authority) 135 and 30,000 participants enrolled in the study received one intramuscular (IM) injection of 100 micrograms (ug) mRNA-1273 on Day 1 and Day 29, and some participants received 1 IM injection of mRNA-1273-matching placebo on Day 1 and Day 29 136. On November 16, Moderna informed that in the first interim review of the Phase 3 COVE trial, mRNA-1273 reached its primary efficacy endpoint with a vaccine efficacy of 94.5 percent 135.
2. The Jenner Institute at Oxford University developed AZD1222 (formerly ChAdOx1 nCoV-19), with AstraZeneca in charge of global production, processing, and marketing. A Phase I / II clinical trial of AZD1222 began in April 2020 to determine the vaccine's safety, immunogenicity, and efficacy in over 1000 healthy volunteers across multiple trial centers in southern England 137. A total of 23,848 participants were enrolled between 23 April and 4 November 2020, and in the interim primary efficacy assessment, 11,636 participants were included (7,548 in the UK, 4088 in Brazil). Vaccine efficacy was 62.1 percent in participants who received two standard doses (95 percent CI 41.0–75.7; 27 [0.6 percent] of 4440 in the ChAdOx1 nCoV-19 group vs71 [1.6 percent] of 4455 in the control group) and 90.0 percent in participants who received a low dose followed by a standard dose 138. The overall effectiveness of the vaccine in both groups was 70.4 percent. ChAdOx1 nCoV-19 has an appropriate safety profile and is effective against symptomatic COVID-19 in this interim study of ongoing clinical trials 138.
3. "Ad5-nCoV" is a recombinant new coronavirus vaccine (Adenovirus Type 5 Vector) developed and approved by Biologics Inc. with the Beijing Institute of Biotechnology (BIB), perform a Phase 1 clinical trial in China. The first-in-human study in 153 healthy adults showed that the Ad5nCoV vaccine is tolerable and immunogenic. After post-vaccination, on day 28, it showed specific antibody-mediated immunity responses to coronavirus-2019, and on day 14, it observed rapid T-cell response 139. A global phase III clinical trial begins on 15 September 2020 involving 40,000 healthy adults aged 18 years and older participants to examine the efficacy, safety, and immunogenicity of Ad5-nCoV developed by the Cansino and Beijing Institute of Biotechnology. The immunization protocol is one dose of intramuscular injections (deltoid) 140.
4. INO-4800: Inovio Pharmaceuticals was the first to carry out a human evaluation of its MERS-CoV 151 related coronavirus vaccine (INO-4700) 141, and for prevention of COVID-19 infection company developed a DNA vaccine (INO-4800). Inovio also performed an open-label trial 142 in healthy adult volunteers using a CELLECTRA ® 2000 device to evaluate the effectiveness, tolerability, and immunological profile of INO-4800 delivered by intradermal (ID) injection accompanied by EP. A Phase 2/3 randomized, blinded, placebo-controlled study begins on November 30, 2020, with 6578 participants enrolled to determine the safety, immunogenicity, and efficacy of INO-4800, a prophylactic vaccine against COVID-19 disease, injected intradermally with electroporation in healthy seronegative adults at high risk of SARS-CoV-2 exposure 143.
5. NVX-CoV2373: Novavax, Inc. announced the commencement of clinical trial phase I/II for the development of coronavirus vaccine, NVX‐CoV2373, a robust perfusion protein produced using its patented nanoparticles adjuvant (Matrix‐MTM) technology to boost immunity and trigger a high level of antibody neutralization. In the southern hemisphere, the first human trial to be carried out, with approximately 130 participants spanning two Australian locations 144. A Phase 3 trial began on December 27, 2020, and enrolled 30,000 participants with mild, moderate, or severe symptomatic coronaviral disease 2019 (COVID-19). This trial will focus on the body’s immune response and the safety of SARS-CoV-2 rS with Matrix-M1 adjuvant in participants. The enrolled participants in the problem will be randomized to receive SARS-CoV-2 rS with either Matrix-M1 adjuvant or placebo. A total of 2 intramuscular injections will be given to each of the participants in the study 145.
6. LV-SMENP-DC and pathogen-specific APC vaccine developed by Shenzhen Geno-Immune medical institute are in phase 1 clinical trials to determine security, effectiveness, and immune response 146,147.
7. For the development of an investigative vaccine against COVID-19, Merck has announced a collaboration with the International AIDS Vaccine Initiative (IAVI), using recombinant vesicular stomatitis virus (rVSV) technology, which forms the framework for its Ebola Zaire Virus (Ervebo) 148. A randomised, placebo-controlled, double-blind study is being performed at seven locations in the U.S. and will involve up to 252 participants aged 18 and up, including 149 adults.
8. BNT162: Pfizer Inc. & BioNTech SE reported that in phase I/II clinical trial for the BNT162 vaccination system to combat COVID-19, the first patients were administered in the U.S. the first patients in phase I/II clinical trial for the BNT162 vaccination system to combat COVID-19 assess the vaccine's safety and immunogenicity 150. Another mRNA vaccine (BNT162), approved for clinical trial phase ½, has, been produced collaboratively by BioNTech and Pfizer to evaluate the appropriate dose for further studies and to assess the safety and immunogenicity of the vaccine 59. A Phase I/II trial, 2-Part, begins on September 9, 2020, sequentially administering increasing doses of an investigational drug to examine the safety and immunogenicity of prophylactic SARS-CoV-2 RNA vaccine (BNT162b3) against COVID-19 utilizing different dosing protocols in healthy adults 150.
9. India's first indigenous COVID-19 ‘COVAXIN’ vaccine, developed by city-based Bharat Biotech in partnership with ICMR and NIV, has received the nod from India's Drug Controller General for human clinical trials, the company said on June 29, 2020. The phase I/II clinical trials of the SARS-CoV-2 vaccine, which were approved after pre-clinical studies showed safety and immune response, would begin nationwide in July 2020 151. Following the successful completion of the interim phase, 1 & 2 clinical trials of COVAXINTM, Bharat Biotech enrolled 26,000 participants after getting DCGI clearance for phase 3 clinical trials over 25 centres across India 149.

India has completed 1 billion vaccine doses and is now capable of testing approximately 1 million samples each day. Existing Virus Research Diagnostic Laboratory for viral diagnosis and INSACOG labs for genome sequencing are already in place to identify and monitor the present Omicron or any novel variant. Early and active surveillance, as well as whole-genome sequencing, will aid in understanding circulating variants and tracing their progression. According to COVID-19, appropriate behavior such as social distancing, hand hygiene, mask use, and vaccines remain the most critical elements in viral transmission management.

### **Immune responses in the human body against SARS-CoV-2**

Pathogenic cell infection can cause development of humoral and cellular immunities in the host, which are essential for removing the viral infection 152. However, uncontrolled or ineffective immune reactions can cause adverse effects on the patients 153. New immunotherapies will be created by proper understanding of the immune response induced by infection with SARS-CoV-2, along with reducing the possible risk of inflammation 152. About 80 percent of COVID-19-infected people record moderate to negligible side effects, taking into account immunopathological aspects 5,6. In severe COVID-19 cases, patients suffering from lymphopenia and interstitial pneumonia have elevated levels of proinflammatory cytokines, including interleukin-10 (IL-10), interleukin-2 (IL-2), interleukin-6 (IL-6), interleukin-7 (IL-7), colony-stimulating factor-3 (CSF-3), interferon gamma-induced protein-10 (IP-10), monocyte chemoattractant protein-1 (MCP-1), macrophage inflammatory protein-1 (MIP-1), macrophage inflammatory protein-1 alpha (MIP-1α) and tumor necrosis factor-alpha (TNFα) 5. Immune cells and their subsets play a vital role in the process of protecting from infected cells 5. Mild SARS CoV-2 infection can lead to neutralizing antibodies, sufficient immune cell response, and rapid viral clearance, whereas severe SARS CoV-2 infection can induce serious illness and delayed viral clearance 154 (Figure 3). Due to viral infectious diseases, immune cells like lymphocyte subsets may also be dysfunctional 155–157. Innate immune response and cell-mediated immune response against viral infection involve the following cells: CD3 +, CD4 +, CD8 +, CD16 +, CD56 +, and CD19 + label T-helper cells (CD4+CD3 +) and T-cells (CD3+CD8 +), B cells (CD19 +) and natural killer cells (CD16+CD56+) 155. Many studies are currently identifying the adverse effect of COVID-19 infection on immunity and immune cells. Most of these have revealed that infected people develop an uncontrolled immune response during the infection, triggered by highly active macrophages and monocytes 5,158. Recently several epitopes have been reported through immuno-informatics, including 5 CTL epitopes, 3 sequential B cell epitopes, 5 discontinuous B cell epitopes of immune cells 4,159, and 13 MHC-I and 3 MHC-II antigenic epitopes 4,160, and some of these epitopes are expected to help in the manufacturing of potential vaccine for COVID-2019 4.

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**Figure 3.** Distinct responses to mild and severe SARS-CoV-2 infections.

**Response of T-cell and B-cell**

Immune cells such as T-lymphocytes for antibody-independent directed immunity and B-cells for antibody-mediated immunity play a significant role in the adaptive immune response to all viral infections. Nonetheless, helper T lymphocyte activation of Th1 / Th17 results in the alleviation of the inflammatory effect, while B lymphocytes process the production of specific SARS CoV-2 antibodies for virus neutralization 5,161. After reaching the host, the virus binds and invades the cells expressing its particular replication receptor. The human leukocyte antigen (HLA) system, such as class I and II, displays viral antigenic peptides to appropriate T-cells, such as CD8+ and CD4+. Class II HLA system is represented only on professional antigen-presenting cells (APC) such as dendritic cells and macrophages 161 to display processed antigenic protein or whole viruses to helper T-lymphocyte CD4+ molecule 154,162. In the case of COVID-19, once the coronavirus is within the lung epithelial cells, viral peptides are introduced to CD8+ cytotoxic T cells using Class I major histocompatibility complex (MHC) proteins, followed by Class II major histocompatibility complex (MHC) with larger peptides 154,163,164. When CD8+ cells start dividing and showing clonal expansion, virus-specific effectors and T-cell memory develop. Cytotoxic T-cells that express CD8+ receptors are lysed with tissue cells infected with the virus. B cells can interact with CD4+ T cells and directly recognize and activate viruses. In the first week of symptoms, IgM isotype primary virus-specific antibody response is observed 154,164.

After dealing with the study of past events or situations of COVID-19 patients, the outcomes displayed a significant reduction in total lymphocyte counts, B lymphocytes, Natural Killer (NK) cells, CD4+ and CD8+ T cells 154,155,165,166, particularly in patients requiring emergency unit. Patient survival was inversely associated with total T cells, cytotoxic T-lymphocytes (CD8+) cells, and helper-T lymphocytes (CD4+) cells below 800/μL, 300/μL, and 400/μL, respectively. T lymphocytes counts were inversely correlated with concentrations of serum interleukin-6, interleukin-10, and tumor necrosis factor-alpha (TNF-α), with patients showing reduced concentrations of interleukin-6, interleukin-10, and tumor necrosis factor-alpha, and restored T lymphocytes counts during disease resolution period. T-lymphocytes are found to have substantially more significant levels of the depleted PD-1 marker in the infected patients. As the patient’s condition worsened from prodromal to excessively symptomatic, increased levels of programmed cell death protein 1 (PD-1) and HAVCR 2 (Tim-3) were seen in T-lymphocytes 166,167.

Cytotoxic T-cells are essential for enabling virus removal after many acute pulmonary infections. Therefore, it is believed that combined activation of virus-specific cytotoxic T-lymphocytes (CD8+) and antibodies would provide optimum immune defense. Moreover, cytotoxic T-cells (CD8+) protect from secondary infections 154,155,168. In the recovered patients, virus-specific helper T-cell (CD4+) responses have more frequent evidence than virus-specific cytotoxic T-cell (CD8+) responses. In contrast, pre-existing helper CD4+ T-cell responses to other coronaviruses are also found in patients without SARS-CoV 2 exposures 169,170. In extreme cases, the mechanisms for a significant decrease in lymphocytes remain unexplained. The type of death of lymphocytes in COVID-19 should be thoroughly investigated 154.

**Response of Antibodies**

As of today, the convalescent plasma of more than one million patients has been considered clinically appropriate for treating infected patients with SARS-CoV-2, which provides passive immunisation. It is significant to mention that if a patient with severe pneumonia is found to have a heavy virus load, convalescent serum containing virus-specific immunoglobulin G1‐, immunoglobulin G2‐ and immunoglobulin G3‐type antibodies that induce local injury in the lung via complement activation and maximize damages of tissues, is observed. Immunoglobulin G4 must be context-specific for treatment for such cases, as no additional activating properties have been identified to date 154,171. At the onset of initiation of infection in COVID-19 patients, total antibodies (IgA, IgG, and IgM) and complement (C3/C4) protein levels were found within the normal range 154,158. However, high concentrations of IgG or IgM antibodies to COVID-19 N protein or receptor binding domain (RBD) were observed after infection of 10 days or later. A typical antibody response is an early increase in IgM followed by a noticeable increase in IgG after a few days. However, the serum level of the common IgG can be identified by shooting up earlier than that of IgM against SARS-CoV-2 154,172. These results have shown that, due to the presence of other coronaviruses, cross-reactivity of antigens with existing specific IgG may not be helpful for detecting COVID-19 154. In different studies, COVID-19 virus-specific IgG and IgM antibodies were found to reach their peak levels in 17-19 days and 20-22 days after the appearance of symptoms. Another observation was that in the severe COVID-19 category, the IgG and IgM titers were relatively higher than the non-serious category 172,173. It is widely recognized that during viral infection, IgM provides the first line of protection compared to the high-affinity IgG, which offers long-term immunological memory and immunity 5. The presence of IgM in the serum indicates the recent interaction with the virus, while the presence of IgG implies the interaction that occurred a few days ago. Nonetheless, there is still a lack of in-depth information on the human defense system's response to novel coronavirus infection, most of which relies on information obtained during the most recent outbreak of coronaviruses such as SARS 2002 and MERS 2012 76,161.

### **Vaccines associated antibody-dependent enhancement (ADE)**

The development of antibodies following a vaccine is typically regarded as a positive result. Antibodies (from immunizations or recovering from a past infection) are essential for our immune systems to fight a virus effectively. When we produce antibodies to the disease, our immune system can sometimes overreact the next time we are exposed to the disease. This is a highly uncommon condition known as ADE (Antibody-Dependent-Enhancement). The antibodies involved in ADE do not aid the body's immunological response and may even exacerbate it. When a person is infected, ADE raises the likelihood that they may develop severe disease symptoms. Antibodies that cause ADE behave like a "Trojan horse," allowing the virus into the cells and activating the immune response. They will enable the virus to bind to our cells, resulting in inflammation and an increased immunological response. Nevertheless, there have been no confirmed reports of ADE caused by COVID-19 vaccinations.

The virus particles adhering to the cytoplasmic membrane on the cell surface, where viral surface proteins attach to the host cell with specific receptors, is the first step in viral infection 174. Antibodies mainly targeting viral surface proteins are released to prevent viruses from attaching to target cells. These antibodies bind and destroy the viruses, reducing their infective ability. In some viruses, antibodies, instead of binding to viral surface proteins, can promote viral invasion into specific cell type, maximizing viral infection 174. This effect is referred to as antibody-dependent enhancement (ADE) 175. ADE happens both ways: (1) when a pathogen specific antibody facilitates entry of the virus into phagocytic cells (macrophages, monocytes, and granulocytes) and (2) when it enhances infection in cells via contact with the Fc receptor (FcR) and complement receptor. In most circumstances, enhancing viral attachment to target cells is critical. Over 40 different viruses have been found to have ADE. There are a range of antigenic determinants (epitopes) in these viruses, some of which trigger neutralizing antibodies and others which induce enhanced antibodies 174. Conventional vaccinations have only had a minor preventive and therapeutic effect on these viruses 176. In some cases, they have been proven to increase the susceptibility of those who have been vaccinated.

The likelihood of ADE, i.e., antibody-dependent enhancement in Covid-19, making the severity of conditions worsen, cannot be ignored entirely; this is considered one of the significant threats around antibody-based vaccines and treatments 177. Two Viruses, RSV (respiratory syncytial virus) 178,179 and measles 180,181, can go in worse states by antibody-dependent enhancement (ADE), which makes it risky. Increased respiratory disease (ERD) is a broader category that includes ADE in it, and non-antibody-based processes such as cell-mediated immunopathology and cytokine cascades are part of ERD 177. Often, when viral replication is boosted, then macrophage-infecting viruses, such as dengue virus 182,183 and feline infectious peritonitis virus (FIPV) 184, have been observed to produce ADE. SARS-CoV and MERS-CoV have been closely linked with ADE and ERD in both vitro and in vivo. Also, the role and impact of ADE in the immunopathology of COVID-19 remain to be explored 177.

# **14.1** **ADE in human coronavirus infections**

There is no conclusive evidence that ADE has a role in human coronavirus illnesses. Concerns regarding ADE were raised when seroconversion and neutralizing antibody responses were found to correlate with clinical severity and mortality in SARS patients 185. According to the findings in COVID-19 patients, more significant SARS-CoV-2 antibody titers are linked to more severe illness 186. Increased antibody titers in chronic COVID-19 patients are thought to result from more and more prolonged antigen exposure due to increased viral loads 187,188. A new analysis showed that asymptomatic and symptomatic COVID-19 189 individuals had similar viral shedding in the upper respiratory tract. Anti-SARS-CoV-2 antibody titers were noticeably higher in symptomatic people, and they cleared the virus from the upper respiratory tract faster, refuting the idea that higher viral loads cause higher antibody titers. According to other studies, anti-SARS-CoV-2 T-cell responses were seen at high levels in both symptomatic and asymptomatic infections 170,190. The data suggest that patients with a broad range of clinical symptoms can have potent T-cell responses, although high antibody titers are more strongly linked to chronic COVID-19. The fact that viral shedding was discovered in the upper respiratory system rather than the lower respiratory tract is a critical issue 189. The lower respiratory tract is likely more essential for severe COVID-19 lung pathology, and it's unclear how closely SARS-CoV-2 viral shedding in the upper and lower respiratory tracts correlates with chronic infection 177. Apart from the host response to specific SARS-CoV-2 illnesses, another potential worry is the probability of pre-existing antibodies against other human coronavirus strains triggering ADE in COVID-19 patients 191. Human antibodies elicited by endemic coronavirus variants (like OC43, NL63, 229E, and HKU1) could potentially enhance ADE by increasing SARS-CoV-2 cross-reactive identification in the lack of viral neutralization. According to early findings, antibodies (with high reactivity) from SARS-CoV-2-naive donors to seasonal human coronavirus strains showed slight cross-reactivity against the nucleocapsid and S2 components of SARS-COV-2 192. It would be interesting to know if such cross-reactive antibodies have a role in SARS-COV-2-triggered ADE.

**14.2 Mechanisms of ADE (Antibody-Dependent Enhancement of Coronavirus)**

ADE can be triggered by several different molecular pathways. According to one concept, the complex of antibody/Fc-receptor mimics viral receptors in function, allowing some phagocytic cells to engage in expanded host cell trophism 193. Wan et al. show that antibody dose affects whether a virus promotes disease or suppresses it. Antibodies to one virus strain are reported to be sub-neutralizing or non-neutralizing for viral infections of other variants 182,194,195. Infected cells expressing Fc-gamma were shown to be infected with SARS-CoV-1 196. A case of ADE was detected in a patient with a second SARS-CoV-2 infection 197. ADE has been demonstrated to arise in viral infections in two ways: FcR-mediated ADE is the first mechanism. FcRs are receptors that target the Fc regions of antibodies and are mostly found on immune cells. Enhanced antibody-mediated virus intake into Fc gamma receptor IIa (FcRIIa)-expressing phagocytic cells leads to increased viral infection and replication or increased antibody Fc-mediated effector activities or immune complex creation, resulting in increased inflammation and immunopathology.

Both ADE processes can happen when non-neutralizing antibodies or sub-neutralizing levels latch to viral antigens without suppressing or clearing the infection. In vitro assays (more frequently performed for the first mechanism, which involves FcRIIa-mediated enhancement of infection in phagocytes), immunopathology or pulmonary pathology can all be used to detect ADE. For macrophage-tropic viruses such as dengue virus in humans 198 and FIPV in cats 184, ADE via FcRIIa-mediated endocytosis into phagocytic cells has been reported and intensively studied in vitro. Non-neutralizing antibodies or binding antibodies bind to the surface of the virus and transport virions to macrophages, which uptake the virions and get infected. According to a recent dengue vaccine trial, because many antibodies against different dengue serotypes are cross-reactive but non-neutralizing, repeated infections with heterologous strains can increase viral multiplication and more severe disease, offering considerable safety risks 182,183. Cats immunized against the FIPV S protein or passively infused with anti-FIPV antibodies outlived control groups when challenged with FIPV in other vaccination studies 199.

The spreading of the infection, severe symptoms, and more dangerous sickness outcomes 200 could be caused by antibodies with sub-neutralizing levels or non-neutralizing antibodies. These antibodies have shown capabilities to promote the entry into alveolar and peritoneal macrophages, which support this spread and worsening of sickness 201. Antibody effector functions that are Fc-mediated can intensify respiratory sickness that leads to detectable and intensified lung damage by establishing a robust immunological reaction in the second known ADE mechanism; this is defined more perfectly as a respiratory infection 202,203. Fc-mediated activation of local and circulating innate immune cells such as monocytes, macrophages, neutrophils, dendritic cells, and natural killer cells, despite their potential efficacy in removing virus-infected cells and debris, may result in overactive inflammatory responses. Non-neutralizing antibodies have been shown to cause ADE and ERD in non-macrophage tropic respiratory viruses like RSV and measles by forming immune complexes that accumulate in airway tissues and initiate cytokine and complement mechanisms, resulting in infection, respiratory failure, and, in extreme cases, acute respiratory distress syndrome 178,180,204,205.

These prior observations of ADE with RSV and measles are strikingly identical to the clinical symptoms of COVID-19. In COVID-19 and SARS, for example, over-activation of the complement cascade has been shown to lead to inflammatory lung injury 206,207. S- and RBD-specific immunoglobulin G (IgG) antibodies in COVID-19 patients have lower levels of a fucosylation within their Fc domains, according to two current studies 208,209, a phenotype linked to higher affinity for FcRIIIa. Increased affinity can be beneficial in some instances due to more robust FcRIIIa-mediated effector responses210,211, while fucosylated non-neutralizing IgG (immunoglobulin-G) antibodies against dengue virus were linked to worse clinical outcomes 212. Larsen et al. also discovered that patients with COVID-19 and acute respiratory distress syndrome had lower S-specific IgG levels than those with asymptomatic or moderate illnesses 209. It's unknown if decreased fucosylation of SARS-CoV-2-specific antibodies caused COVID-19 immunopathology. On the other hand, SARS-CoV-2 (severe acute respiratory syndrome coronavirus-2) has not been shown to infect macrophages in a productive manner 213,214. As a result, the available evidence suggests that the most plausible ADE mechanism related to COVID-19 disease is the creation of antibodies–antigen immune complexes, which results in immune cascade hyperactivation in lung tissue.

Early immunization findings reveal substantial antibody responses by day 14 215, indicating memory B-cell responses with cross-reactivity antibodies from different coronavirus variants (s). SARS 216 and COVID-19 188,217–221, early high antibody responses are associated with higher illness severity. Wu et al. found that antibodies from COVID-19 patients allowed SARS-CoV-2 infections of Raji cells (lymphoma cells derived from B lymphocytes), K562 cells (derived from monocytes), and primary B cells 222. Infection of some phagocytic cells (macrophages) with SARS-CoV-2 may be a significant step in illness development for some patients.

**14.3 Vaccine-associated Antibody-Dependent Enhancement (ADE) risks-**

Virus vaccine manufacturing can be done by using live-disarmed virus strains, inactivated (dead) viruses, protein components, messenger ribonucleic acid (mRNA), and deoxyribonucleic acid (DNA). Vaccines produce antibodies that can fall either in the neutralizing category or non-neutralizing category. We have three known possible processes to support non-neutralizing antibodies in their role towards antiviral activities; they are (CDC), i.e., antibody-mediated complement-dependent cytotoxicity, (ADCC), i.e., antibody-dependent cellular cytotoxicity, and ADCP, i.e., antibody-dependent cellular phagocytosis 223. The annual influenza vaccine is beneficial against the strains of vaccines and other closely linked strains as it provides protection by producing neutralizing and non-neutralizing antibodies.

Vaccine-associated enhanced disease (VAED) can occur when a virus has many circularizing serotypes [e.g., Dengue fever 182,194,195, or when the virus leverages antibodies for increased host cell tropism of phagocytic immune cells. Cell membrane fusion processes are seen in many viruses implicated in ADE 224. Vaccine-induced cross-reactive anti-HA2 antibodies expedite viral fusion in a pig model of influenza A H1N1, resulting in vaccine-associated increased respiratory disease (VAERD) 225. The respiratory syncytial virus (RSV) caused ADE in the Bonnet monkey model 226. To avoid ADE, Van Erp et al. advised against producing respiratory syncytial virus (RSV) non-neutralizing or sub-neutralizing antibodies. ADE has been observed in several SARS-CoV-1 animal models. Attempts to produce SARS-CoV-1 vaccines in a mouse model resulted in pulmonary immunopathology after SARS-CoV-1 problem 227,228; these vaccines included inactivated whole viruses, inactivated viruses with adjuvant and a virus-like particle (VLP) vaccine containing a recombinant DNA spike (S) protein vaccine. Animals immunized with nucleocapsid protein suffered severe pneumonia after exposure to SARS-CoV-1 229.

A vaccine comprising recombinant modified vaccinia virus Ankara (rMVA) expressing the SARS-CoV-1 spike protein exacerbated hepatitis in a ferret model 230. In rhesus macaques, vaccination with SARS-CoV-1 resulted in ADE 176. SARS-CoV-1 ADE is mediated by spike protein antibodies 231. Antibodies to the SARS-CoV-1 spike protein can facilitate viral entry via Fc receptor-exposing cells in a dose-dependent manner 193. Because Fc mediates infection of immune cells, Jaume et al. 232 pointed out the potential downsides of vaccines against SARS-CoV-1 spike protein.

As a result, future attempts to develop SARS-CoV-1, MERS-CoV 233, or SARS-CoV-2 vaccines could increase the danger of causing ADE in people, which is increased by antibody infection of phagocytic immune cells. This possible ADE risk is independent of the vaccination technique 234 or targeting strategy selected due to anticipated phagocytic-resistant cell infections upon antibody absorption. In MERS patients, the rate of seroconversion increased as the disease progressed 235. Severe clinical worsening for SARS patients occurs concurrently with the timing of IgG seroconversion 236. Early high immunoglobulin-G (IgG) responses in SARS patients are linked to illness progression 237 and seriousness 217–221,238. Antibody therapy for severely ill COVID-19 patients has been halted due to a potential safety signal and an inadequate risk-benefit profile 239. With high antibody titers, the existing COVID-19 vaccinations appear to give protection; however, the possibility of ADE risks associated with declining antibody titers over time is unknown.

**14.4 The future management for ADE in coronavirus**

According to preliminary studies, ADE in coronavirus illness can be treated in various methods. The first approach is to keep the dose under control. MERS-CoV's ADE can be blocked with a large antibody dose without compromising the virus's antiviral activity 240. Altering the antibody specificity is the second option. Anti-spike antibodies make it easier to induce ADE, despite the fact that inhibiting the binding of coronavirus spike proteins is a good treatment strategy due to its high efficiency in decreasing viral load. The use of specific inhibitors is the third technique 174. In MERS-CoV) and SARS-CoV, protease inhibitors and Fc inhibitors, for example, have a role in inhibiting ADE 240,241. An adjuvant was observed to increase Th2-type (T helper type 2) immunity and minimize immunopathology in previous SARS-CoV (severe acute respiratory syndrome coronavirus) research, implying the adjuvant's latent importance 242. Furthermore, the dengue virus case can be utilized as a model, with changes to the FcgR binding location on the antibody Fc component lowering the probability of ADE. In these conditions, ensuring that classical viral entry is prevented via antibodies while resolving ADE is another difficulty. Combining Cyclospora A and Chinese medicine pharmaceutics with immunosuppressive qualities with colloidal sub-particles, which can improve macrophage targeting and induce an immunosuppressive effect, could be a viable alternative. This may be effective against viruses and bacteria as well as immune-injury inflammation.

**Conclusion and future perspective**

Coronavirus 2019 (SARS-CoV-2) causes respiratory illness in humans, including vomiting, loss of taste, sneezing, and coughing, while diarrhea and upper respiratory disease occur in animals. According to the WHO guidelines, avoiding contact with infected people and limiting visits to markets or public spaces as much as possible can help break the chain of transmission of coronavirus 2. Essentially, both SARS-CoV and SARS-CoV-2 share common symptoms and exhibit a similar mechanism of action, creating a war-like situation worldwide. Since no specific antiviral medicine or therapy is available, many potential drugs for repurposing to treat COVID-19 patients are being explored worldwide. Such efforts have led to at least managing and controlling the seriousness of illness and death rate of SARS-CoV-2 patients. In parallel, clinical trials of various medicines for repurposing are in progress with due approval from the respective authorities of the country. Also, vaccine development has started taking its shape as many of them are currently undergoing the phases of clinical trials. A few vaccines got approval for usage purposes, such as the AZD1222 vaccine produced by Oxford University and AstraZeneca and Covaxin developed by Bharat Biotech in collaboration with the Indian Council of Medical Research (ICMR). Stem cell-based therapy also shows hope as a potential treatment measure for managing this uncured disease; however, more intensive study of this novel coronavirus is needed to find its permanent cure. ADE has been documented in SARS, MERS, and other human respiratory viral illnesses such as RSV and measles, implying that SARS-CoV-2 vaccinations and antibody-based therapies pose a real risk of ADE. However, Clinical studies have yet to prove ADE's function in human COVID-19 pathology. Animal and human clinical investigations are now underway to learn more about the mechanisms of ADE in SARS-CoV-2. Since December 2019, the world has been coping with various strains of SARS-CoV-2 and the first and second waves of the COVID-19 pandemic. However, a few nations are battling the third wave due to the Delta variety. In the midst of this, the introduction of a new Omicron variety may have a devastating impact on human life and livelihood. The tireless efforts of scientists, medical professionals, front-line workers, and policymakers dealing with this pandemic are commendable. In the previous two years, there has been a significant improvement in our understanding of SARS-CoV-2 and its variations, origin and structure, pathogenesis, and related symptoms in various patient groups. The availability of effective FDA-approved vaccinations, treatment, and management regimes, as well as improved diagnostic and treatment infrastructure and skilled healthcare workers, may aid in better handling of the novel Omicron strain.

**Conflicts of interest**

The authors declare no conflict of interest.

**References**

1. *Coronavirus Disease (COVID-19) Situation Reports*. https://www.who.int/emergencies/diseases/novel-coronavirus-2019/situation-reports (accessed 2020-12-08).
2. Zhu, N.; Zhang, D.; Wang, W.; Li, X.; Yang, B.; Song, J.; Zhao, X.; Huang, B.; Shi, W.; Lu, R.; Niu, P.; Zhan, F.; Ma, X.; Wang, D.; Xu, W.; Wu, G.; Gao, G. F.; Tan, W.; China Novel Coronavirus Investigating and Research Team. A Novel Coronavirus from Patients with Pneumonia in China, 2019. *N. Engl. J. Med.* **2020**, *382* (8), 727–733. https://doi.org/10.1056/NEJMoa2001017.
3. *Coronavirus Update (Live): 72,893,236 Cases and 1,623,425 Deaths from COVID-19 Virus Pandemic - Worldometer*. https://www.worldometers.info/coronavirus/ (accessed 2020-12-14).
4. Golchin, A.; Seyedjafari, E.; Ardeshirylajimi, A. Mesenchymal Stem Cell Therapy for COVID-19: Present or Future. *Stem Cell Rev. Rep.* **2020**, 1–7. https://doi.org/10.1007/s12015-020-09973-w.
5. di Mauro, G.; Scavone, C.; Rafaniello, C.; Rossi, F.; Capuano, A. SARS-Cov-2 Infection: Response of Human Immune System and Possible Implications for the Rapid Test and Treatment. *Int. Immunopharmacol.* **2020**, *84*, 106519. https://doi.org/10.1016/j.intimp.2020.106519.
6. Anderson, R. M.; Heesterbeek, H.; Klinkenberg, D.; Hollingsworth, T. D. How Will Country-Based Mitigation Measures Influence the Course of the COVID-19 Epidemic? *The Lancet* **2020**, *395* (10228), 931–934. https://doi.org/10.1016/S0140-6736(20)30567-5.
7. Shereen, M. A.; Khan, S.; Kazmi, A.; Bashir, N.; Siddique, R. COVID-19 Infection: Origin, Transmission, and Characteristics of Human Coronaviruses. *J. Adv. Res.* **2020**, *24*, 91–98. https://doi.org/10.1016/j.jare.2020.03.005.
8. Harapan, H.; Itoh, N.; Yufika, A.; Winardi, W.; Keam, S.; Te, H.; Megawati, D.; Hayati, Z.; Wagner, A. L.; Mudatsir, M. Coronavirus Disease 2019 (COVID-19): A Literature Review. *J. Infect. Public Health* **2020**, *13* (5), 667–673. https://doi.org/10.1016/j.jiph.2020.03.019.
9. *Fenner and White’s Medical Virology - 5th Edition*. https://www.elsevier.com/books/fenner-and-whites-medical-virology/burrell/978-0-12-375156-0 (accessed 2020-12-14).
10. Mousavizadeh, L.; Ghasemi, S. Genotype and Phenotype of COVID-19: Their Roles in Pathogenesis. *J. Microbiol. Immunol. Infect. Wei Mian Yu Gan Ran Za Zhi* **2020**. https://doi.org/10.1016/j.jmii.2020.03.022.
11. WWeiss, S. R.; Leibowitz, J. L. Coronavirus Pathogenesis. *Adv. Virus Res.* **2011**, *81*, 85–164. https://doi.org/10.1016/B978-0-12-385885-6.00009-2.
12. Hoffmann, M.; Kleine-Weber, H.; Schroeder, S.; Krüger, N.; Herrler, T.; Erichsen, S.; Schiergens, T. S.; Herrler, G.; Wu, N.-H.; Nitsche, A.; Müller, M. A.; Drosten, C.; Pöhlmann, S. SARS-CoV-2 Cell Entry Depends on ACE2 and TMPRSS2 and Is Blocked by a Clinically Proven Protease Inhibitor. *Cell* **2020**, *181* (2), 271-280.e8. https://doi.org/10.1016/j.cell.2020.02.052.
13. Leng, Z.; Zhu, R.; Hou, W.; Feng, Y.; Yang, Y.; Han, Q.; Shan, G.; Meng, F.; Du, D.; Wang, S.; Fan, J.; Wang, W.; Deng, L.; Shi, H.; Li, H.; Hu, Z.; Zhang, F.; Gao, J.; Liu, H.; Li, X.; Zhao, Y.; Yin, K.; He, X.; Gao, Z.; Wang, Y.; Yang, B.; Jin, R.; Stambler, I.; Lim, L. W.; Su, H.; Moskalev, A.; Cano, A.; Chakrabarti, S.; Min, K.-J.; Ellison-Hughes, G.; Caruso, C.; Jin, K.; Zhao, R. C. Transplantation of ACE2- Mesenchymal Stem Cells Improves the Outcome of Patients with COVID-19 Pneumonia. *Aging Dis.* **2020**, *11* (2), 216–228. https://doi.org/10.14336/AD.2020.0228.
14. Callaway, E. Delta Coronavirus Variant: Scientists Brace for Impact. *Nature* **2021**, *595* (7865), 17–18. https://doi.org/10.1038/d41586-021-01696-3.
15. Liu, Y.; Rocklöv, J. The Reproductive Number of the Delta Variant of SARS-CoV-2 Is Far Higher Compared to the Ancestral SARS-CoV-2 Virus. *J. Travel Med.* **2021**, *28* (7), taab124. https://doi.org/10.1093/jtm/taab124.
16. Rambaut, A.; Holmes, E. C.; O’Toole, Á.; Hill, V.; McCrone, J. T.; Ruis, C.; du Plessis, L.; Pybus, O. G. A Dynamic Nomenclature Proposal for SARS-CoV-2 Lineages to Assist Genomic Epidemiology. *Nat. Microbiol.* **2020**, *5* (11), 1403–1407. https://doi.org/10.1038/s41564-020-0770-5.
17. Shang, J.; Wan, Y.; Luo, C.; Ye, G.; Geng, Q.; Auerbach, A.; Li, F. Cell Entry Mechanisms of SARS-CoV-2. *Proc. Natl. Acad. Sci.* **2020**, *117* (21), 11727–11734. https://doi.org/10.1073/pnas.2003138117.
18. Starr, T. N.; Greaney, A. J.; Dingens, A. S.; Bloom, J. D. Complete Map of SARS-CoV-2 RBD Mutations That Escape the Monoclonal Antibody LY-CoV555 and Its Cocktail with LY-CoV016. *BioRxiv Prepr. Serv. Biol.* **2021**, 2021.02.17.431683. https://doi.org/10.1101/2021.02.17.431683.
19. Eichler, N.; Thornley, C.; Swadi, T.; Devine, T.; McElnay, C.; Sherwood, J.; Brunton, C.; Williamson, F.; Freeman, J.; Berger, S.; Ren, X.; Storey, M.; de Ligt, J.; Geoghegan, J. L. Transmission of Severe Acute Respiratory Syndrome Coronavirus 2 during Border Quarantine and Air Travel, New Zealand (Aotearoa). *Emerg. Infect. Dis.* **2021**, *27* (5), 1274–1278. https://doi.org/10.3201/eid2705.210514.
20. He, X.; He, C.; Hong, W.; Zhang, K.; Wei, X. The Challenges of COVID-19 Delta Variant: Prevention and Vaccine Development. *MedComm* **2021**, *2* (4), 846–854. https://doi.org/10.1002/mco2.95.
21. Kenyon, G. Australia’s Struggle with the Delta Variant. *Lancet Infect. Dis.* **2021**, *21* (10), 1358. https://doi.org/10.1016/S1473-3099(21)00579-X.
22. Lopez Bernal, J.; Andrews, N.; Gower, C.; Gallagher, E.; Simmons, R.; Thelwall, S.; Stowe, J.; Tessier, E.; Groves, N.; Dabrera, G.; Myers, R.; Campbell, C. N. J.; Amirthalingam, G.; Edmunds, M.; Zambon, M.; Brown, K. E.; Hopkins, S.; Chand, M.; Ramsay, M. Effectiveness of Covid-19 Vaccines against the B.1.617.2 (Delta) Variant. *N. Engl. J. Med.* **2021**, *385* (7), 585–594. https://doi.org/10.1056/NEJMoa2108891.
23. Mishra, R.; Goel, S. K.; Gupta, K. C.; Kumar, A. Biocomposite Cryogels as Tissue-Engineered Biomaterials for Regeneration of Critical-Sized Cranial Bone Defects. *Tissue Eng. Part A* **2014**, *20* (3–4), 751–762. https://doi.org/10.1089/ten.TEA.2013.0072.
24. Mlcochova, P.; Kemp, S. A.; Dhar, M. S.; Papa, G.; Meng, B.; Ferreira, I. A. T. M.; Datir, R.; Collier, D. A.; Albecka, A.; Singh, S.; Pandey, R.; Brown, J.; Zhou, J.; Goonawardane, N.; Mishra, S.; Whittaker, C.; Mellan, T.; Marwal, R.; Datta, M.; Sengupta, S.; Ponnusamy, K.; Radhakrishnan, V. S.; Abdullahi, A.; Charles, O.; Chattopadhyay, P.; Devi, P.; Caputo, D.; Peacock, T.; Wattal, C.; Goel, N.; Satwik, A.; Vaishya, R.; Agarwal, M.; Mavousian, A.; Lee, J. H.; Bassi, J.; Silacci-Fegni, C.; Saliba, C.; Pinto, D.; Irie, T.; Yoshida, I.; Hamilton, W. L.; Sato, K.; Bhatt, S.; Flaxman, S.; James, L. C.; Corti, D.; Piccoli, L.; Barclay, W. S.; Rakshit, P.; Agrawal, A.; Gupta, R. K. SARS-CoV-2 B.1.617.2 Delta Variant Replication and Immune Evasion. *Nature* **2021**, *599* (7883), 114–119. https://doi.org/10.1038/s41586-021-03944-y.
25. Shiehzadegan, S.; Alaghemand, N.; Fox, M.; Venketaraman, V. Analysis of the Delta Variant B.1.617.2 COVID-19. *Clin. Pract.* **2021**, *11* (4), 778–784. https://doi.org/10.3390/clinpract11040093.
26. Thakur, V.; Ratho, R. K. OMICRON (B.1.1.529): A New SARS-CoV-2 Variant of Concern Mounting Worldwide Fear. *J. Med. Virol.* **2022**, *94* (5), 1821–1824. https://doi.org/10.1002/jmv.27541.
27. Kupferschmidt, K. Where Did “weird” Omicron Come From? *Science* **2021**, *374* (6572), 1179. https://doi.org/10.1126/science.acx9738.
28. Wang, L.; Cheng, G. Sequence Analysis of the Emerging SARS-CoV-2 Variant Omicron in South Africa. *J. Med. Virol.* **2022**, *94* (4), 1728–1733. https://doi.org/10.1002/jmv.27516.
29. Msomi, N.; Lessells, R.; Mlisana, K.; de Oliveira, T. Africa: Tackle HIV and COVID-19 Together. *Nature* **2021**, *600* (7887), 33–36. https://doi.org/10.1038/d41586-021-03546-8.
30. de Wit, E.; van Doremalen, N.; Falzarano, D.; Munster, V. J. SARS, and MERS: Recent Insights into Emerging Coronaviruses. *Nat. Rev. Microbiol.* **2016**, *14* (8), 523–534. https://doi.org/10.1038/nrmicro.2016.81.
31. Lee, N.; Hui, D.; Wu, A.; Chan, P.; Cameron, P.; Joynt, G. M.; Ahuja, A.; Yung, M. Y.; Leung, C. B.; To, K. F.; Lui, S. F.; Szeto, C. C.; Chung, S.; Sung, J. J. Y. A Major Outbreak of Severe Acute Respiratory Syndrome in Hong Kong. *N. Engl. J. Med.* **2003**, *348* (20), 1986–1994. https://doi.org/10.1056/NEJMoa030685.
32. Guan, Y.; Peiris, J.; Zheng, B.; Poon, L.; Chan, K.; Zeng, F.; Chan, C.; Chan, M.; Chen, J.; Chow, K.; Hon, C.; Hui, K.; Li, J.; Li, V.; Wang, Y.; Leung, S.; Yuen, K.; Leung, F. Molecular Epidemiology of the Novel Coronavirus That Causes Severe Acute Respiratory Syndrome. *Lancet Lond. Engl.* **2004**, *363* (9403), 99–104. https://doi.org/10.1016/S0140-6736(03)15259-2.
33. Drosten, C.; Günther, S.; Preiser, W.; van der Werf, S.; Brodt, H.-R.; Becker, S.; Rabenau, H.; Panning, M.; Kolesnikova, L.; Fouchier, R. A. M.; Berger, A.; Burguière, A.-M.; Cinatl, J.; Eickmann, M.; Escriou, N.; Grywna, K.; Kramme, S.; Manuguerra, J.-C.; Müller, S.; Rickerts, V.; Stürmer, M.; Vieth, S.; Klenk, H.-D.; Osterhaus, A. D. M. E.; Schmitz, H.; Doerr, H. W. Identification of a Novel Coronavirus in Patients with Severe Acute Respiratory Syndrome. *N. Engl. J. Med.* **2003**, *348* (20), 1967–1976. https://doi.org/10.1056/NEJMoa030747.
34. Dj, C.; Aj, R.; Jr, V. Severe Acute Respiratory Syndrome (SARS). *Infect. Dis. Clin. North Am.* **2010**, *24* (1), 175–202. https://doi.org/10.1016/j.idc.2009.10.005.
35. Hijawi, B.; Abdallat, M.; Sayaydeh, A.; Alqasrawi, S.; Haddadin, A.; Jaarour, N.; Alsheikh, S.; Alsanouri, T. Novel Coronavirus Infections in Jordan, April 2012: Epidemiological Findings from a Retrospective Investigation. *East. Mediterr. Health J. Rev. Sante Mediterr. Orient. Al-Majallah Al-Sihhiyah Li-Sharq Al-Mutawassit* **2013**, *19 Suppl 1*, S12-18.
36. *WHO | Middle East respiratory syndrome coronavirus (MERS-CoV) – Saudi Arabia*. WHO. http://www.who.int/csr/don/26-april-2016-mers-saudi-arabia/en/ (accessed 2020-12-15).
37. *WHO | Pneumonia of unknown cause – China*. WHO. http://www.who.int/csr/don/05-january-2020-pneumonia-of-unkown-cause-china/en/ (accessed 2020-12-15).
38. Zhang, Y.-Z.; Holmes, E. C. A Genomic Perspective on the Origin and Emergence of SARS-CoV-2. *Cell* **2020**, *181* (2), 223–227. https://doi.org/10.1016/j.cell.2020.03.035.
39. Zhou, P.; Yang, X.-L.; Wang, X.-G.; Hu, B.; Zhang, L.; Zhang, W.; Si, H.-R.; Zhu, Y.; Li, B.; Huang, C.-L.; Chen, H.-D.; Chen, J.; Luo, Y.; Guo, H.; Jiang, R.-D.; Liu, M.-Q.; Chen, Y.; Shen, X.-R.; Wang, X.; Zheng, X.-S.; Zhao, K.; Chen, Q.-J.; Deng, F.; Liu, L.-L.; Yan, B.; Zhan, F.-X.; Wang, Y.-Y.; Xiao, G.-F.; Shi, Z.-L. A Pneumonia Outbreak Associated with a New Coronavirus of Probable Bat Origin. *Nature* **2020**, *579* (7798), 270–273. https://doi.org/10.1038/s41586-020-2012-7.
40. Lu, R.; Zhao, X.; Li, J.; Niu, P.; Yang, B.; Wu, H.; Wang, W.; Song, H.; Huang, B.; Zhu, N.; Bi, Y.; Ma, X.; Zhan, F.; Wang, L.; Hu, T.; Zhou, H.; Hu, Z.; Zhou, W.; Zhao, L.; Chen, J.; Meng, Y.; Wang, J.; Lin, Y.; Yuan, J.; Xie, Z.; Ma, J.; Liu, W. J.; Wang, D.; Xu, W.; Holmes, E. C.; Gao, G. F.; Wu, G.; Chen, W.; Shi, W.; Tan, W. Genomic Characterisation and Epidemiology of 2019 Novel Coronavirus: Implications for Virus Origins and Receptor Binding. *The Lancet* **2020**, *395* (10224), 565–574. https://doi.org/10.1016/S0140-6736(20)30251-8.
41. CDCMMWR. SARS-CoV-2 B.1.1.529 (Omicron) Variant — United States, December 1–8, 2021. *MMWR Morb. Mortal. Wkly. Rep.* **2021**, *70*. https://doi.org/10.15585/mmwr.mm7050e1.
42. Chappell, B. Omicron Spreads Faster than Any Other Variant, WHO Says. It’s Now in 77 Countries. *NPR*. December 15, 2021. https://www.npr.org/sections/coronavirus-live-updates/2021/12/15/1064432010/omicron-spread-variant-coronavirus (accessed 2022-11-19).
43. CDC. *Coronavirus Disease 2019 (COVID-19) – Symptoms*. Centers for Disease Control and Prevention. https://www.cdc.gov/coronavirus/2019-ncov/symptoms-testing/symptoms.html (accessed 2020-12-17).
44. *Question and answers hub*. https://www.who.int/emergencies/diseases/novel-coronavirus-2019/question-and-answers-hub (accessed 2020-12-17).
45. *What Is R0? Gauging Contagious Infections*. Healthline. https://www.healthline.com/health/r-nought-reproduction-number (accessed 2020-12-17).
46. Li, Q.; Guan, X.; Wu, P.; Wang, X.; Zhou, L.; Tong, Y.; Ren, R.; Leung, K. S. M.; Lau, E. H. Y.; Wong, J. Y.; Xing, X.; Xiang, N.; Wu, Y.; Li, C.; Chen, Q.; Li, D.; Liu, T.; Zhao, J.; Liu, M.; Tu, W.; Chen, C.; Jin, L.; Yang, R.; Wang, Q.; Zhou, S.; Wang, R.; Liu, H.; Luo, Y.; Liu, Y.; Shao, G.; Li, H.; Tao, Z.; Yang, Y.; Deng, Z.; Liu, B.; Ma, Z.; Zhang, Y.; Shi, G.; Lam, T. T. Y.; Wu, J. T.; Gao, G. F.; Cowling, B. J.; Yang, B.; Leung, G. M.; Feng, Z. Early Transmission Dynamics in Wuhan, China, of Novel Coronavirus–Infected Pneumonia. *N. Engl. J. Med.* **2020**, *382* (13), 1199–1207. https://doi.org/10.1056/NEJMoa2001316.
47. Wu, J. T.; Leung, K.; Leung, G. M. Nowcasting and Forecasting the Potential Domestic and International Spread of the 2019-NCoV Outbreak Originating in Wuhan, China: A Modelling Study. *The Lancet* **2020**, *395* (10225), 689–697. https://doi.org/10.1016/S0140-6736(20)30260-9.
48. *COVID-19 Projections Using Machine Learning*. COVID-19 Projections Using Machine Learning. https://covid19-projections.com/ (accessed 2020-12-17).
49. He, X.; Lau, E. H. Y.; Wu, P.; Deng, X.; Wang, J.; Hao, X.; Lau, Y. C.; Wong, J. Y.; Guan, Y.; Tan, X.; Mo, X.; Chen, Y.; Liao, B.; Chen, W.; Hu, F.; Zhang, Q.; Zhong, M.; Wu, Y.; Zhao, L.; Zhang, F.; Cowling, B. J.; Li, F.; Leung, G. M. Temporal Dynamics in Viral Shedding and Transmissibility of COVID-19. *Nat. Med.* **2020**, *26* (5), 672–675. https://doi.org/10.1038/s41591-020-0869-5.
50. Gao, Z.; Xu, Y.; Sun, C.; Wang, X.; Guo, Y.; Qiu, S.; Ma, K. A Systematic Review of Asymptomatic Infections with COVID-19. *J. Microbiol. Immunol. Infect.* **2020**. https://doi.org/10.1016/j.jmii.2020.05.001.
51. *50 Percent of People with COVID-19 Aren’t Aware They Have Virus*. Healthline. https://www.healthline.com/health-news/50-percent-of-people-with-covid19-not-aware-have-virus (accessed 2020-12-17).
52. CDC. *Coronavirus Disease 2019 (COVID-19)*. Centers for Disease Control and Prevention. https://www.cdc.gov/coronavirus/2019-ncov/hcp/planning-scenarios.html (accessed 2020-12-17).
53. *COVID-19: Prevention & Investigational Treatments*. Drugs.com. https://www.drugs.com/condition/covid-19.html (accessed 2020-12-17).
54. Chen, J.; Wang, R.; Gilby, N. B.; Wei, G.-W. Omicron (B.1.1.529): Infectivity, Vaccine Breakthrough, and Antibody Resistance. *ArXiv* **2021**, arXiv:2112.01318v1.
55. Pulliam, J. R. C.; van Schalkwyk, C.; Govender, N.; von Gottberg, A.; Cohen, C.; Groome, M. J.; Dushoff, J.; Mlisana, K.; Moultrie, H. Increased Risk of SARS-CoV-2 Reinfection Associated with Emergence of Omicron in South Africa. *Science* **2022**, *376* (6593), eabn4947. https://doi.org/10.1126/science.abn4947.
56. Zhang, L.; Li, Q.; Liang, Z.; Li, T.; Liu, S.; Cui, Q.; Nie, J.; Wu, Q.; Qu, X.; Huang, W.; Wang, Y. The Significant Immune Escape of Pseudotyped SARS-CoV-2 Variant Omicron. *Emerg. Microbes Infect.* **2022**, *11* (1), 1–5. https://doi.org/10.1080/22221751.2021.2017757.
57. Burki, T. K. Omicron Variant and Booster COVID-19 Vaccines. *Lancet Respir. Med.* **2022**, *10* (2), e17. https://doi.org/10.1016/S2213-2600(21)00559-2.
58. Campbell, F.; Archer, B.; Laurenson-Schafer, H.; Jinnai, Y.; Konings, F.; Batra, N.; Pavlin, B.; Vandemaele, K.; Van Kerkhove, M. D.; Jombart, T.; Morgan, O.; le Polain de Waroux, O. Increased Transmissibility and Global Spread of SARS-CoV-2 Variants of Concern as at June 2021. *Euro Surveill. Bull. Eur. Sur Mal. Transm. Eur. Commun. Dis. Bull.* **2021**, *26* (24). https://doi.org/10.2807/1560-7917.ES.2021.26.24.2100509.
59. Liu, X.; Liu, C.; Liu, G.; Luo, W.; Xia, N. COVID-19: Progress in Diagnostics, Therapy and Vaccination. *Theranostics* **2020**, *10* (17), 7821–7835. https://doi.org/10.7150/thno.47987.
60. *Technical guidance publications*. https://www.who.int/emergencies/diseases/novel-coronavirus-2019/technical-guidance-publications (accessed 2020-12-17).
61. Organization, W. H. *Laboratory Testing of 2019 Novel Coronavirus (‎‎‎‎‎‎‎‎‎‎‎2019-NCoV)‎‎‎‎‎‎‎‎‎‎‎ in Suspected Human Cases: Interim Guidance, 17 January 2020*; World Health Organization, 2020.
62. *How is the COVID-19 Virus Detected using Real-Time RT-PCR?* https://www.iaea.org/newscenter/news/how-is-the-covid-19-virus-detected-using-real-time-rt-pcr (accessed 2020-12-17).
63. Bustin, S. A.; Benes, V.; Garson, J. A.; Hellemans, J.; Huggett, J.; Kubista, M.; Mueller, R.; Nolan, T.; Pfaffl, M. W.; Shipley, G. L.; Vandesompele, J.; Wittwer, C. T. The MIQE Guidelines: Minimum Information for Publication of Quantitative Real-Time PCR Experiments. *Clin. Chem.* **2009**, *55* (4), 611–622. https://doi.org/10.1373/clinchem.2008.112797.
64. Corman, V. M.; Landt, O.; Kaiser, M.; Molenkamp, R.; Meijer, A.; Chu, D. K.; Bleicker, T.; Brünink, S.; Schneider, J.; Schmidt, M. L.; Mulders, D. G.; Haagmans, B. L.; van der Veer, B.; van den Brink, S.; Wijsman, L.; Goderski, G.; Romette, J.-L.; Ellis, J.; Zambon, M.; Peiris, M.; Goossens, H.; Reusken, C.; Koopmans, M. P.; Drosten, C. Detection of 2019 Novel Coronavirus (2019-NCoV) by Real-Time RT-PCR. *Eurosurveillance* **2020**, *25* (3). https://doi.org/10.2807/1560-7917.ES.2020.25.3.2000045.
65. *The Basics: RT-PCR - IN*. //www.thermofisher.com/in/en/home/references/ambion-tech-support/rtpcr-analysis/general-articles/rt--pcr-the-basics.html (accessed 2020-12-17).
66. CDC. *Information for Laboratories about Coronavirus (COVID-19)*. Centers for Disease Control and Prevention. https://www.cdc.gov/coronavirus/2019-ncov/lab/serology-testing.html (accessed 2020-12-17).
67. CDC. *COVID-19 and Your Health*. Centers for Disease Control and Prevention. https://www.cdc.gov/coronavirus/2019-ncov/testing/serology-overview.html (accessed 2020-12-17).
68. administrator, J. website. *Global Progress on COVID-19 Serology-Based Testing*. Johns Hopkins Center for Health Security. https://www.centerforhealthsecurity.org/resources/COVID-19/serology/Serology-based-tests-for-COVID-19.html (accessed 2020-12-17).
69. *ICMR-NIV, Pune, develops indigenous ELISA test kit for COVID-19 antibody detection*. http://newsonair.com/Main-News-Details.aspx?id=388679 (accessed 2020-12-23).
70. Infantino, M.; Grossi, V.; Lari, B.; Bambi, R.; Perri, A.; Manneschi, M.; Terenzi, G.; Liotti, I.; Ciotta, G.; Taddei, C.; Benucci, M.; Casprini, P.; Veneziani, F.; Fabbri, S.; Pompetti, A.; Manfredi, M. Diagnostic Accuracy of an Automated Chemiluminescent Immunoassay for Anti-SARS-CoV-2 IgM and IgG Antibodies: An Italian Experience. *J. Med. Virol.* **2020**, *92* (9), 1671–1675. https://doi.org/10.1002/jmv.25932.
71. Revised Guidelines for TrueNat Testing for COVID-19. 1.
72. *Truenat TB Test - Diagnosis, resistance testing, COVID-19*. TBFacts. https://tbfacts.org/truenat/ (accessed 2020-12-18).
73. *Information for COVID-19 Testing Labs*. https://www.icmr.gov.in/ctestlab.html (accessed 2020-12-24).
74. *World Health Organization endorses Truenat tests for initial diagnosis of tuberculosis and detection of rifampicin resistance*. FIND. https://www.finddx.org/newsroom/pr-02jul20/ (accessed 2020-12-23).
75. *World Economic Outlook, April 2020: The Great Lockdown*. IMF. https://www.imf.org/en/Publications/WEO/Issues/2020/04/14/weo-april-2020 (accessed 2020-12-20).
76. *WHO Director-General’s opening remarks at the media briefing on COVID-19 - 20 March 2020*. https://www.who.int/director-general/speeches/detail/who-director-general-s-opening-remarks-at-the-media-briefing-on-covid-19---20-march-2020 (accessed 2020-12-20).
77. *The Coronavirus App*. https://coronavirus.app (accessed 2023-02-01).
78. *Coronavirus in India: Latest Map and Case Count*. https://www.covid19india.org (accessed 2023-02-01).
79. Menon, G. R.; Singh, L.; Sharma, P.; Yadav, P.; Sharma, S.; Kalaskar, S.; Singh, H.; Adinarayanan, S.; Joshua, V.; Kulothungan, V.; Yadav, J.; Watson, L. K.; Fadel, S. A.; Suraweera, W.; Rao, M. V. V.; Dhaliwal, R. S.; Begum, R.; Sati, P.; Jamison, D. T.; Jha, P. National Burden Estimates of Healthy Life Lost in India, 2017: An Analysis Using Direct Mortality Data and Indirect Disability Data. *Lancet Glob. Health* **2019**, *7* (12), e1675–e1684. https://doi.org/10.1016/S2214-109X(19)30451-6.
80. *India - REPORT ON VITAL STATISTICS OF INDIA BASED ON THE CIVIL REGISTRATION SYSTEM-2019*. https://censusindia.gov.in/nada/index.php/catalog/42541 (accessed 2023-02-01).
81. CDC. *COVID Data Tracker*. Centers for Disease Control and Prevention. https://covid.cdc.gov/covid-data-tracker (accessed 2023-02-01).
82. Ahmad, F. B.; Anderson, R. N. The Leading Causes of Death in the US for 2020. *JAMA* **2021**, *325* (18), 1829–1830. https://doi.org/10.1001/jama.2021.5469.
83. (83) *CDC - NCHS - National Center for Health Statistics*. https://www.cdc.gov/nchs/index.htm (accessed 2023-02-01).
84. Velumani, A.; Nikam, C.; Suraweera, W.; Fu, S. H.; Gelband, H.; Brown, P.; Bogoch, I.; Nagelkerke, N.; Jha, P. SARS-CoV-2 Seroprevalence in 12 Cities of India from July-December 2020. *medRxiv* **2021**, 2021.03.19.21253429. https://doi.org/10.1101/2021.03.19.21253429.
85. Pattabiraman, C.; Prasad, P.; George, A. K.; Sreenivas, D.; Rasheed, R.; Reddy, N. V. K.; Desai, A.; Vasanthapuram, R. Importation, Circulation, and Emergence of Variants of SARS-CoV-2 in the South Indian State of Karnataka. *Wellcome Open Res.* **2021**, *6*, 110. https://doi.org/10.12688/wellcomeopenres.16768.2.
86. *Advice for the public on COVID-19 – World Health Organization*. https://www.who.int/emergencies/diseases/novel-coronavirus-2019/advice-for-public (accessed 2020-12-20).
87. *vikaspedia Domains*. https://vikaspedia.in/news/ayurvedas-immunity-boosting-measures-for-self-care-during-covid-19-crisis (accessed 2020-12-20).
88. *Advisory for Coronavirus*. pib.gov.in/Pressreleaseshare.aspx?PRID=1600895 (accessed 2020-12-20).
89. Commissioner, O. of the. Hydroxychloroquine or Chloroquine for COVID-19: Drug Safety Communication - FDA Cautions Against Use Outside of the Hospital Setting or a Clinical Trial Due to Risk of Heart Rhythm Problems. *FDA* **2020**.
90. Yao, X.; Ye, F.; Zhang, M.; Cui, C.; Huang, B.; Niu, P.; Liu, X.; Zhao, L.; Dong, E.; Song, C.; Zhan, S.; Lu, R.; Li, H.; Tan, W.; Liu, D. 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). *Clin. Infect. Dis. Off. Publ. Infect. Dis. Soc. Am.* **2020**, *71* (15), 732–739. https://doi.org/10.1093/cid/ciaa237.
91. Tang, C.; Wang, Y.; Lv, H.; Guan, Z.; Gu, J. Caution against Corticosteroid-Based COVID-19 Treatment. *Lancet Lond. Engl.* **2020**, *395* (10239), 1759–1760. https://doi.org/10.1016/S0140-6736(20)30749-2.
92. *Clinical management of COVID-19*. https://www.who.int/publications-detail-redirect/clinical-management-of-covid-19 (accessed 2020-12-20).
93. Ye, Z.; Wang, Y.; Colunga-Lozano, L. E.; Prasad, M.; Tangamornsuksan, W.; Rochwerg, B.; Yao, L.; Motaghi, S.; Couban, R. J.; Ghadimi, M.; Bala, M. M.; Gomaa, H.; Fang, F.; Xiao, Y.; Guyatt, G. H. Efficacy and Safety of Corticosteroids in COVID-19 Based on Evidence for COVID-19, Other Coronavirus Infections, Influenza, Community-Acquired Pneumonia, and Acute Respiratory Distress Syndrome: A Systematic Review and Meta-Analysis. *CMAJ Can. Med. Assoc. J. J. Assoc. Medicale Can.* **2020**, *192* (27), E756–E767. https://doi.org/10.1503/cmaj.200645.
94. Horby, P.; Lim, W. S.; Emberson, J.; Mafham, M.; Bell, J.; Linsell, L.; Staplin, N.; Brightling, C.; Ustianowski, A.; Elmahi, E.; Prudon, B.; Green, C.; Felton, T.; Chadwick, D.; Rege, K.; Fegan, C.; Chappell, L. C.; Faust, S. N.; Jaki, T.; Jeffery, K.; Montgomery, A.; Rowan, K.; Juszczak, E.; Baillie, J. K.; Haynes, R.; Landray, M. J.; Group, R. C. Effect of Dexamethasone in Hospitalized Patients with COVID-19: Preliminary Report. *medRxiv* **2020**, 2020.06.22.20137273. https://doi.org/10.1101/2020.06.22.20137273.
95. Ni, Y.-N.; Chen, G.; Sun, J.; Liang, B.-M.; Liang, Z.-A. The Effect of Corticosteroids on Mortality of Patients with Influenza Pneumonia: A Systematic Review and Meta-Analysis. *Crit. Care Lond. Engl.* **2019**, *23* (1), 99. https://doi.org/10.1186/s13054-019-2395-8.
96. Huanzhong, S. *Efficacy and Safety of Corticosteroids in COVID-19: A Prospective Randomized Controlled Trails*; Clinical trial registration NCT04273321; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04273321 (accessed 2020-12-19).
97. Wang, Y.; Jiang, W.; He, Q.; Wang, C.; Wang, B.; Zhou, P.; Dong, N.; Tong, Q. A Retrospective Cohort Study of Methylprednisolone Therapy in Severe Patients with COVID-19 Pneumonia. *Signal Transduct. Target. Ther.* **2020**, *5* (1), 57. https://doi.org/10.1038/s41392-020-0158-2.
98. Liu, L.-H.; Zhang, Q.-Y.; Sun, W.; Li, Z.-R.; Gao, F.-Q. Corticosteroid-Induced Osteonecrosis of the Femoral Head: Detection, Diagnosis, and Treatment in Earlier Stages. *Chin. Med. J. (Engl.)* **2017**, *130* (21), 2601–2607. https://doi.org/10.4103/0366-6999.217094.
99. Cao, B.; Wang, Y.; Wen, D.; Liu, W.; Wang, J.; Fan, G.; Ruan, L.; Song, B.; Cai, Y.; Wei, M.; Li, X.; Xia, J.; Chen, N.; Xiang, J.; Yu, T.; Bai, T.; Xie, X.; Zhang, L.; Li, C.; Yuan, Y.; Chen, H.; Li, H.; Huang, H.; Tu, S.; Gong, F.; Liu, Y.; Wei, Y.; Dong, C.; Zhou, F.; Gu, X.; Xu, J.; Liu, Z.; Zhang, Y.; Li, H.; Shang, L.; Wang, K.; Li, K.; Zhou, X.; Dong, X.; Qu, Z.; Lu, S.; Hu, X.; Ruan, S.; Luo, S.; Wu, J.; Peng, L.; Cheng, F.; Pan, L.; Zou, J.; Jia, C.; Wang, J.; Liu, X.; Wang, S.; Wu, X.; Ge, Q.; He, J.; Zhan, H.; Qiu, F.; Guo, L.; Huang, C.; Jaki, T.; Hayden, F. G.; Horby, P. W.; Zhang, D.; Wang, C. A Trial of Lopinavir–Ritonavir in Adults Hospitalized with Severe Covid-19. *N. Engl. J. Med.* **2020**, *382* (19), 1787–1799. https://doi.org/10.1056/NEJMoa2001282.
100. Yamamoto, M.; Kiso, M.; Sakai-Tagawa, Y.; Iwatsuki-Horimoto, K.; Imai, M.; Takeda, M.; Kinoshita, N.; Ohmagari, N.; Gohda, J.; Semba, K.; Matsuda, Z.; Kawaguchi, Y.; Kawaoka, Y.; Inoue, J. The Anticoagulant Nafamostat Potently Inhibits SARS-CoV-2 Infection in Vitro: An Existing Drug with Multiple Possible Therapeutic Effects. *bioRxiv* **2020**, 2020.04.22.054981. https://doi.org/10.1101/2020.04.22.054981.
101. Chupp, G. *The Effect of Camostat Mesylate on COVID-19 Infection in Ambulatory Patients: An Investigator-Initiated Randomized, Placebo-Controlled, Phase IIa Trial*; Clinical trial registration NCT04353284; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04353284 (accessed 2020-12-19).
102. FACC, G. P. R., MD, FAHA. *Randomized Clinical Trial in COvid19 Patients to Assess the Efficacy of the Transmembrane Protease Serine 2 (TMPRSS2) Inhibitor NAfamostat (RACONA Study)*; Clinical trial registration NCT04352400; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04352400 (accessed 2020-12-19).
103. Freedberg, D. E.; Conigliaro, J.; Wang, T. C.; Tracey, K. J.; Callahan, M. V.; Abrams, J. A. Famotidine Use Is Associated With Improved Clinical Outcomes in Hospitalized COVID-19 Patients: A Propensity Score Matched Retrospective Cohort Study. *Gastroenterology* **2020**, *159* (3), 1129-1131.e3. https://doi.org/10.1053/j.gastro.2020.05.053.
104. Malone, R. W.; Tisdall, P.; Fremont-Smith, P.; Liu, Y.; Huang, X.-P.; White, K. M.; Miorin, L.; Olmo, E. M. D.; Alon, A.; Delaforge, E.; Hennecker, C. D.; Wang, G.; Pottel, J.; Smith, N.; Hall, J. M.; Shapiro, G.; Mittermaier, A.; Kruse, A. C.; García-Sastre, A.; Roth, B. L.; Glasspool-Malone, J.; Ricke, D. O. COVID-19: Famotidine, Histamine, Mast Cells, and Mechanisms. *Res. Sq.* **2020**. https://doi.org/10.21203/rs.3.rs-30934/v2.
105. Conigliaro, J. *A Multi-Site, Randomized, Double-Blind, Comparative Trial of the Safety and Efficacy of Standard of Care (SOC) Plus Famotidine vs. SOC Plus Placebo for the Treatment of COVID-19 in Hospitalized Adults*; Clinical trial registration NCT04370262; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04370262 (accessed 2020-12-19).
106. Researcher, S. S. N. I., MD, MPH, MBA, Senior. *Efficacy and Safety of Umifenovir as an Adjuvant Therapy Compared to the Control Therapeutic Regiment of Interferon Beta 1a, Lopinavir / Ritonavir and a Single Dose of Hydroxychloroquine in Moderate to Severe COVID-19: A Randomized, Double-Blind, Placebo-Controlled, Clinical Trial*; Clinical trial registration NCT04350684; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04350684 (accessed 2020-12-19).
107. Leneva, I. A.; Russell, R. J.; Boriskin, Y. S.; Hay, A. J. Characteristics of Arbidol-Resistant Mutants of Influenza Virus: Implications for the Mechanism of Anti-Influenza Action of Arbidol. *Antiviral Res.* **2009**, *81* (2), 132–140. https://doi.org/10.1016/j.antiviral.2008.10.009.
108. Zeron, H. M. *Treatment With Hydroxychloroquine vs Nitazoxanide + Hydroxychloroquine in Patients With COVID-19 With Risk Factors for Poor Outcome*; Clinical trial registration NCT04341493; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04341493 (accessed 2020-12-19).
109. Caly, L.; Druce, J. D.; Catton, M. G.; Jans, D. A.; Wagstaff, K. M. The FDA-Approved Drug Ivermectin Inhibits the Replication of SARS-CoV-2 in Vitro. *Antiviral Res.* **2020**, *178*, 104787. https://doi.org/10.1016/j.antiviral.2020.104787.
110. National Cancer Institute, Naples. *Multicenter Study on the Efficacy and Tolerability of Tocilizumab in the Treatment of Patients With COVID-19 Pneumonia*; Clinical trial registration NCT04317092; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04317092 (accessed 2020-12-19).
111. Fu, B.; Xu, X.; Wei, H. Why Tocilizumab Could Be an Effective Treatment for Severe COVID-19? *J. Transl. Med.* **2020**, *18* (1), 164. https://doi.org/10.1186/s12967-020-02339-3.
112. Regeneron Pharmaceuticals. *An Adaptive Phase 2/3, Randomized, Double-Blind, Placebo-Controlled Study Assessing Efficacy and Safety of Sarilumab for Hospitalized Patients With COVID-19*; Clinical trial registration NCT04315298; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04315298 (accessed 2020-12-19).
113. Xu, X.; Han, M.; Li, T.; Sun, W.; Wang, D.; Fu, B.; Zhou, Y.; Zheng, X.; Yang, Y.; Li, X.; Zhang, X.; Pan, A.; Wei, H. Effective Treatment of Severe COVID-19 Patients with Tocilizumab. *Proc. Natl. Acad. Sci. U. S. A.* **2020**, *117* (20), 10970–10975. https://doi.org/10.1073/pnas.2005615117.
114. Wu, R.; Wang, L.; Kuo, H.-C. D.; Shannar, A.; Peter, R.; Chou, P. J.; Li, S.; Hudlikar, R.; Liu, X.; Liu, Z.; Poiani, G. J.; Amorosa, L.; Brunetti, L.; Kong, A.-N. An Update on Current Therapeutic Drugs Treating COVID-19. *Curr. Pharmacol. Rep.* **2020**, 1–15. https://doi.org/10.1007/s40495-020-00216-7.
115. Zhou, F.; Yu, T.; Du, R.; Fan, G.; Liu, Y.; Liu, Z.; Xiang, J.; Wang, Y.; Song, B.; Gu, X.; Guan, L.; Wei, Y.; Li, H.; Wu, X.; Xu, J.; Tu, S.; Zhang, Y.; Chen, H.; Cao, B. Clinical Course and Risk Factors for Mortality of Adult Inpatients with COVID-19 in Wuhan, China: A Retrospective Cohort Study. *Lancet Lond. Engl.* **2020**, *395* (10229), 1054–1062. https://doi.org/10.1016/S0140-6736(20)30566-3.
116. Qilu Hospital of Shandong University. *The Efficacy and Safety of Bevacizumab in Severe or Critical Patients With COVID-19--a Multicenter Randomized Controlled Clinical Trial*; Clinical trial registration NCT04305106; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04305106 (accessed 2020-12-20).
117. Lenze, E. J.; Mattar, C.; Zorumski, C. F.; Stevens, A.; Schweiger, J.; Nicol, G. E.; Miller, J. P.; Yang, L.; Yingling, M.; Avidan, M. S.; Reiersen, A. M. Fluvoxamine vs Placebo and Clinical Deterioration in Outpatients With Symptomatic COVID-19: A Randomized Clinical Trial. *JAMA* **2020**, *324* (22), 2292–2300. https://doi.org/10.1001/jama.2020.22760.
118. PubChem. *Favipiravir*. https://pubchem.ncbi.nlm.nih.gov/compound/492405 (accessed 2020-12-21).
119. Shiraki, K.; Daikoku, T. Favipiravir, an Anti-Influenza Drug against Life-Threatening RNA Virus Infections. *Pharmacol. Ther.* **2020**, *209*, 107512. https://doi.org/10.1016/j.pharmthera.2020.107512.
120. Ebola Drug From Japan May Emerge Among Key Candidates. *Bloomberg.com*. August 7, 2014. https://www.bloomberg.com/news/articles/2014-08-07/ebola-drug-from-japan-may-emerge-among-key-candidates (accessed 2020-12-21).
121. Ltd, G. P. *Glenmark Becomes the First Pharmaceutical Company in India to Receive Regulatory Approval for Oral Antiviral Favipiravir for the Treatment of Mild to Moderate COVID-19*. https://www.prnewswire.com/in/news-releases/glenmark-becomes-the-first-pharmaceutical-company-in-india-to-receive-regulatory-approval-for-oral-antiviral-favipiravir-for-the-treatment-of-mild-to-moderate-covid-19-855346546.html (accessed 2020-12-21).
122. Chen, C.; Zhang, Y.; Huang, J.; Yin, P.; Cheng, Z.; Wu, J.; Chen, S.; Zhang, Y.; Chen, B.; Lu, M.; Luo, Y.; Ju, L.; Zhang, J.; Wang, X. Favipiravir versus Arbidol for COVID-19: A Randomized Clinical Trial. *medRxiv* **2020**, 2020.03.17.20037432. https://doi.org/10.1101/2020.03.17.20037432.
123. Metcalfe, S. M. Mesenchymal Stem Cells and Management of COVID-19 Pneumonia. *Med. Drug Discov.* **2020**, *5*, 100019. https://doi.org/10.1016/j.medidd.2020.100019.
124. Qingsong, Y. *Safety and Efficacy Study of Allogeneic Human Dental Pulp Mesenchymal Stem Cells to Treat Severe Pneumonia of COVID-19：a Single-Center, Prospective, Randomised Clinical Trial*; Clinical trial registration NCT04336254; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04336254 (accessed 2021-01-11).
125. Andalusian Network for Design and Translation of Advanced Therapies. *Phase I / II Clinical Trial, Multicenter, Randomized, and Controlled, to Assess the Safety and Efficacy of Intravenous Administration of Allogeneic Adult Mesenchymal Stem Cells of Expanded Adipose Tissue in Patients With Severe Pneumonia Due to COVID-19*; Clinical trial registration NCT04366323; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04366323 (accessed 2021-01-11).
126. Royan Institute. *Mesenchymal Stem Cell Therapy for Acute Respiratory Distress Syndrome in Coronavirus Infection: A Phase 2-3 Clinical Trial*; Clinical trial registration NCT04366063; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04366063 (accessed 2021-01-13).
127. SBÜ Dr. Sadi Konuk Eğitim ve Araştırma Hastanesi. *What Is the Effect of Mesenchymal Stem Cell Therapy on Seriously Ill Patients With Covid 19 in Intensive Care? (Prospective Double Controlled Study)*; Clinical trial registration NCT04392778; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04392778 (accessed 2021-01-13).
128. Ricordi, C. *Umbilical Cord-Derived Mesenchymal Stem Cells for COVID-19 Patients With Acute Respiratory Distress Syndrome (ARDS)*; Clinical trial registration NCT04355728; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04355728 (accessed 2021-01-13).
129. McAuley, P. D. *Repair of Acute Respiratory Distress Syndrome by Stromal Cell Administration (REALIST): An Open Label Dose Escalation Phase 1 Trial Followed by a Randomized, Double-Blind, Placebo-Controlled Phase 2 Trial (COVID-19)*; Clinical trial registration NCT03042143; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT03042143 (accessed 2021-01-13).
130. Wang, F.-S. *A Phase II, Multicenter, Randomized, Double-Blind, Placebo-Controlled Trial to Evaluate the Efficacy and Safety of Human Umbilical Cord-Derived Mesenchymal Stem Cells in the Treatment of Severe COVID-19 Patients*; Clinical trial registration NCT04288102; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04288102 (accessed 2021-01-13).
131. Chen, J.; Hu, C.; Chen, L.; Tang, L.; Zhu, Y.; Xu, X.; Chen, L.; Gao, H.; Lu, X.; Yu, L.; Dai, X.; Xiang, C.; Li, L. Clinical Study of Mesenchymal Stem Cell Treatment for Acute Respiratory Distress Syndrome Induced by Epidemic Influenza A (H7N9) Infection: A Hint for COVID-19 Treatment. *Eng. Beijing China* **2020**, *6* (10), 1153–1161. https://doi.org/10.1016/j.eng.2020.02.006.
132. Liang, B.; Chen, J.; Li, T.; Wu, H.; Yang, W.; Li, Y.; Li, J.; Yu, C.; Nie, F.; Ma, Z.; Yang, M.; Xiao, M.; Nie, P.; Gao, Y.; Qian, C.; Hu, M. Clinical Remission of a Critically Ill COVID-19 Patient Treated by Human Umbilical Cord Mesenchymal Stem Cells. *Medicine (Baltimore)* **2020**, *99* (31). https://doi.org/10.1097/MD.0000000000021429.
133. Liu, S.; Peng, D.; Qiu, H.; Yang, K.; Fu, Z.; Zou, L. Mesenchymal Stem Cells as a Potential Therapy for COVID-19. *Stem Cell Res. Ther.* **2020**, *11* (1), 169. https://doi.org/10.1186/s13287-020-01678-8.
134. National Institute of Allergy and Infectious Diseases (NIAID). *Phase I, Open-Label, Dose-Ranging Study of the Safety and Immunogenicity of 2019-NCoV Vaccine (MRNA-1273) in Healthy Adults*; Clinical trial registration NCT04283461; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04283461 (accessed 2020-12-20).
135. *Moderna’s Work on a COVID-19 Vaccine Candidate | Moderna, Inc.* https://www.modernatx.com/modernas-work-potential-vaccine-against-covid-19 (accessed 2021-01-06).
136. ModernaTX, Inc. *A Phase 3, Randomized, Stratified, Observer-Blind, Placebo-Controlled Study to Evaluate the Efficacy, Safety, and Immunogenicity of MRNA-1273 SARS-CoV-2 Vaccine in Adults Aged 18 Years and Older*; Clinical trial registration NCT04470427; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04470427 (accessed 2021-01-04).
137. *AZD1222 SARS-CoV-2 Vaccine*. https://www.precisionvaccinations.com/vaccines/azd1222-sars-cov-2-vaccine (accessed 2020-12-22).
138. Voysey, M.; Clemens, S. A. C.; Madhi, S. A.; Weckx, L. Y.; Folegatti, P. M.; Aley, P. K.; Angus, B.; Baillie, V. L.; Barnabas, S. L.; Bhorat, Q. E.; Bibi, S.; Briner, C.; Cicconi, P.; Collins, A. M.; Colin-Jones, R.; Cutland, C. L.; Darton, T. C.; Dheda, K.; Duncan, C. J. A.; Emary, K. R. W.; Ewer, K. J.; Fairlie, L.; Faust, S. N.; Feng, S.; Ferreira, D. M.; Finn, A.; Goodman, A. L.; Green, C. M.; Green, C. A.; Heath, P. T.; Hill, C.; Hill, H.; Hirsch, I.; Hodgson, S. H. C.; Izu, A.; Jackson, S.; Jenkin, D.; Joe, C. C. D.; Kerridge, S.; Koen, A.; Kwatra, G.; Lazarus, R.; Lawrie, A. M.; Lelliott, A.; Libri, V.; Lillie, P. J.; Mallory, R.; Mendes, A. V. A.; Milan, E. P.; Minassian, A. M.; McGregor, A.; Morrison, H.; Mujadidi, Y. F.; Nana, A.; O’Reilly, P. J.; Padayachee, S. D.; Pittella, A.; Plested, E.; Pollock, K. M.; Ramasamy, M. N.; Rhead, S.; Schwarzbold, A. V.; Singh, N.; Smith, A.; Song, R.; Snape, M. D.; Sprinz, E.; Sutherland, R. K.; Tarrant, R.; Thomson, E. C.; Török, M. E.; Toshner, M.; Turner, D. P. J.; Vekemans, J.; Villafana, T. L.; Watson, M. E. E.; Williams, C. J.; Douglas, A. D.; Hill, A. V. S.; Lambe, T.; Gilbert, S. C.; Pollard, A. J.; Oxford COVID Vaccine Trial Group. Safety and Efficacy of the ChAdOx1 NCoV-19 Vaccine (AZD1222) against SARS-CoV-2: An Interim Analysis of Four Randomised Controlled Trials in Brazil, South Africa, and the UK. *Lancet Lond. Engl.* **2020**. https://doi.org/10.1016/S0140-6736(20)32661-1.
139. Zhu, F.-C.; Li, Y.-H.; Guan, X.-H.; Hou, L.-H.; Wang, W.-J.; Li, J.-X.; Wu, S.-P.; Wang, B.-S.; Wang, Z.; Wang, L.; Jia, S.-Y.; Jiang, H.-D.; Wang, L.; Jiang, T.; Hu, Y.; Gou, J.-B.; Xu, S.-B.; Xu, J.-J.; Wang, X.-W.; Wang, W.; Chen, W. 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. *Lancet Lond. Engl.* **2020**, *395* (10240), 1845–1854. https://doi.org/10.1016/S0140-6736(20)31208-3.
140. CanSino Biologics Inc. *A Global Multicenter, Randomized, Double-Blind, Placebo -Controlled, Adaptive Designed Phase Ⅲ Clinical Trial to Evaluate the Efficacy, Safety, and Immunogenicity of Ad5-NCoV in Adults 18 Years of Age and Older*; Clinical trial registration NCT04526990; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04526990 (accessed 2021-01-04).
141. Modjarrad, K.; Roberts, C. C.; Mills, K. T.; Castellano, A. R.; Paolino, K.; Muthumani, K.; Reuschel, E. L.; Robb, M. L.; Racine, T.; Oh, M.; Lamarre, C.; Zaidi, F. I.; Boyer, J.; Kudchodkar, S. B.; Jeong, M.; Darden, J. M.; Park, Y. K.; Scott, P. T.; Remigio, C.; Parikh, A. P.; Wise, M. C.; Patel, A.; Duperret, E. K.; Kim, K. Y.; Choi, H.; White, S.; Bagarazzi, M.; May, J. M.; Kane, D.; Lee, H.; Kobinger, G.; Michael, N. L.; Weiner, D. B.; Thomas, S. J.; Maslow, J. N. Safety and Immunogenicity of an Anti-Middle East Respiratory Syndrome Coronavirus DNA Vaccine: A Phase 1, Open-Label, Single-Arm, Dose-Escalation Trial. *Lancet Infect. Dis.* **2019**, *19* (9), 1013–1022. https://doi.org/10.1016/S1473-3099(19)30266-X.
142. Inovio Pharmaceuticals. *Phase 1 Open-Label Study to Evaluate the Safety, Tolerability, and Immunogenicity of INO-4800, a Prophylactic Vaccine Against SARS-CoV-2, Administered Intradermally Followed by Electroporation in Healthy Volunteers*; Clinical trial registration NCT04336410; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04336410 (accessed 2020-12-20).
143. Inovio Pharmaceuticals. *Phase 2/3 Randomized, Blinded, Placebo-Controlled Trial to Evaluate the Safety, Immunogenicity, and Efficacy of INO-4800, a Prophylactic Vaccine Against COVID-19 Disease, Administered Intradermally Followed by Electroporation in Healthy Seronegative Adults at High Risk of SARS-CoV-2 Exposure*; Clinical trial registration NCT04642638; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04642638 (accessed 2021-01-04).
144. *Novavax Initiates Phase 1/2 Clinical Trial of COVID-19 Vaccine | Novavax Inc. - IR Site*. https://ir.novavax.com/news-releases/news-release-details/novavax-initiates-phase-12-clinical-trial-covid-19-vaccine (accessed 2020-12-22).
145. Novavax. *A Phase 3, Randomized, Observer-Blinded, Placebo-Controlled Study to Evaluate the Efficacy, Safety, and Immunogenicity of a SARS-CoV-2 Recombinant Spike Protein Nanoparticle Vaccine (SARS-CoV-2 RS) With Matrix-M1TM Adjuvant in Adult Participants ≥ 18 Years*; Clinical trial registration NCT04611802; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04611802 (accessed 2021-01-04).
146. Shenzhen Geno-Immune Medical Institute. *Phase I/II Multicenter Trial of Lentiviral Minigene Vaccine (LV-SMENP) of Covid-19 Coronavirus*; Clinical trial registration NCT04276896; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04276896 (accessed 2020-12-20).
147. Chang, L.-J. *Safety and Immunity Evaluation of A Covid-19 Coronavirus Artificial Antigen Presenting Cell Vaccine*; Clinical trial registration NCT04299724; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04299724 (accessed 2020-12-20).
148. *Our COVID-19 Response*. IAVI. https://www.iavi.org/covid-19 (accessed 2020-12-22).
149. *Participant Enrollment Begins for Phase I Trial of IAVI-Merck COVID-19 Vaccine Candidate*. IAVI. https://www.iavi.org/news-resources/features/participant-enrollment-begins-for-phase-i-trial-of-iavi-merck-covid-19-vaccine-candidate (accessed 2021-01-06).
150. BioNTech RNA Pharmaceuticals GmbH. *A Multi-Site, Phase I/II, 2-Part, Dose-Escalation Trial Investigating the Safety and Immunogenicity of Four Prophylactic SARS-CoV-2 RNA Vaccines Against COVID-2019 Using Different Dosing Regimens in Healthy Adults*; Clinical trial registration NCT04380701; clinicaltrials.gov, 2020. https://clinicaltrials.gov/ct2/show/NCT04380701 (accessed 2020-12-20).
151. *COVAXIN - India’s First Indigenous Covid-19 Vaccine | Bharat Biotech*. https://www.bharatbiotech.com/covaxin.html (accessed 2020-12-22).
152. Florindo, H. F.; Kleiner, R.; Vaskovich-Koubi, D.; Acúrcio, R. C.; Carreira, B.; Yeini, E.; Tiram, G.; Liubomirski, Y.; Satchi-Fainaro, R. Immune-Mediated Approaches against COVID-19. *Nat. Nanotechnol.* **2020**, *15* (8), 630–645. https://doi.org/10.1038/s41565-020-0732-3.
153. Vellingiri, B.; Jayaramayya, K.; Iyer, M.; Narayanasamy, A.; Govindasamy, V.; Giridharan, B.; Ganesan, S.; Venugopal, A.; Venkatesan, D.; Ganesan, H.; Rajagopalan, K.; Rahman, P. K. S. M.; Cho, S.-G.; Kumar, N. S.; Subramaniam, M. D. COVID-19: A Promising Cure for the Global Panic. *Sci. Total Environ.* **2020**, *725*, 138277. https://doi.org/10.1016/j.scitotenv.2020.138277.
154. Azkur, A. K.; Akdis, M.; Azkur, D.; Sokolowska, M.; Veen, W. van de; Brüggen, M.-C.; O’Mahony, L.; Gao, Y.; Nadeau, K.; Akdis, C. A. Immune Response to SARS-CoV-2 and Mechanisms of Immunopathological Changes in COVID-19. *Allergy* **2020**, *75* (7), 1564–1581. https://doi.org/10.1111/all.14364.
155. Wang, F.; Nie, J.; Wang, H.; Zhao, Q.; Xiong, Y.; Deng, L.; Song, S.; Ma, Z.; Mo, P.; Zhang, Y. Characteristics of Peripheral Lymphocyte Subset Alteration in COVID-19 Pneumonia. *J. Infect. Dis.* **2020**, *221* (11), 1762–1769. https://doi.org/10.1093/infdis/jiaa150.
156. Su, R.; Li, Z.; Wang, Y.; Liu, Y.; Zheng, X.; Gao, C.; Li, X.; Wang, C. Imbalance between Th17 and Regulatory T Cells in Patients with Systemic Lupus Erythematosus Combined EBV/CMV Viraemia. *Clin. Exp. Rheumatol.* **2020**, *38* (5), 864–873.
157. Chan, M. H. M.; Wong, V. W. S.; Wong, C. K.; Chan, P. K. S.; Chu, C. M.; Hui, D. S. C.; Suen, M. W. M.; Sung, J. J. Y.; Chung, S. S. C.; Lam, C. W. K. Serum LD1 Isoenzyme and Blood Lymphocyte Subsets as Prognostic Indicators for Severe Acute Respiratory Syndrome. *J. Intern. Med.* **2004**, *255* (4), 512–518. https://doi.org/10.1111/j.1365-2796.2004.01323.x.
158. Qin, C.; Zhou, L.; Hu, Z.; Zhang, S.; Yang, S.; Tao, Y.; Xie, C.; Ma, K.; Shang, K.; Wang, W.; Tian, D.-S. Dysregulation of Immune Response in Patients With Coronavirus 2019 (COVID-19) in Wuhan, China. *Clin. Infect. Dis. Off. Publ. Infect. Dis. Soc. Am.* **2020**, *71* (15), 762–768. https://doi.org/10.1093/cid/ciaa248.
159. Baruah, V.; Bose, S. Immunoinformatics-Aided Identification of T Cell and B Cell Epitopes in the Surface Glycoprotein of 2019-NCoV. *J. Med. Virol.* **2020**, *92* (5), 495–500. https://doi.org/10.1002/jmv.25698.
160. Bhattacharya, M.; Sharma, A. R.; Patra, P.; Ghosh, P.; Sharma, G.; Patra, B. C.; Lee, S.-S.; Chakraborty, C. Development of Epitope-Based Peptide Vaccine against Novel Coronavirus 2019 (SARS-COV-2): Immunoinformatics Approach. *J. Med. Virol.* **2020**, *92* (6), 618–631. https://doi.org/10.1002/jmv.25736.
161. Prompetchara, E.; Ketloy, C.; Palaga, T. Immune Responses in COVID-19 and Potential Vaccines: Lessons Learned from SARS and MERS Epidemic. *Asian Pac. J. Allergy Immunol.* **2020**, *38* (1), 1–9. https://doi.org/10.12932/AP-200220-0772.
162. Kumar, S.; Nyodu, R.; Maurya, V. K.; Saxena, S. K. Host Immune Response and Immunobiology of Human SARS-CoV-2 Infection. In *Coronavirus Disease 2019 (COVID-19): Epidemiology, Pathogenesis, Diagnosis, and Therapeutics*; Saxena, S. K., Ed.; Medical Virology: From Pathogenesis to Disease Control; Springer: Singapore, 2020; pp 43–53. https://doi.org/10.1007/978-981-15-4814-7\_5.
163. Jansen, J. M.; Gerlach, T.; Elbahesh, H.; Rimmelzwaan, G. F.; Saletti, G. Influenza Virus-Specific CD4+, and CD8+ T Cell-Mediated Immunity Induced by Infection and Vaccination. *J. Clin. Virol. Off. Publ. Pan Am. Soc. Clin. Virol.* **2019**, *119*, 44–52. https://doi.org/10.1016/j.jcv.2019.08.009.
164. Yaqinuddin, A. Cross-Immunity between Respiratory Coronaviruses May Limit COVID-19 Fatalities. *Med. Hypotheses* **2020**, *144*, 110049. https://doi.org/10.1016/j.mehy.2020.110049.
165. *Immune responses and immunity to SARS-CoV-2*. European Centre for Disease Prevention and Control. https://www.ecdc.europa.eu/en/covid-19/latest-evidence/immune-responses (accessed 2020-12-22).
166. Diao, B.; Wang, C.; Tan, Y.; Chen, X.; Liu, Y.; Ning, L.; Chen, L.; Li, M.; Liu, Y.; Wang, G.; Yuan, Z.; Feng, Z.; Zhang, Y.; Wu, Y.; Chen, Y. Reduction and Functional Exhaustion of T Cells in Patients With Coronavirus Disease 2019 (COVID-19). *Front. Immunol.* **2020**, *11*. https://doi.org/10.3389/fimmu.2020.00827.
167. Shurin, M. R.; Morris, A.; Wells, A.; Wheeler, S. E. Assessing Immune Response to SARS-CoV-2 Infection. *ImmunoTargets Ther.* **2020**, *9*, 111–114. https://doi.org/10.2147/ITT.S264138.
168. Schmidt, M. E.; Varga, S. M. The CD8 T Cell Response to Respiratory Virus Infections. *Front. Immunol.* **2018**, *9*. https://doi.org/10.3389/fimmu.2018.00678.
169. Grifoni, A.; Weiskopf, D.; Ramirez, S. I.; Mateus, J.; Dan, J. M.; Moderbacher, C. R.; Rawlings, S. A.; Sutherland, A.; Premkumar, L.; Jadi, R. S.; Marrama, D.; de Silva, A. M.; Frazier, A.; Carlin, A. F.; Greenbaum, J. A.; Peters, B.; Krammer, F.; Smith, D. M.; Crotty, S.; Sette, A. Targets of T Cell Responses to SARS-CoV-2 Coronavirus in Humans with COVID-19 Disease and Unexposed Individuals. *Cell* **2020**, *181* (7), 1489-1501.e15. https://doi.org/10.1016/j.cell.2020.05.015.
170. Mathew, D.; Giles, J. R.; Baxter, A. E.; Oldridge, D. A.; Greenplate, A. R.; Wu, J. E.; Alanio, C.; Kuri-Cervantes, L.; Pampena, M. B.; D’Andrea, K.; Manne, S.; Chen, Z.; Huang, Y. J.; Reilly, J. P.; Weisman, A. R.; Ittner, C. A. G.; Kuthuru, O.; Dougherty, J.; Nzingha, K.; Han, N.; Kim, J.; Pattekar, A.; Goodwin, E. C.; Anderson, E. M.; Weirick, M. E.; Gouma, S.; Arevalo, C. P.; Bolton, M. J.; Chen, F.; Lacey, S. F.; Ramage, H.; Cherry, S.; Hensley, S. E.; Apostolidis, S. A.; Huang, A. C.; Vella, L. A.; UPenn COVID Processing Unit; Betts, M. R.; Meyer, N. J.; Wherry, E. J. Deep Immune Profiling of COVID-19 Patients Reveals Distinct Immunotypes with Therapeutic Implications. *Science* **2020**, *369* (6508). https://doi.org/10.1126/science.abc8511.
171. van de Veen, W.; Akdis, M. Tolerance Mechanisms of Allergen Immunotherapy. *Allergy* **2020**, *75* (5), 1017–1018. https://doi.org/10.1111/all.14126.
172. Long, Q.-X.; Liu, B.-Z.; Deng, H.-J.; Wu, G.-C.; Deng, K.; Chen, Y.-K.; Liao, P.; Qiu, J.-F.; Lin, Y.; Cai, X.-F.; Wang, D.-Q.; Hu, Y.; Ren, J.-H.; Tang, N.; Xu, Y.-Y.; Yu, L.-H.; Mo, Z.; Gong, F.; Zhang, X.-L.; Tian, W.-G.; Hu, L.; Zhang, X.-X.; Xiang, J.-L.; Du, H.-X.; Liu, H.-W.; Lang, C.-H.; Luo, X.-H.; Wu, S.-B.; Cui, X.-P.; Zhou, Z.; Zhu, M.-M.; Wang, J.; Xue, C.-J.; Li, X.-F.; Wang, L.; Li, Z.-J.; Wang, K.; Niu, C.-C.; Yang, Q.-J.; Tang, X.-J.; Zhang, Y.; Liu, X.-M.; Li, J.-J.; Zhang, D.-C.; Zhang, F.; Liu, P.; Yuan, J.; Li, Q.; Hu, J.-L.; Chen, J.; Huang, A.-L. Antibody Responses to SARS-CoV-2 in Patients with COVID-19. *Nat. Med.* **2020**, *26* (6), 845–848. https://doi.org/10.1038/s41591-020-0897-1.
173. Lee, Y.-L.; Liao, C.-H.; Liu, P.-Y.; Cheng, C.-Y.; Chung, M.-Y.; Liu, C.-E.; Chang, S.-Y.; Hsueh, P.-R. Dynamics of Anti-SARS-Cov-2 IgM and IgG Antibodies among COVID-19 Patients. *J. Infect.* **2020**, *81* (2), e55–e58. https://doi.org/10.1016/j.jinf.2020.04.019.
174. Wen, J.; Cheng, Y.; Ling, R.; Dai, Y.; Huang, B.; Huang, W.; Zhang, S.; Jiang, Y. Antibody-Dependent Enhancement of Coronavirus. *Int. J. Infect. Dis. IJID Off. Publ. Int. Soc. Infect. Dis.* **2020**, *100*, 483–489. https://doi.org/10.1016/j.ijid.2020.09.015.
175. Taylor, A.; Foo, S.-S.; Bruzzone, R.; Dinh, L. V.; King, N. J. C.; Mahalingam, S. Fc Receptors in Antibody-Dependent Enhancement of Viral Infections. *Immunol. Rev.* **2015**, *268* (1), 340–364. https://doi.org/10.1111/imr.12367.
176. Wang, Q.; Zhang, L.; Kuwahara, K.; Li, L.; Liu, Z.; Li, T.; Zhu, H.; Liu, J.; Xu, Y.; Xie, J.; Morioka, H.; Sakaguchi, N.; Qin, C.; Liu, G. Immunodominant SARS Coronavirus Epitopes in Humans Elicited Both Enhancing and Neutralizing Effects on Infection in Non-Human Primates. *ACS Infect. Dis.* **2016**, *2* (5), 361–376. https://doi.org/10.1021/acsinfecdis.6b00006.
177. Lee, W. S.; Wheatley, A. K.; Kent, S. J.; DeKosky, B. J. Antibody-Dependent Enhancement and SARS-CoV-2 Vaccines and Therapies. *Nat. Microbiol.* **2020**, *5* (10), 1185–1191. https://doi.org/10.1038/s41564-020-00789-5.
178. Graham, B. S. Vaccines against Respiratory Syncytial Virus: The Time Has Finally Come. *Vaccine* **2016**, *34* (30), 3535–3541. https://doi.org/10.1016/j.vaccine.2016.04.083.
179. Kim, H. W.; Canchola, J. G.; Brandt, C. D.; Pyles, G.; Chanock, R. M.; Jensen, K.; Parrott, R. H. Respiratory Syncytial Virus Disease in Infants despite Prior Administration of Antigenic Inactivated Vaccine. *Am. J. Epidemiol.* **1969**, *89* (4), 422–434. https://doi.org/10.1093/oxfordjournals.aje.a120955.
180. Nader, P. R.; Horwitz, M. S.; Rousseau, J. Atypical Exanthem Following Exposure to Natural Measles. Eleven Cases in Children Previously Inoculated with Killed Vaccine. *J. Pediatr.* **1968**, *72* (1), 22–28.
181. Polack, F. P. Atypical Measles and Enhanced Respiratory Syncytial Virus Disease (ERD) Made Simple. *Pediatr. Res.* **2007**, *62* (1), 111–115. https://doi.org/10.1203/PDR.0b013e3180686ce0.
182. Dejnirattisai, W.; Jumnainsong, A.; Onsirisakul, N.; Fitton, P.; Vasanawathana, S.; Limpitikul, W.; Puttikhunt, C.; Edwards, C.; Duangchinda, T.; Supasa, S.; Chawansuntati, K.; Malasit, P.; Mongkolsapaya, J.; Screaton, G. Cross-Reacting Antibodies Enhance Dengue Virus Infection in Humans. *Science* **2010**, *328* (5979), 745–748. https://doi.org/10.1126/science.1185181.
183. Sridhar, S.; Luedtke, A.; Langevin, E.; Zhu, M.; Bonaparte, M.; Machabert, T.; Savarino, S.; Zambrano, B.; Moureau, A.; Khromava, A.; Moodie, Z.; Westling, T.; Mascareñas, C.; Frago, C.; Cortés, M.; Chansinghakul, D.; Noriega, F.; Bouckenooghe, A.; Chen, J.; Ng, S.-P.; Gilbert, P. B.; Gurunathan, S.; DiazGranados, C. A. Effect of Dengue Serostatus on Dengue Vaccine Safety and Efficacy. *N. Engl. J. Med.* **2018**, *379* (4), 327–340. https://doi.org/10.1056/NEJMoa1800820.
184. Hohdatsu, T.; Yamada, M.; Tominaga, R.; Makino, K.; Kida, K.; Koyama, H. Antibody-Dependent Enhancement of Feline Infectious Peritonitis Virus Infection in Feline Alveolar Macrophages and Human Monocyte Cell Line U937 by Serum of Cats Experimentally or Naturally Infected with Feline Coronavirus. *J. Vet. Med. Sci.* **1998**, *60* (1), 49–55. https://doi.org/10.1292/jvms.60.49.
185. Ho, M.-S.; Chen, W.-J.; Chen, H.-Y.; Lin, S.-F.; Wang, M.-C.; Di, J.; Lu, Y.-T.; Liu, C.-L.; Chang, S.-C.; Chao, C.-L.; King, C.-C.; Chiou, J.-M.; Su, I.-J.; Yang, J.-Y. Neutralizing Antibody Response and SARS Severity. *Emerg. Infect. Dis.* **2005**, *11* (11), 1730–1737. https://doi.org/10.3201/eid1111.040659.
186. Zhao, J.; Yuan, Q.; Wang, H.; Liu, W.; Liao, X.; Su, Y.; Wang, X.; Yuan, J.; Li, T.; Li, J.; Qian, S.; Hong, C.; Wang, F.; Liu, Y.; Wang, Z.; He, Q.; Li, Z.; He, B.; Zhang, T.; Fu, Y.; Ge, S.; Liu, L.; Zhang, J.; Xia, N.; Zhang, Z. Antibody Responses to SARS-CoV-2 in Patients With Novel Coronavirus Disease 2019. *Clin. Infect. Dis. Off. Publ. Infect. Dis. Soc. Am.* **2020**, *71* (16), 2027–2034. https://doi.org/10.1093/cid/ciaa344.
187. Liu, Y.; Yan, L.-M.; Wan, L.; Xiang, T.-X.; Le, A.; Liu, J.-M.; Peiris, M.; Poon, L. L. M.; Zhang, W. Viral Dynamics in Mild and Severe Cases of COVID-19. *Lancet Infect. Dis.* **2020**, *20* (6), 656–657. https://doi.org/10.1016/S1473-3099(20)30232-2.
188. Zheng, S.; Fan, J.; Yu, F.; Feng, B.; Lou, B.; Zou, Q.; Xie, G.; Lin, S.; Wang, R.; Yang, X.; Chen, W.; Wang, Q.; Zhang, D.; Liu, Y.; Gong, R.; Ma, Z.; Lu, S.; Xiao, Y.; Gu, Y.; Zhang, J.; Yao, H.; Xu, K.; Lu, X.; Wei, G.; Zhou, J.; Fang, Q.; Cai, H.; Qiu, Y.; Sheng, J.; Chen, Y.; Liang, T. Viral Load Dynamics and Disease Severity in Patients Infected with SARS-CoV-2 in Zhejiang Province, China, January-March 2020: Retrospective Cohort Study. *BMJ* **2020**, *369*, m1443. https://doi.org/10.1136/bmj.m1443.
189. Long, Q.-X.; Tang, X.-J.; Shi, Q.-L.; Li, Q.; Deng, H.-J.; Yuan, J.; Hu, J.-L.; Xu, W.; Zhang, Y.; Lv, F.-J.; Su, K.; Zhang, F.; Gong, J.; Wu, B.; Liu, X.-M.; Li, J.-J.; Qiu, J.-F.; Chen, J.; Huang, A.-L. Clinical and Immunological Assessment of Asymptomatic SARS-CoV-2 Infections. *Nat. Med.* **2020**, *26* (8), 1200–1204. https://doi.org/10.1038/s41591-020-0965-6.
190. Sekine, T.; Perez-Potti, A.; Rivera-Ballesteros, O.; Strålin, K.; Gorin, J.-B.; Olsson, A.; Llewellyn-Lacey, S.; Kamal, H.; Bogdanovic, G.; Muschiol, S.; Wullimann, D. J.; Kammann, T.; Emgård, J.; Parrot, T.; Folkesson, E.; Karolinska COVID-19 Study Group; Rooyackers, O.; Eriksson, L. I.; Henter, J.-I.; Sönnerborg, A.; Allander, T.; Albert, J.; Nielsen, M.; Klingström, J.; Gredmark-Russ, S.; Björkström, N. K.; Sandberg, J. K.; Price, D. A.; Ljunggren, H.-G.; Aleman, S.; Buggert, M. Robust T Cell Immunity in Convalescent Individuals with Asymptomatic or Mild COVID-19. *Cell* **2020**, *183* (1), 158-168.e14. https://doi.org/10.1016/j.cell.2020.08.017.
191. Tetro, J. A. Is COVID-19 Receiving ADE from Other Coronaviruses? *Microbes Infect.* **2020**, *22* (2), 72–73. https://doi.org/10.1016/j.micinf.2020.02.006.
192. Khan, S.; Nakajima, R.; Jain, A.; de Assis, R. R.; Jasinskas, A.; Obiero, J. M.; Adenaiye, O.; Tai, S.; Hong, F.; Milton, D. K.; Davies, H.; Felgner, P. L. Analysis of Serologic Cross-Reactivity Between Common Human Coronaviruses and SARS-CoV-2 Using Coronavirus Antigen Microarray. *bioRxiv* **2020**, 2020.03.24.006544. https://doi.org/10.1101/2020.03.24.006544.
193. Wan, Y.; Shang, J.; Sun, S.; Tai, W.; Chen, J.; Geng, Q.; He, L.; Chen, Y.; Wu, J.; Shi, Z.; Zhou, Y.; Du, L.; Li, F. Molecular Mechanism for Antibody-Dependent Enhancement of Coronavirus Entry. *J. Virol.* **2020**, *94* (5), e02015-19. https://doi.org/10.1128/JVI.02015-19.
194. Guzman, M. G.; Alvarez, M.; Rodriguez-Roche, R.; Bernardo, L.; Montes, T.; Vazquez, S.; Morier, L.; Alvarez, A.; Gould, E. A.; Kourí, G.; Halstead, S. B. Neutralizing Antibodies after Infection with Dengue 1 Virus. *Emerg. Infect. Dis.* **2007**, *13* (2), 282–286. https://doi.org/10.3201/eid1302.060539.
195. Katzelnick, L. C.; Gresh, L.; Halloran, M. E.; Mercado, J. C.; Kuan, G.; Gordon, A.; Balmaseda, A.; Harris, E. Antibody-Dependent Enhancement of Severe Dengue Disease in Humans. *Science* **2017**, *358* (6365), 929–932. https://doi.org/10.1126/science.aan6836.
196. Yeh, C.-S.; Yang, J.-Y.; Liu, W.-T.; Huang, J. C.; Chen, Y.-M. A.; Wang, S.-F. SARS Coronavirus Has Antibody-Dependent Enhancement (ADE) Effect through the Autologous Antibodies against Envelope Spikes on Fcγ Receptor Expressing Cells. *J. Virus Erad.* **2016**, *2*, 48. https://doi.org/10.1016/S2055-6640(20)31216-4.
197. Tillett, R. L.; Sevinsky, J. R.; Hartley, P. D.; Kerwin, H.; Crawford, N.; Gorzalski, A.; Laverdure, C.; Verma, S. C.; Rossetto, C. C.; Jackson, D.; Farrell, M. J.; Hooser, S. V.; Pandori, M. Genomic Evidence for Reinfection with SARS-CoV-2: A Case Study. *Lancet Infect. Dis.* **2021**, *21* (1), 52–58. https://doi.org/10.1016/S1473-3099(20)30764-7.
198. Halstead, S. B.; O’Rourke, E. J. Dengue Viruses and Mononuclear Phagocytes. I. Infection Enhancement by Non-Neutralizing Antibody. *J. Exp. Med.* **1977**, *146* (1), 201–217. https://doi.org/10.1084/jem.146.1.201.
199. Vennema, H.; de Groot, R. J.; Harbour, D. A.; Dalderup, M.; Gruffydd-Jones, T.; Horzinek, M. C.; Spaan, W. J. Early Death after Feline Infectious Peritonitis Virus Challenge Due to Recombinant Vaccinia Virus Immunization. *J. Virol.* **1990**, *64* (3), 1407–1409.
200. Weiss, R. C.; Scott, F. W. Antibody-Mediated Enhancement of Disease in Feline Infectious Peritonitis: Comparisons with Dengue Hemorrhagic Fever. *Comp. Immunol. Microbiol. Infect. Dis.* **1981**, *4* (2), 175–189. https://doi.org/10.1016/0147-9571(81)90003-5.
201. Hohdatsu, T.; Nakamura, M.; Ishizuka, Y.; Yamada, H.; Koyama, H. A Study on the Mechanism of Antibody-Dependent Enhancement of Feline Infectious Peritonitis Virus Infection in Feline Macrophages by Monoclonal Antibodies. *Arch. Virol.* **1991**, *120* (3–4), 207–217. https://doi.org/10.1007/BF01310476.
202. Winarski, K. L.; Tang, J.; Klenow, L.; Lee, J.; Coyle, E. M.; Manischewitz, J.; Turner, H. L.; Takeda, K.; Ward, A. B.; Golding, H.; Khurana, S. Antibody-Dependent Enhancement of Influenza Disease Promoted by Increase in Hemagglutinin Stem Flexibility and Virus Fusion Kinetics. *Proc. Natl. Acad. Sci.* **2019**, *116* (30), 15194–15199. https://doi.org/10.1073/pnas.1821317116.
203. Ye, Z.-W.; Yuan, S.; Poon, K.-M.; Wen, L.; Yang, D.; Sun, Z.; Li, C.; Hu, M.; Shuai, H.; Zhou, J.; Zhang, M.-Y.; Zheng, B.-J.; Chu, H.; Yuen, K.-Y. Antibody-Dependent Cell-Mediated Cytotoxicity Epitopes on the Hemagglutinin Head Region of Pandemic H1N1 Influenza Virus Play Detrimental Roles in H1N1-Infected Mice. *Front. Immunol.* **2017**, *8*, 317. https://doi.org/10.3389/fimmu.2017.00317.
204. Polack, F. P.; Teng, M. N.; Collins, P. L.; Prince, G. A.; Exner, M.; Regele, H.; Lirman, D. D.; Rabold, R.; Hoffman, S. J.; Karp, C. L.; Kleeberger, S. R.; Wills-Karp, M.; Karron, R. A. A Role for Immune Complexes in Enhanced Respiratory Syncytial Virus Disease. *J. Exp. Med.* **2002**, *196* (6), 859–865. https://doi.org/10.1084/jem.20020781.
205. Polack, F. P.; Hoffman, S. J.; Crujeiras, G.; Griffin, D. E. A Role for Nonprotective Complement-Fixing Antibodies with Low Avidity for Measles Virus in Atypical Measles. *Nat. Med.* **2003**, *9* (9), 1209–1213. https://doi.org/10.1038/nm918.
206. Gao, T.; Hu, M.; Zhang, X.; Li, H.; Zhu, L.; Liu, H.; Dong, Q.; Zhang, Z.; Wang, Z.; Hu, Y.; Fu, Y.; Jin, Y.; Li, K.; Zhao, S.; Xiao, Y.; Luo, S.; Li, L.; Zhao, L.; Liu, J.; Zhao, H.; Liu, Y.; Yang, W.; Peng, J.; Chen, X.; Li, P.; Liu, Y.; Xie, Y.; Song, J.; Zhang, L.; Ma, Q.; Bian, X.; Chen, W.; Liu, X.; Mao, Q.; Cao, C. Highly Pathogenic Coronavirus N Protein Aggravates Lung Injury by MASP-2-Mediated Complement over-Activatio. 2020. https://doi.org/10.1101/2020.03.29.20041962.
207. Gralinski, L. E.; Sheahan, T. P.; Morrison, T. E.; Menachery, V. D.; Jensen, K.; Leist, S. R.; Whitmore, A.; Heise, M. T.; Baric, R. S. Complement Activation Contributes to Severe Acute Respiratory Syndrome Coronavirus Pathogenesis. *mBio* **2018**, *9* (5), e01753-18. https://doi.org/10.1128/mBio.01753-18.
208. Chakraborty, S.; Edwards, K.; Buzzanco, A. S.; Memoli, M. J.; Sherwood, R.; Mallajosyula, V.; Xie, M. M.; Gonzalez, J.; Buffone, C.; Kathale, N.; Providenza, S.; Jagannathan, P.; Andrews, J. R.; Blish, C. A.; Krammer, F.; Dugan, H.; Wilson, P. C.; Pham, T. D.; Boyd, S. D.; Zhang, S.; Taubenberger, J. K.; Morales, T.; Schapiro, J. M.; Parsonnet, J.; Wang, T. T. Symptomatic SARS-CoV-2 Infections Display Specific IgG Fc Structures. *medRxiv*. https://doi.org/10.1101/2020.05.15.20103341.
209. Larsen, M. D.; Graaf, E. L. de; Sonneveld, M. E.; Plomp, H. R.; Linty, F.; Visser, R.; Brinkhaus, M.; Šuštić, T.; Taeye, S. W. de; Bentlage, A. E. H.; Nouta, J.; Natunen, S.; Koeleman, C. A. M.; Sainio, S.; Kootstra, N. A.; Brouwer, P. J. M.; Sanders, R. W.; Gils, M. J. van; Bruin, S. de; Vlaar, A. P. J.; Group, A. U. C.-19 biobank study; Zaaijer, H. L.; Wuhrer, M.; Schoot, C. E. van der; Vidarsson, G. Afucosylated Immunoglobulin G Responses Are a Hallmark of Enveloped Virus Infections and Show an Exacerbated Phenotype in COVID-19. *bioRxiv* **2020**, 2020.05.18.099507. https://doi.org/10.1101/2020.05.18.099507.
210. Hiatt, A.; Bohorova, N.; Bohorov, O.; Goodman, C.; Kim, D.; Pauly, M. H.; Velasco, J.; Whaley, K. J.; Piedra, P. A.; Gilbert, B. E.; Zeitlin, L. Glycan Variants of a Respiratory Syncytial Virus Antibody with Enhanced Effector Function and in Vivo Efficacy. *Proc. Natl. Acad. Sci.* **2014**, *111* (16), 5992–5997. https://doi.org/10.1073/pnas.1402458111.
211. Zeitlin, L.; Pettitt, J.; Scully, C.; Bohorova, N.; Kim, D.; Pauly, M.; Hiatt, A.; Ngo, L.; Steinkellner, H.; Whaley, K. J.; Olinger, G. G. Enhanced Potency of a Fucose-Free Monoclonal Antibody Being Developed as an Ebola Virus Immunoprotectant. *Proc. Natl. Acad. Sci.* **2011**, *108* (51), 20690–20694. https://doi.org/10.1073/pnas.1108360108.
212. Wang, T. T.; Sewatanon, J.; Memoli, M. J.; Wrammert, J.; Bournazos, S.; Bhaumik, S. K.; Pinsky, B. A.; Chokephaibulkit, K.; Onlamoon, N.; Pattanapanyasat, K.; Taubenberger, J. K.; Ahmed, R.; Ravetch, J. V. IgG Antibodies to Dengue Enhanced for FcγRIIIA Binding Determine Disease Severity. *Science* **2017**, *355* (6323), 395. https://doi.org/10.1126/science.aai8128.
213. Hui, K. P. Y.; Cheung, M.-C.; Perera, R. A. P. M.; Ng, K.-C.; Bui, C. H. T.; Ho, J. C. W.; Ng, M. M. T.; Kuok, D. I. T.; Shih, K. C.; Tsao, S.-W.; Poon, L. L. M.; Peiris, M.; Nicholls, J. M.; Chan, M. C. W. 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. *Lancet Respir Med* **2020**, 687–695.
214. Yip, M. S.; Leung, N. H. L.; Cheung, C. Y.; Li, P. H.; Lee, H. H. Y.; Daëron, M.; Peiris, J. S. M.; Bruzzone, R.; Jaume, M. Antibody-Dependent Infection of Human Macrophages by Severe Acute Respiratory Syndrome Coronavirus. *Virol. J.* **2014**, *11*, 82. https://doi.org/10.1186/1743-422X-11-82.
215. Jackson, L. A.; Anderson, E. J.; Rouphael, N. G.; Roberts, P. C.; Makhene, M.; Coler, R. N.; McCullough, M. P.; Chappell, J. D.; Denison, M. R.; Stevens, L. J.; Pruijssers, A. J.; McDermott, A.; Flach, B.; Doria-Rose, N. A.; Corbett, K. S.; Morabito, K. M.; O’Dell, S.; Schmidt, S. D.; Swanson, P. A.; Padilla, M.; Mascola, J. R.; Neuzil, K. M.; Bennett, H.; Sun, W.; Peters, E.; Makowski, M.; Albert, J.; Cross, K.; Buchanan, W.; Pikaart-Tautges, R.; Ledgerwood, J. E.; Graham, B. S.; Beigel, J. H. An MRNA Vaccine against SARS-CoV-2 — Preliminary Report. *N. Engl. J. Med.* **2020**, NEJMoa2022483. https://doi.org/10.1056/NEJMoa2022483.
216. Lee, N.; Chan, P. K. S.; Ip, M.; Wong, E.; Ho, J.; Ho, C.; Cockram, C. S.; Hui, D. S. Anti-SARS-CoV IgG Response about Disease Severity of Severe Acute Respiratory Syndrome. *J. Clin. Virol. Off. Publ. Pan Am. Soc. Clin. Virol.* **2006**, *35* (2), 179–184. https://doi.org/10.1016/j.jcv.2005.07.005.
217. Chen, W.; Zhang, J.; Qin, X.; Wang, W.; Xu, M.; Wang, L.-F.; Xu, C.; Tang, S.; Liu, P.; Zhang, L.; Liu, X.; Zhang, Y.; Yi, C.; Hu, Z.; Yi, Y. SARS-CoV-2 Neutralizing Antibody Levels Are Correlated with Severity of COVID-19 Pneumonia. *Biomed. Pharmacother.* **2020**, *130*, 110629. https://doi.org/10.1016/j.biopha.2020.110629.
218. Fajnzylber, J.; Regan, J.; Coxen, K.; Corry, H.; Wong, C.; Rosenthal, A.; Worrall, D.; Giguel, F.; Piechocka-Trocha, A.; Atyeo, C.; Fischinger, S.; Chan, A.; Flaherty, K. T.; Hall, K.; Dougan, M.; Ryan, E. T.; Gillespie, E.; Chishti, R.; Li, Y.; Jilg, N.; Hanidziar, D.; Baron, R. M.; Baden, L.; Tsibris, A. M.; Armstrong, K. A.; Kuritzkes, D. R.; Alter, G.; Walker, B. D.; Yu, X.; Li, J. Z. SARS-CoV-2 Viral Load Is Associated with Increased Disease Severity and Mortality. *Nat. Commun.* **2020**, *11* (1), 5493. https://doi.org/10.1038/s41467-020-19057-5.
219. Liu, X.; Wang, J.; Xu, X.; Liao, G.; Chen, Y.; Hu, C.-H. Patterns of IgG and IgM Antibody Response in COVID-19 Patients. *Emerg. Microbes Infect.* **2020**, *9* (1), 1269–1274. https://doi.org/10.1080/22221751.2020.1773324.
220. Luo, Y. R.; Chakraborty, I.; Yun, C.; Wu, A. H. B.; Lynch, K. L. Kinetics of SARS-CoV-2 Antibody Avidity Maturation and Association with Disease Severity. *Clin. Infect. Dis. Off. Publ. Infect. Dis. Soc. Am.* **2020**, ciaa1389. https://doi.org/10.1093/cid/ciaa1389.
221. Pujadas, E.; Chaudhry, F.; McBride, R.; Richter, F.; Zhao, S.; Wajnberg, A.; Nadkarni, G.; Glicksberg, B. S.; Houldsworth, J.; Cordon-Cardo, C. SARS-CoV-2 Viral Load Predicts COVID-19 Mortality. *Lancet Respir. Med.* **2020**, *8* (9), e70. https://doi.org/10.1016/S2213-2600(20)30354-4.
222. Wu, F.; Yan, R.; Liu, M.; Liu, Z.; Wang, Y.; Luan, D.; Wu, K.; Song, Z.; Sun, T.; Ma, Y.; Zhang, Y.; Wang, Q.; Li, X.; Ji, P.; Li, Y.; Li, C.; Wu, Y.; Ying, T.; Wen, Y.; Jiang, S.; Zhu, T.; Lu, L.; Zhang, Y.; Zhou, Q.; Huang, J. Antibody-Dependent Enhancement (ADE) of SARS-CoV-2 Infection in Recovered COVID-19 Patients: Studies Based on Cellular and Structural Biology Analysis. *medRxiv* **2020**, 2020.10.08.20209114. https://doi.org/10.1101/2020.10.08.20209114.
223. Sedova, E. S.; Scherbinin, D. N.; Lysenko, A. A.; Alekseeva, S. V.; Artemova, E. A.; Shmarov, M. M. Non-Neutralizing Antibodies Directed at Conservative Influenza Antigens. *Acta Naturae* **2019**, *11* (4), 22–32. https://doi.org/10.32607/20758251-2019-11-4-22-32.
224. Smatti, M. K.; Al Thani, A. A.; Yassine, H. M. Viral-Induced Enhanced Disease Illness. *Front. Microbiol.* **2018**, *9*, 2991. https://doi.org/10.3389/fmicb.2018.02991.
225. Khurana, S.; Loving, C. L.; Manischewitz, J.; King, L. R.; Gauger, P. C.; Henningson, J.; Vincent, A. L.; Golding, H. Vaccine-Induced Anti-HA2 Antibodies Promote Virus Fusion and Enhance Influenza Virus Respiratory Disease. *Sci. Transl. Med.* **2013**, *5* (200), 200ra114. https://doi.org/10.1126/scitranslmed.3006366.
226. van Erp, E. A.; van Kasteren, P. B.; Guichelaar, T.; Ahout, I. M. L.; de Haan, C. A. M.; Luytjes, W.; Ferwerda, G.; Wicht, O. In Vitro, Enhancement of Respiratory Syncytial Virus Infection by Maternal Antibodies Does Not Explain Disease Severity in Infants. *J. Virol.* **2017**, *91* (21), e00851-17. https://doi.org/10.1128/JVI.00851-17.
227. Bolles, M.; Deming, D.; Long, K.; Agnihothram, S.; Whitmore, A.; Ferris, M.; Funkhouser, W.; Gralinski, L.; Totura, A.; Heise, M.; Baric, R. S. A Double-Inactivated Severe Acute Respiratory Syndrome Coronavirus Vaccine Provides Incomplete Protection in Mice and Induces Increased Eosinophilic Proinflammatory Pulmonary Response upon Challenge. *J. Virol.* **2011**, *85* (23), 12201–12215. https://doi.org/10.1128/JVI.06048-11.
228. Tseng, C.-T.; Sbrana, E.; Iwata-Yoshikawa, N.; Newman, P. C.; Garron, T.; Atmar, R. L.; Peters, C. J.; Couch, R. B. Immunization with SARS Coronavirus Vaccines Leads to Pulmonary Immunopathology on Challenge with the SARS Virus. *PloS One* **2012**, *7* (4), e35421. https://doi.org/10.1371/journal.pone.0035421.
229. Yasui, F.; Kai, C.; Kitabatake, M.; Inoue, S.; Yoneda, M.; Yokochi, S.; Kase, R.; Sekiguchi, S.; Morita, K.; Hishima, T.; Suzuki, H.; Karamatsu, K.; Yasutomi, Y.; Shida, H.; Kidokoro, M.; Mizuno, K.; Matsushima, K.; Kohara, M. Prior Immunization with Severe Acute Respiratory Syndrome (SARS)-Associated Coronavirus (SARS-CoV) Nucleocapsid Protein Causes Severe Pneumonia in Mice Infected with SARS-CoV1. *J. Immunol.* **2008**, *181* (9), 6337–6348. https://doi.org/10.4049/jimmunol.181.9.6337.
230. Weingartl, H.; Czub, M.; Czub, S.; Neufeld, J.; Marszal, P.; Gren, J.; Smith, G.; Jones, S.; Proulx, R.; Deschambault, Y.; Grudeski, E.; Andonov, A.; He, R.; Li, Y.; Copps, J.; Grolla, A.; Dick, D.; Berry, J.; Ganske, S.; Manning, L.; Cao, J. Immunization with Modified Vaccinia Virus Ankara-Based Recombinant Vaccine against Severe Acute Respiratory Syndrome Is Associated with Enhanced Hepatitis in Ferrets. *J. Virol.* **2004**, *78* (22), 12672–12676. https://doi.org/10.1128/JVI.78.22.12672-12676.2004.
231. Wang, S.-F.; Tseng, S.-P.; Yen, C.-H.; Yang, J.-Y.; Tsao, C.-H.; Shen, C.-W.; Chen, K.-H.; Liu, F.-T.; Liu, W.-T.; Chen, Y.-M. A.; Huang, J. C. Antibody-Dependent SARS Coronavirus Infection Is Mediated by Antibodies against Spike Proteins. *Biochem. Biophys. Res. Commun.* **2014**, *451* (2), 208–214. https://doi.org/10.1016/j.bbrc.2014.07.090.
232. Jaume, M.; Yip, M. S.; Cheung, C. Y.; Leung, H. L.; Li, P. H.; Kien, F.; Dutry, I.; Callendret, B.; Escriou, N.; Altmeyer, R.; Nal, B.; Daëron, M.; Bruzzone, R.; Peiris, J. S. M. Anti-Severe Acute Respiratory Syndrome Coronavirus Spike Antibodies Trigger Infection of Human Immune Cells via a PH- and Cysteine Protease-Independent FcγR Pathway. *J. Virol.* **2011**, *85* (20), 10582–10597. https://doi.org/10.1128/JVI.00671-11.
233. Agrawal, A. S.; Tao, X.; Algaissi, A.; Garron, T.; Narayanan, K.; Peng, B.-H.; Couch, R. B.; Tseng, C.-T. K. Immunization with Inactivated Middle East Respiratory Syndrome Coronavirus Vaccine Leads to Lung Immunopathology on Challenge with Live Virus. *Hum. Vaccines Immunother.* **2016**, *12* (9), 2351–2356. https://doi.org/10.1080/21645515.2016.1177688.
234. Rauch, S.; Jasny, E.; Schmidt, K. E.; Petsch, B. New Vaccine Technologies to Combat Outbreak Situations. *Front. Immunol.* **2018**, *9*.
235. Ko, J.-H.; Müller, M. A.; Seok, H.; Park, G. E.; Lee, J. Y.; Cho, S. Y.; Ha, Y. E.; Baek, J. Y.; Kim, S. H.; Kang, J.-M.; Kim, Y.-J.; Jo, I. J.; Chung, C. R.; Hahn, M.-J.; Drosten, C.; Kang, C.-I.; Chung, D. R.; Song, J.-H.; Kang, E.-S.; Peck, K. R. Serologic Responses of 42 MERS-Coronavirus-Infected Patients According to the Disease Severity. *Diagn. Microbiol. Infect. Dis.* **2017**, *89* (2), 106–111. https://doi.org/10.1016/j.diagmicrobio.2017.07.006.
236. Peiris, J. S. M.; Chu, C. M.; Cheng, V. C. C.; Chan, K. S.; Hung, I. F. N.; Poon, L. L. M.; Law, K. I.; Tang, B. S. F.; Hon, T. Y. W.; Chan, C. S.; Chan, K. H.; Ng, J. S. C.; Zheng, B. J.; Ng, W. L.; Lai, R. W. M.; Guan, Y.; Yuen, K. Y.; HKU/UCH SARS Study Group. Clinical Progression and Viral Load in a Community Outbreak of Coronavirus-Associated SARS Pneumonia: A Prospective Study. *Lancet Lond. Engl.* **2003**, *361* (9371), 1767–1772. https://doi.org/10.1016/s0140-6736(03)13412-5.
237. Hsueh, P.-R.; Hsiao, C.-H.; Yeh, S.-H.; Wang, W.-K.; Chen, P.-J.; Wang, J.-T.; Chang, S.-C.; Kao, C.-L.; Yang, P.-C. Microbiologic Characteristics, Serologic Responses, and Clinical Manifestations in Severe Acute Respiratory Syndrome, Taiwan1. *Emerg. Infect. Dis.* **2003**, *9* (9), 1163–1167. https://doi.org/10.3201/eid0909.030367.
238. Be, Y.; Swx, O.; Lfp, N.; De, A.; Wn, C.; Py, C.; Lw, A.; Tm, M.; S, K.; Lya, C.; S, P.; Sy, T.; L, S.; P, P.; Sw, F.; Yh, C.; Cw, T.; B, L.; O, R.; Y, D.; P, T.; Jgh, L.; L, C.; T, B.; Rtp, L.; Ys, L.; L, R.; Lf, W.; Dc, L.; undefined. Viral Dynamics and Immune Correlates of Coronavirus Disease 2019 (COVID-19) Severity. *Clin. Infect. Dis. Off. Publ. Infect. Dis. Soc. Am.* **2021**, *73* (9), e2932–e2942. https://doi.org/10.1093/cid/ciaa1280.
239. *Regeneron halts enrolment of severely ill patients in COVID-19 antibody trial*. PMLive. https://www.pmlive.com/pharma\_news/regeneron\_halts\_enrolment\_of\_severely\_ill\_patients\_in\_covid-19\_antibody\_trial\_1355837 (accessed 2023-01-11).
240. Wan, Y.; Shang, J.; Graham, R.; Baric, R. S.; Li, F. Receptor Recognition by the Novel Coronavirus from Wuhan: An Analysis Based on Decade-Long Structural Studies of SARS Coronavirus. *J. Virol.* **2020**, *94* (7). https://doi.org/10.1128/JVI.00127-20.
241. Liu, L.; Wei, Q.; Lin, Q.; Fang, J.; Wang, H.; Kwok, H.; Tang, H.; Nishiura, K.; Peng, J.; Tan, Z.; Wu, T.; Cheung, K.-W.; Chan, K.-H.; Alvarez, X.; Qin, C.; Lackner, A.; Perlman, S.; Yuen, K.-Y.; Chen, Z. Anti-Spike IgG Causes Severe Acute Lung Injury by Skewing Macrophage Responses during Acute SARS-CoV Infection. *JCI Insight* **2019**, *4* (4), e123158, 123158. https://doi.org/10.1172/jci.insight.123158.
242. Hotez, P. J.; Corry, D. B.; Bottazzi, M. E. COVID-19 Vaccine Design: The Janus Face of Immune Enhancement. *Nat. Rev. Immunol.* **2020**, *20* (6), 347–348. https://doi.org/10.1038/s41577-020-0323-4.