Abstract In December 2019, Wuhan city in the Hubei province of China reported for the first time a cluster of patients infected with a novel coronavirus, since then there has been an outburst of this disease across the globe affecting millions of human inhabitants. Severe acute respiratory syndrome coronavirus type-2 (SARS-CoV-2), is a member of beta coronavirus family which upon exposure caused a highly infectious disease called novel coronavirus disease-2019 (COVID-19). COVID-19, a probably bat originated disease was declared by World Health Organization (WHO) as a global pandemic in March 2020. Since then, despite rigorous global containment and quarantine efforts, the disease has affected nearly 56,261,952 laboratory confirmed human population and caused deaths of over 1,349,506 lives worldwide. Virus passes in majority through respiratory droplets and then enters lung epithelial cells by binding to angiotensin converting enzyme 2 (ACE2) receptor and there it undergoes replication and targeting host cells causing severe pathogenesis. Majority of human population exposed to SARS-CoV-2 having fully functional immune system undergo asymptomatic infection while 5–10% are symptomatic and only 1–2% are critically affected and requires ventilation support. Older people or people with co-morbidities are severely affected by COVID-19. These categories of patients also display cytokine storm due to dysfunctional immune response which brutally destroys the affected organs and may lead to death in some. Real time PCR is still considered as standard method of diagnosis along with other serology, radiological and biochemical investigations. Till date, no specific validated medication is available for the treatment of COVID-19 patients. Thus, this review provides detailed knowledge about the different landscapes of disease incidence, etiopathogenesis, involvement of various organs, diagnostic criteria’s and treatment guidelines followed for management of COVID-19 infection since its inception. In conclusion, extensive research to recognize novel pathways and their cross talk to combat this virus in precarious settings is our future positive hope.

Keywords Severe acute respiratory syndrome coronavirus-2 (SARS-COV-2) · Novel coronavirus disease-2019 (COVID-19) · Etiopathogenesis · Clinical trials

Introduction

An outbreak of highly infectious novel coronavirus disease-2019 (COVID-19) emerged in December 2019 in Wuhan city in the Hubei province of China which rapidly spread across worldwide and was later declared pandemic by WHO on March 11, 2020 [1]. The disease was caused by a novel coronavirus, severe acute respiratory syndrome coronavirus type-2 (SARS-CoV-2) classified on the same day by International Committee on Taxonomy of Viruses [2]. Since then, disease is continuously spreading and as per the WHO report till 20th November, it has nearly targeted 56,261,952 infected cases and cost over 1,349,506 human lives [3]. In India, first case of COVID-19 was reported on January 30, 2020, which originated from
China. According to Indian Council of Medical Research (ICMR) and Ministry of Health and Family Welfare (MOHFW), total 9,004,365 confirmed cases were reported, whereas 132,162 deaths occurred till November 20, 2020 [4, 5].

Shanghai Public Health Clinical Center and School of Public Health researchers on January 7, 2020 jointly revealed the origin of this coronavirus from a seafood market in Wuhan city [6]. Number of cases started increasing since then, even in people not having any exposure to animal market, and hence human-to-human transmission was also reported to occur [7]. Coronavirus (CoVs) encompass a large virus family affecting mainly humans and different animal species including cats, cattles, camels and also bats [8]. Not so often, but animal coronaviruses have also infected human, recently introduced SARS-CoV2 has joined the list of middle east respiratory syndrome coronavirus (MERS-CoV) and severe acute respiratory syndrome coronavirus (SARS-CoV). SARS-CoV2, a beta coronavirus, has probably originated from bats just like MERS-CoV and SARS-CoV [9]. Human coronaviruses in majority causes respiratory and enteric infections and SARS-CoV-2 infection causes flu like symptoms ranging from fever, cough, headache, asthenia etc. Once the virus reaches to respiratory epithelial cells, it enters by binding to angiotensin converting enzyme-2 through its spike proteins and is then endocytosed inside the host cells. Once inside the cells, the virus starts replicating and targets variety of host cells thereby causing severe pathologies. However, SARS-CoV-2 infection affects people of all ages, in some high-risk individuals, such as older people or those having co-morbidities, the virus might cause severe infections such as interstitial pneumonia, acute respiratory distress syndrome (ARDS), progressing to multi organ failure and ultimately causing respiratory failure leading to death [10]. Cytokine storm is one of the characteristics of COVID-19 infection majorly occurring in persons having dysfunctional immune responses leading to secretion of pro-inflammatory cytokines such IL-16, IL-1β, TNF-α, IL-2, IL-7, MCP1, G-CSF, IP-10, IL-10, MIP1α and others. This review, focusses on current incidence, etiopathogenesis, clinical characteristics with multiorgan involvements associated infections, diagnostic criteria’s, its management and present and future prospects of clinical trials to prevent, manage and control COVID-19.

Structural Aspects of Coronavirus

Coronavirus (CoVs) are the largest group of viruses belongs to the Nidovirales order that comprises three families i.e. arteriviridae, roniviridae and coronaviridae [11]. On the basis of genome structure, coronaviridae comprises of four genera- alpha coronavirus, beta coronavirus, gamma coronavirus and delta coronavirus. Alpha coronavirus and beta coronavirus transmission are limited to mammals and in humans they cause respiratory illnesses such as SARS and MERS, whereas gamma coronaviruses and delta coronaviruses infect birds as well as mammals [12, 13]. Coronaviruses are 60 to 140 nm in diameter and have 26–32 kilobases positive sense single stranded RNA genome connected to a nucleoprotein surrounded by capsid [14]. A typical CoV has at least six open reading frames (ORF) and all proteins including structural and accessory are encoded by a single guide RNA [15]. Two major overlapping open reading frames (ORFs) i.e. ORF1a and ORF1b comprise of two-third of the whole genome and the third ORF codes for four structural proteins i.e. spike (S), envelope (E), membrane (M), and nucleocapsid (N). SARS-CoV-2 genome encodes, a very large polyproteins pp1ab, four structural proteins and six accessory proteins (3a, 6, 7a, 7b, 8, and 10) which is equivalent to the structure of SARS-CoV and MERS-CoV [16, 17]. (Figs. 1 and 2).

The Spike (S) proteins has been divided into functional subunits i.e. N-terminal S1 subunit and a C-terminal S2 subunit. The size of each monomer of S protein is 180 kDa and is involved in attachment and membrane fusion. Depending on the virus, both N and C terminal domain are involved in the membrane attachment and entry of viral genome inside the host cells. Apart from this, an additional furin like cleavage site is present at the junction of S1 and S2 subunits of S protein in SARS-CoV-2, which is absent in the genome of SARS-CoV. This additional subunit has been speculated to be involved in higher infectivity of SARS-CoV-2 than SARS-COV [18]. Further, using phylogenetic tree investigation that SARS-CoV-2 genome shares 79% genetic similarity with SARS-CoV [19] whereas it demonstrates 98% genetic similarities with bat coronavirus RaTG13 [20] and is 85.98% identical to pangolin-CoV genome [21] thereby suggesting its zoonotic origin.

Further, whole genome sequencing analyses have revealed that mutations occurring in the spike protein are
important in the evolution of SARS-CoV-2. Phylogenetic and alignment study of 591 novel coronaviruses of different clades from Group I to Group V identified several mutations and related amino acid changes and majority were distributed over spike protein [22]. In a major study involving investigation of 3617 available whole genome sequence of SARS-COV-2 maintained in NCBI globally identified that E protein possess several non-synonymous mutations [23]. Additionally, from five different isolates identified in the state of West Bengal (Eastern state of India), India, two major mutations one at position 723 and other at 1124 were identified in the S2 domain of Spike (S) protein. Another mutation at 614 position in S1 domain of S protein was also identified which is similar to mutation found in the isolates of state of Gujarat (Western state of India). These mutations were crucial in modulating the affinity of receptor binding [24]. Thus, these whole genome study revealed that SARS-CoV-2 is continuously evolving virus with crucial mutation identified in the receptor binding regions.

**SARS CoV-2 Transmission and Its Etiopathogenesis**

Virus spreads mainly from person to person through respiratory droplets or bits of liquid, mostly through sneezing or coughing [25–28]. According to National Institutes of Health (NIH), SARS-CoV-2 virus is stable in aerosols and on surfaces for several hours to days depending upon the surface materials and also ambient environmental conditions. Virus has been reported to be detectable up to three hours in aerosols, up to four hours on copper, up to 24 h on cardboard and up to two to three days on plastic and stainless steel [29]. It has also been testified in stool and may contaminate water supply possibly and subsequently results in aerosol and/or feco-oral based route of transmission [30]. Another study reported the presence of SARS-CoV-2 virus in the tears and conjunctival secretions [31].

Virus can pass through nasal and larynx mucous membranes and enter into the lungs through respiratory tract. First step of any viral infection is the binding of some viral proteins with receptors expressed by host cells followed by fusion with host cell membrane. ACE2 has been speculated to be the ultimate target of spike protein of SARS-CoV-2. Initial target of virus include lung epithelial cells, where virus attach through its spikes to cellular angiotensin converting enzymes 2 (ACE2) receptor [32].

ACE2 is a membrane bound metalloproteinase and was discovered in 2000. It is a homolog of ACE, though its activity cannot be stopped by angiotensin-converting enzyme inhibitors (ACEIs) [33]. Further study reported that SARS-CoV-2 has nearly 10–20 folds higher affinity for ACE2 than SARS-CoV. ACE2 is also significantly expressed in other tissues like heart, renal, and gastrointestinal tract [34, 35]. It is an important regulator of renin-angiotensin system (RAS) and renin angiotensin aldosterone system (RAAS). RAS is an indispensable system of human body which is regulated and counter-regulated by two main pathways: classical RAS pathway ACE-angiotensin II-angiotensin I receptor pathway, and counter-regulatory RAS axis ACE2-angiotensin 1–7–MasR-based pathway, which plays a negative role in regulation [36, 37]. In classical RAS pathway, angiotensinogen is cleaved into angiotensin (Ang) by renin, which is then cleaved by ACE into angiotensin II. This Ang II after binding to type 1 angiotensin II receptor (AT1R) mediate functions like increases secretion of aldosterone and anti-diuretic hormone, increases blood pressure by decreasing sensitivity to baroreflex, increases vasoconstriction, decreases production of nitric oxide (NO), increases fibrosis, reactive oxygen species production and inflammation. Whereas ang II can also be cleaved by ACE2 in counter-regulated pathway into Ang-(1–9) and Ang-(1–7), Ang (1–9) which then binds to AT2R and triggers NO production thus mediating vasodilatation and reducing blood pressure. In addition to that, it also reduces inflammation, fibrosis and cardiac hypertrophy. Further, Ang-(1–7) binds to proto-oncogene Mas receptor (MasR) and reduces the blood pressure and reverses all its function performed during classical RAS pathway.
SARS-CoV-2 attaches to the receptor of host cell through S1 domains of its spike glycoprotein (S), and this spike protein is proteolytically cleaved by transmembrane serine protease 2 (TMPRSS2) into S1 and S2 subunits and then S2 induces membrane fusion and virus internalization by endocytosis [38]. S1 regulates virus-host range and cellular tropism with the receptor binding domain (RBD), where S2 helps in virus cell membrane fusion with the help of two tandem domains, heptad repeats 1 and 2 (HR1 & HR2) [39, 40]. After the fusion it releases viral genome RNA into cytoplasm via endocytosis and replication and transcription of virus occur in cytoplasm. Positive RNA genome acts as a messenger RNA (mRNA). Once viral genome enters into host cell, it translates into pp1a and pp1ab, two large precursor polyproteins [41] and these polyproteins by ORF 1a-encoded viral proteinases, 3C-like proteinases (3CLpro) and papain-like proteinase (PLpro) get processed into 16 mature nonstructural proteins (nsp1–nsp16) [9, 42]. During viral RNA replication and transcription these nonstructural proteins (nsp) perform vital functions [43] (Fig. 3). This viral replication and cell to cell transmission plays important role in the suppression of ACE2 expression. This suppression of ACE2 leads to decrease in Ang (1–7) synthesis and enhanced levels of Ang II which drives the Ang II–AT1R dependent inflammatory pathways in the lungs and causes parenchymal injury [44]. SARS-CoV-2 enhanced the apoptosis and p53 signaling pathways in lymphocytes thereby causing the lymphopenic situation in COVID-19 patients [45]. Additionally, coagulopathy has also been observed in COVID-19 infection, as elevated levels of plasmin(ogen) has been observed in patients. This plasmin along with other proteases might play an important role in the cleavage of furin site in the S protein of SARS-CoV-2, thus increasing its virulence and infectivity and is also associated with hyperfibrinolysis [46]. Among the existing coronaviruses, RNA recombination along with its potential proof-reading capacity has been involved in the development and occurrence of novel coronaviruses [47]. Organs presenting higher ACE2 expression are presented as possible targets.
of SARS-CoV-2 virus infection such as lung being an important and primary infection target organ. Further, ACE2 expression and its organ wise distribution is significantly associated with COVID-19 clinical symptoms [25, 48]. (Fig. 4).

**Symptoms And Etiological Factors Associated With Pathogenesis**

Nearly, 80% of the SARS-CoV-2 infected patients display mild symptoms, whereas 15% cases come in severe category and rest 5% fall under critical category requiring ventilation support [49]. The course of infection of SARS-CoV-2 spans from mild disease limiting itself to upper respiratory, non-severe pneumonia, could be severe pneumonia involving ARDS, multiple organ failure and ultimately death [50]. Common clinical symptoms include fever, dry cough, headache, sore throat, breathlessness, diarrhea, vomiting and abdominal pain [51, 52]. Olfactory and taste desensitization have also been reported in COVID-19 [53].

People who have some chronic medical comorbidities such as cardiovascular disease, any lung problems or diabetes or who are older and have higher risk of severe illness (chart out in Tables 1 and 2). Children who are infected with SARS-CoV-2 generally display mild symptoms which might be due to altered ACE2 activity and active innate immune response. Whereas adult patients display suppressed adaptive immunity and dysfunctional immune response. Further, it is also observed that women are less vulnerable to viral infection compared to men which may be due to incongruent immune system, steroid hormone or sex chromosome associated factors. In the next few paragraphs, we will observe that how infection with SARS-CoV-2 affects different organs of the body and also on the dysfunctional immune response.

**Immune Response With Cytokine Storm in COVID-19**

Immune system of an individual is highly responsible to depict asymptomatic or symptomatic clinico manifestations of COVID-19. Geriatric population with poor immune function along with comorbidities is at increased risk with more susceptibility to various viral and bacterial infections due to less efficient, less coordinated and slower immune response. Cytokine storm noted in critical cases of COVID-19 is a response of an uncontrolled immune response.
mechanism with an uncontrolled release of cytokines. Multiple factors are involved in triggering cytokine storm which includes virus, bacterial components, sepsis, super antigens, toxins, chimeric antigen receptor T cells and others [60]. It is a life threatening condition leading to detrimental changes including capillaries leakage, edema, tissue toxicity, organ failure and even shock. Increased levels of IL-6 were significantly observed with clinical manifestation in critical COVID-19 cases [61]. A trend has been observed where, SARS-CoV-2 infection rapidly activates CD4 T lymphocytes thereby forming pathogenic T helper (Th1) cells and release various cytokines including GM-CSF. This cytokine environment, induces an inflammatory environment with production of CD14 +, CD16 + monocytes along with enhanced secretion of IL-6. Finally, pathogenic T-cells along with monocytes enter pulmonary circulation where monocytes induce to become macrophages [62]. Acute respiratory distress syndrome (ARDS) is the most severe form of acute lung injury. ACE2 expression is very high in lung tissue and is associated with protection of lung tissue injury induced by sepsis [67]. Decreased expression of ACE2 is inversely correlated with angiotensin II levels thereby enhanced its production. Further, this angiotensin II stimulates its type 1a receptor, which then enhances lung vascular permeability and leads to its pathological phenotype [14].

Single-cell RNA sequencing data of 43,134 human lung cells reported that only 0.64% of total cells expressed ACE2 and out of which 83% of total ACE2 expression was present on type II apical surface of epithelial cells (AT2) [41]. Other cellular types including type I epithelial cells, airway epithelial cells, endothelial and others also expresses ACE2 but at levels considerably lower than AT2 cells [68]. Thus, these results indicated AT2 cells as primary targets of SARS-CoV-2 in lungs. Hypoxia is one of the clinical symptoms associated with various pulmonary diseases. During initial phase of hypoxia, enhanced expression of ACE2 was determine in pulmonary smooth muscle cells whereas during later phase. Hypoxia-Inducible Factor-1 (HIF-1α), induces ACE expression (one of the targets of HIF-1α) with concomitant decreased in the expression of ACE2 [69].

**Lung Injury in COVID-19**

Coronavirus infection frequently occurs in patients with pre-existing lung disease such as asthma and COPD [64, 65]. Lung tissue has high renin-angiotensin system (RAS) activity which is chief site of Ang II synthesis. Ang II is an effective pulmonary vasoconstrictor [66]. Ang II over activity also promotes pulmonary edema and impairs lung function [67]. Acute respiratory distress syndrome (ARDS) is the most severe form of acute lung injury. ACE2 expression is very high in lung tissue and is associated with protection of lung tissue injury induced by sepsis [67]. Decreased expression of ACE2 is inversely correlated with angiotensin II levels thereby enhanced its production. Further, this angiotensin II stimulates its type 1a receptor, which then enhances lung vascular permeability and leads to its pathological phenotype [14].

**Kidney Injury in COVID-19**

Kidneys are presented as another major targets of SARS-CoV-2 viral infection as virus was detected in the urine samples of few COVID-19 positive cases [25]. In these patients, acute kidney injury is presented as a critical
| S. no | Author          | Country | Sample size | Disease and symptoms       | Study outcomes                                                                                                                                                                                                                                                                                                                                 |
|-------|-----------------|---------|-------------|----------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| 1     | Zhu 2019        | China   | Patients with pneumonia                  | Human airway epithelial cells were used to isolate a novel coronavirus, because they appear to be a valuable research tool for analysis of human respiratory pathogens. Study showed that initial propagation of human respiratory secretions onto human airway epithelial cell cultures, followed by transmission electron microscopy and whole genome sequencing of culture supernatant, was successfully used for visualization and detection of new human coronavirus. |
| 2     | Guan 2019       | China   | Common symptoms were fever (43.8% on admission and 88.7% during hospitalization) and cough (67.8%). Diarrhea was uncommon (3.8%). Lymphocytopenia was present in 83.2% | Patients often presented without fever, and many did not have abnormal radiologic findings.                                                                                                                                                                                                                                                                 |
| 3     | Huang 2019      | Wuhan, China | Common symptoms fever (40 of 41 patients), cough (31), and myalgia or fatigue (18); less common symptoms were sputum production (11 of 39), headache (three of 38), haemoptysis (two of 39), and diarrhoea (one of 38) | Complications included acute respiratory distress syndrome (12 [29%]), RNAemia (six [15%]), acute cardiac injury (five [12%]) and secondary infection (four [10%]). 13 (32%) patients were admitted to an ICU and six (15%) died. Compared with non-ICU patients, ICU patients had higher plasma levels of IL2, IL7, IL10, GSCF, IP10, MCP1, MIP1A, and TNFα. |
| 4     | Chen 2020       | China   | Patients had clinical manifestations of fever (82 [83%] patients), cough (81 [82%] patients), shortness of breath (31 [31%] patients), muscle ache (11 [11%] patients), confusion (nine [9%] patients), headache (eight [8%] patients), sore throat (five [5%] patients), rhinorrhea (four [4%] patients), chest pain (two [2%] patients), diarrhoea (two [2%] patients), and nausea and vomiting (one [1%] patient) | Characteristics of patients who died were in line with the MuLBSTA score, an early warning model for predicting mortality in viral pneumonia. 74 (75%) patients showed bilateral pneumonia, 14 (14%) patients showed multiple mottling and ground-glass opacity, and one (1%) patient had pneumothorax. 17 (17%) patients developed acute respiratory distress syndrome and, among them, 11 (11%) patients worsened in a short period of time and died of multiple organ failure. |
| 5     | Dawei Wang 2020 | Wuhan, China | Common symptoms included fever (136 [98.6%]), fatigue (96 [69.6%]), and dry cough (82 [59.4%]). Lymphopenia (lymphocyte count, 0.8 × 10^9/L [interquartile range {IQR}, 0.6–1.1]) occurred in 97 patients (70.3%), prolonged prothrombin time (13.0 s [IQR, 12.3–13.7]) in 80 patients (58%), and elevated lactate dehydrogenase (261 U/L [IQR, 182–403]) in 55 patients (39.9%). Bilateral patchy shadows or ground glass opacity in the lungs of all patients | Most patients received antiviral therapy (oseltamivir,124 [89.9%]), and many received antibacterial therapy (moxifloxacin,89 [64.4%]; ceftriaxone, 34 [24.6%]; azithromycin, 25 [18.1%]) and glucocorticoid therapy (62 [44.9%]). In 41% of patients, 26% of patients received ICU care, and mortality was 4.3%. |
complication targeting renal intrinsic cells including proximal tubular epithelial cell thus leading to kidney dysfunction. Kidney expresses very high levels of ACE2, which is very high especially in the renal cortex [70]. During acute kidney disease [71] and other models of chronic kidney diseases, ACE2 expression levels are severely compromised which then deregulated the homeostasis of RAS and causes debilitating pathological changes in the kidneys [72].

Table 2 continued

| S. no | Author         | Country       | Sample size | Disease and symptoms | Study outcomes                                                                                                                                                                                                 | References |
|-------|----------------|---------------|-------------|----------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|
| 6     | Xi Jin         | China         | 651 analyzd COVID-19 cases with GI symptoms | 74 (11.4%) presented with at least one GI symptom (nausea, vomiting or diarrhoea). COVID-19 with GI symptoms, 29 (39.19%), 23 (31.08%), 8 (10.81%) and 16 (21.62%) had significantly higher rates of fever > 38.5 °C, fatigue, shortness of breath and headache, respectively | Sputum production and increased lactate dehydrogenase/glucose levels were risk factors for severe/critical type. Bioinformatics showed sequence mutation of SARS-CoV-2 with m6A methylation and changed binding capacity with ACE2 | [57]       |
| 7     | Lin L          | China         | 95          | 58 cases exhibited GI symptoms of which 11 (11.6%) occurred on admission and 47 (49.5%) developed during hospitalisation. Diarrhoea (24.2%), anorexia (17.9%) and nausea (17.9%) were the main symptoms with five (5.3%), five (5.3%) and three (3.2%) cases occurred on the illness onset, respectively | Patients with GI symptoms were subjected to endoscopy, revealing oesophageal bleeding with erosions and ulcers in one severe patient. SARS-CoV-2 RNA was detected in oesophagus, stomach, duodenum and rectum specimens for both two severe patients. In contrast, only duodenum was positive in one of the four non-severe patients. GI tract may be a potential transmission route and target organ of SARS-CoV-2 | [58]       |
| 8     | Zhenyu Fan     | Shanghai, China | 148        | Patients had clinical manifestations of fever (70.1%), cough (45.3%), expectoration (26.7%) at admission. 75 patients (50.7%) showed abnormal liver functions at admission. Numbers of CD4+ and CD8+ T cells were significantly lower, and utilization rate of lopinavir/ritonavir significantly higher in abnormal liver function group than those in normal liver function group | SARS-CoV-2 may cause the liver function damage and the Lopinavir/ritonavir should be applied carefully for the treatment of COVID-19 | [59]       |

Fig. 5 Diagrammatic representation of Systemic Inflammatory Response Syndrome (SIRS) mediated by SARS-CoV-2
Gastrointestinal Tract (GIT) in COVID-19

Some COVID-19 patients have depicted gastrointestinal symptoms like diarrhea, vomiting and abdominal pain. Previously reported studies suggested relation between coronavirus and gastrointestinal tract tropism [60]. In coronavirus infection, tropism of gastrointestinal may explain an incidence of diarrhea. SARS-CoV RNA might be detected in SARS patients stool specimens [73]. SARS-CoV-2 detected in fecal samples was most likely due to virus entering the blood from lungs and then traveling from blood to intestines [74]. This infected fecal sample can promote fomites-based transmission, especially because of generation of infectious aerosol from toilet plume. Literature has reported very high expression of ACE2 receptor in GIT, particularly in small and large intestines [75, 76]. Positive staining of viral nucleocapsid was detected in cytoplasms of gastric, duodenal and rectal epithelium cells [77]. These evidences provided indispensable information about mode of entry of virus into host cells and further unveiled the possible route of transmission. Some studies reported viral RNA presence in the stool of COVID-19 patients [78–80]. In another study of 73 COVID-19 patients, 39 (53.4%) were positive for SARS-CoV-2 RNA in stool, where duration of viral positivity ranged from 1 to 12 days. Even more, 17 (23.3%) patients that were negative for respiratory samples showed positive results with viral RNA in stool [75].

Liver Injury in COVID-19

Patients with COVID-19 have been reported to have high levels of serum Alanine Transaminase (ALT) and Aspartate Transaminase (AST) [48, 77]. In a study, 56 COVID-19 patients, 54% patients have high levels of serum Gamma-Glutamyl Transferase (GGT) [81]. Chai et al. stated that ACE2 expressed in both liver cells and bile duct cells [82]. Nevertheless, expression of ACE2 in bile duct cells was higher as compared to liver cells. Epithelial cells of bile duct play important roles in liver regeneration and immune response [83]. According to these results in COVID-19, patient’s liver injury occurred may be due to damage of bile duct cells, but not liver cells by virus infection.

Cardiovascular Diseases Associated with COVID-19

Respiratory failure is the main cause of death from COVID-19 patients, mostly in older adults and those who have weaker immune system [84, 85]. Most common cardiovascular (CV) complication is acute myocardial injury. SARS-CoV-2 enters human cells through ACE2 receptor. In normal healthy and in various disease conditions, ACE2 helps in neurohormonal regulation of CV system. When SARS-CoV-2 binds with ACE2R it alters the ACE2 signalling pathways that can lead to acute myocardial and lung injuries [86]. Roughly, 8–12% cases of acute myocardial injury reported significant elevation of cardiac troponin I (cTnI) [86]. In a meta-analysis from China [66, 87], incidences of acute cardiac injury were reported to be 8%; however, in another study, they included only those patients who were discharged from hospital or dead and found 17% incidence of cTnI elevation [88]. Clinical investigations suggested that patients with heart diseases, hypertension or diabetes who were treated with ACE2 increasing drugs including inhibitors and blockers showed increased expression of ACE2 and these were at higher risk of getting SARS-CoV-2 infection [89] (Table 3).

Diabetes Mellitus Associated with COVID-19

ACE2 modifies neutral amino acid transporters expression on epithelial cells surface and growth of pancreatic islet cells along with insulin secretion by pancreatic β-cells [70]. A study involving a mice model of adenovirus mediated human ACE2 expression showed enhanced production of insulin along with decreased apoptosis of pancreatic islets [96]. In addition to that, ACE2 also improves the overall endothelial function of pancreatic islet microvascular unit [97]. Some studies stated that diabetes mellitus (DM) was associated with more severe disease, acute respiratory distress syndrome and increased mortality [25, 98]. In a study by Guan et al., patients with DM showed higher disease severity (16.2%) in contrast, patients without DM showed lesser disease severity (5.7%) [25]. Further, an obvious conclusion was made, where patients having DM higher in age was compared to non-Diabetes mellitus patients. Thus, as age advances it emerged as crucial prognostic factor in determining course of COVID-19 disease. Simultaneously, patients with DM showed lowered ACE2 expression possibly due to glycosylation, which explained increased comorbidity with severe lung injury and ARDS in COVID-19 patients [99, 100]. COVID-19 patients with diabetes as coexisting disorders had a worse clinical outcome.

Diagnosis of COVID-19 Infection

Currently, multiple approaches are efficiently being used for the diagnosis of COVID-19 infection [101]. Two broadly classified techniques include, Real time reverse transcriptase polymerase chain reaction (rRT-PCR) based detection and serology-based detection. RT-PCR technique involves conversion of viral RNA genome into DNA, which is further amplified using specific primers set against defined targeted regions of viral genome. Major targeted regions of SARS-CoV-2 genome include nucleocapsid...
| S. No | Author                  | Study Objective                                                                 | Finding                                                                                                                                                                                                                                                                                                                                 | References |
|------|-------------------------|---------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|
| 1    | Peng Zhou 2020          | Identification and characterization of a new coronavirus (2019-nCoV), which caused an epidemic of acute respiratory syndrome in humans in Wuhan, China | The sequences are almost identical and having sequence identity 79.6% to SARS-CoV. Furthermore, 2019-nCoV is 96% identical at the whole-genome level to a bat coronavirus. Pairwise protein sequence analysis of seven conserved non-structural proteins domains show that this virus belongs to the species of SARSr-CoV. In addition, 2019-nCoV virus isolated from the bronchoalveolar lavage fluid of a critically ill patient could be neutralized by sera from several patients. It confirmed that 2019-nCoV uses the same cell entry receptor—angiotensin converting enzyme II (ACE2)—as SARS-CoV | [20]       |
| 2    | Markus Hoffmann 2020    | 2019-nCoV-S uses the SARS-coronavirus receptor, ACE2, for entry and the cellular protease TMPRSS2 for 2019-nCoV-S priming. A TMPRSS2 inhibitor blocked entry and might constitute a treatment option | The serum forms a convalescent SARS patient neutralized 2019-nCoV-S-driven entry. Our results reveal important commonalities between 2019-nCoV and SARS-coronavirus infection, which might translate into similar transmissibility and disease pathogenesis. Moreover, they identify a target for antiviral intervention | [90]       |
| 3    | Tatsuo Hashimoto 2012   | Report that deficiency in murine angiotensin I converting enzyme (peptidyl-dipeptidase A) 2 (Ace2), which encodes a key regulatory enzyme of the renin-angiotensin system (RAS), results in highly increased susceptibility to intestinal inflammation induced by epithelial damage. The RAS is known to be involved in acute lung failure, cardiovascular functions and SARS infections | Results identify ACE2 as a key regulator of dietary amino acid homeostasis, innate immunity, gut microbial ecology, and transmissible susceptibility to colitis. These results provide a molecular explanation for how amino acid malnutrition can cause intestinal inflammation and diarrhoea | [91]       |
| 4    | Andrew M. South 2020    | Examines the evidence for ACE2 regulation by RAAS blockade and statins, the cardiovascular benefits of ACE2, and whether ACE2 blockade is a viable approach to attenuate COVID-19 | Role of ACE2 in the progression of SARS-CoV-2 infection and, importantly, whether blockade of the peptidase is an appropriate step, at least acutely and targeted to the pulmonary system rather than systemically. Studies of the multi-faceted roles of the RAAS in the setting of infectious disease is warranted, which should not occur in isolation from the other well-known roles of the system in cardiovascular homeostasis, particularly given the recent focus on the RAAS and the balance of Ang II and Ang-in critical care medicine such as in septic shock | [92]       |
| 5    | Mahmoud Gheblawi 2020   | Recombinant ACE2, gene-delivery of Ace2, Ang 1–7 analogs, and Mas receptor agonists enhance ACE2 action and serve as potential therapies for disease conditions associated with an activated RAS (renin-angiotensin system). Recombinant human ACE2 has completed clinical trials and efficiently lowered or increased plasma angiotensin II and angiotensin 1–7 levels, respectively | Highlighting the critical role of ACE2 as the novel SARS-CoV-2 receptor and as the negative regulator of the RAS, together with implications for the COVID-19 pandemic and associated cardiovascular diseases | [93]       |
| 6    | Yi-Ming Yuan 2015       | To explore the role of the Renin–angiotensin–aldosterone system (RAAS) in the pathogenesis of pulmonary arterial hypertension (PAH) induced by chronic exposure to cigarette smoke | Chronic cigarette exposure may result in PAH and affect the protein expression of ACE2 and ACE in lung tissue, suggesting that ACE2 and ACE play an important role in the pathogenesis of smoking-induced PAH | [94]       |
| 7    | Yingxia Liu 2020        | The viral load of 2019-nCoV detected from patient respiratory tracts was positively linked to lung disease severity. Blood biochemistry indexes, albumin (ALB), CRP, LDH, LYM (%), LYM, and NEU (%), may be predictors of disease severity | Angiotensin II level in the plasma sample from 2019-nCoV infected patients was markedly elevated and linearly associated to viral load and lung injury. Results suggest a number of potential diagnosis biomarkers and angiotensin receptor blocker (ARB) drugs for potential repurposing treatment of 2019-nCoV infection | [55]       |
(N) gene, envelope (E) gene and ORF1ab gene regions [102]. RT-PCR based method displays high sensitivity (85–90%) and high specificity for the COVID-19 diagnosis as it is dedicated to the direct amplification of viral genetic material and the turnaround time (TAT) is about 2.5–3.5 h. This technique is quantitative in nature. Sample types for RT-PCR include nasopharyngeal swabs and oropharyngeal swabs. The RT-PCR results generally become positive after 2–8 days of infection and it is able to process large batches of samples. However, there are some limitations of this technique apart from skilled manpower, mutation in the viral genome might give false negative results, and further possible limitation includes improper collection, transportation and handling contributing to false negative results [103]. Rapid PCR by cartridge system (CBNAAT) are also available which are equally efficient and has significantly short response time [104]. Additionally, loop mediated isothermal amplification (LAMP) is another version of CBNAAT, which is highly specific isothermal amplification technique. In this method, transcription of RNA to DNA is then followed by amplification done using 4–6 primers set against multiple targeted regions [105]. This method has got several advantages over RT-PCR in being specific, user friendly in detection, speed (turnaround time is 20 min) and less background. Recently, Sree Chitra Thirunal Medical Institute and Technology in collaboration with Agappe Diagnostics Ltd. developed Chitra Gene LAMP-N that detects two different regions of N gene. In addition to above, CDC has approved a one-step rRT-PCR test kit to quantitatively determine viral molecules in > 90 samples within 45 min [106]. Additionally, Specific High Sensitivity Enzymatic Reporter Unlocking (SHERLOCK) employs the CRISPR-Cas system for the detection of viral RNA genome with greater specificity [107]. Similar technique was employed by Indian scientists who have developed FELUDA, a test kit for SARS-CoV-2 genome detection also employs CRISPR-Cas9 system [108].

Another, crucial diagnostic method is serology-based detection of SARS-CoV-2 infection among COVID-19 patients. This method employs detection of antigens/antibodies in the human blood. COVID-19 infection leads to the generation of IgM antibodies and then IgG antibodies inside the host body. Initially after infection, the titer of IgM increases within a week time. Later on, titer of IgG increases from day 4 post infection and reaches to its peak by day 14. Further, the levels of IgM degrade very rapidly whereas IgG antibodies titer persist in the body for longer duration thus helps in determination of active infection. Apart from antibodies detection, antigen detection (S and N protein) is also being explored for early detection of viral infection. Three major strategies are being employed for detection of Ags/Abs including lateral flow immunoassay, ELISA, and chemiluminescence.

Further apart from above two standardized techniques for detection of COVID-19 infection, other parameters such as high sensitivity CRP levels, lactate dehydrogenase, alanine transaminase, erythrocyte sedimentation rate have been observed [109]. Further, decreased lymphocyte count i.e. depletion of CD4+ and CD8+ cells and decreased IFN-γ expression in CD4+ T cells are linked with severe COVID-19, illustrating the cytokine storm. Another crucial observation is the presence of intravascular coagulation associated with increased D-dimer and fibrinogen levels in some COVID-19 infection. Lastly, chest X-ray (CXR) and computed tomography (CT) scan are considered as an essential tool in the detection of COVID-19 pneumonia during this pandemic.

**Treatment Strategies for COVID-19 Infection**

**ACE2 for Therapeutics**

Despite significant efforts made towards development of vaccines and therapeutic drugs, no significant advances have been observed till today. Currently available drugs are generally categorized according to their targets, one acts on
mode of entry of virus and other acts on inhibition of enzymes responsible for viral genome replication. Combination of lisinopril and losartan treatment in normotensive Lewis rats abolished increase in ACE2 mRNA levels observed individually but retained losartan induced rise in ACE2 activity in heart [110]. Ang II can regulate ACE2 expression through AT1R. Healthy hearts and kidneys displayed high levels of ACE2 mRNA and protein expression, with moderate expression of ACE [82]. RAS over activation in CVD increases AT1R stimulation by Ang II, promoting ERK1/2 and p38 MAPK signalling pathways to downregulate ACE2 while upregulating ACE expression [111]. Promoting ACE2/Ang 1–7/Mas signalling by rhACE2 or Ang 1–7 receptors agonist AVE 0991 can have valuable therapeutic effects in CVD and lung disease from diverse aetiologies [112]. Ang 1–7 receptors agonist AVE 0991 has been shown to exert cardio-renal and pulmonary protective effects, [113] and treatment with rhACE2 improved symptoms of acute lung injury, CVD, and kidney injury in various preclinical models [67, 70, 114]. Maintaining ACE2 levels in patients with or predisposed to common CVD states such as diabetes, hypertension, and obesity wards off the advancement of these comorbidities in instances where, patient contracts SARS-CoV-2 by maintaining levels of ACE2/Ang1–7/MasR negative counter-regulation.

**Vitamin D Response in COVID-19**

Vitamin D (calcitriol or vitamin D3) plays an important role in regulation of serum calcium concentration. Although less highlighted, several reviews have addressed role of vitamin D in prevention of viral infections [115, 116]. Multiple mechanisms are employed by vitamin D in reducing risk of microbial infection and diseases. In our previous papers, we have shown that various polymorphic variants of vitamin D receptor gene modulate the metabolism of blood lead levels along with circulatory levels of vitamin D [117, 118]. Such as in reducing risk of common cold, vitamin D employs three different categories of mechanisms: physical barrier, cellular natural immunity, and adaptive immunity [119, 120]. Some studies reported that administration of an oral dose of 50,000 IU of vitamin D reduces risks of influenza. Vitamin D adequacy also reduces severity of pneumonia, which is associated with coronavirus infection. Taking higher doses of vitamin D supplement or sun exposure is reported to boost immunity and reduce risk. Vitamin D helps in maintenance of various cellular junctions including tight junction, gap junction and adherence junction which are otherwise evaded by several viruses to get inside cells and thereby causing infections [121]. In response to the bacterial and viral infections, innate immune system produces both pro and anti-inflammatory cytokines. Vitamin D improves cellular immunity by minimizing these abrupt increases in cytokines produced mainly by innate immunity [7]. Vitamin D supplementation also stimulates antioxidant genes expression including glutathione reductase and glutamate–cysteine ligase modifier subunit [122]. This increased secretion of glutathione further spares ascorbic acid (vitamin C) which got significant antimicrobial properties [123] and is suggested to be used in prevention and treatment of COVID-19 [124]. SARS-CoV-2 attached to host cells through ACE2R and vitamin D might reduce ACE2R. Recent studies have established that RAS is a major target of vitamin D. Inverse relationship between circulating 25(OH)D levels (a physiological quantifiable form) and plasma renin activity in hypertensive subjects was reported more than two decades ago; however, significance of the same was recognized only after discovery that 1,25(OH)2 D3 is a negative endocrine regulator of renin production [125].

**Other drugs and their targets**

Further, a broad range of antiviral drugs have also been used in various clinical trials against COVID-19 [126]. RNA dependant RNA polymerase, a crucial protease required during viral RNA replication is also an important therapeutic target. Remdesivir, a broad-spectrum antiviral drug initially used for Ebola virus treatment is an analogue adenosine. It effectively blocks viral infection and replication in vitro and in animals and has shown some promising effect against SARS-CoV-2 [127]. Favipiravir, purine nucleic acid analogue initially used for the treatment of influenza is an effective RNA dependant RNA polymerase inhibitor [128]. In addition to above, combination of lopinavir and ritonavir has also been tested for the treatment of COVID-19 infected patients with little clinical significance [129].

Another potential drug is chloroquine, an anti-malarial and anti-parasitic drug, is a promising anti-viral known for neutralising the acidic endosomal pH and thereby blocking the endosomal mediated viral entry [130]. Chloroquine demonstrates potent anti-viral effect by inhibiting SARS-CoV-2 in-vitro with a 90% effective concentration of 6.9 μM. Hydroxychloroquine, an analogue of chloroquine is a less toxic and more potent drug. Chloroquine exhibits potent anti-inflammatory and immuno modulatory effects which might play an important role in reduction of COVID-19 infection probably by hindering cytokine storm [131]. Combination of chloroquine with azithromycin has provided some clinical benefits in COVID-19 infection [132].

Convalescent plasma therapy has long been given to patients infected with deadly viruses like SARS-CoV, H1N1, Spanish flu, Ebola and the MERS. The techniques involved identification of patients who have already
infected with COVID-19 and have completely recovered 14 days prior to be successful donors. Further, they must have sufficient titer of neutralizing antibodies [133]. Various clinical trials have highlighted that the therapy significantly improved the patient’s clinical outcomes and were able to successfully clear the infection and infected cells by activating complement system and phagocytosis [134, 135].

Cytokine storm is one of the characteristic phenomena of SARS-CoV-2 infected patients. Initial evidences revealed that targeting anti-inflammatory molecules such as IL-6, IL-1R and TNF-α might be important in reducing the inflammation and ultimately inflammatory response. Currently, clinical trials have shown that tocilizumab (anti-IL-6), a humanized antibody significantly improves the clinical outcome of COVID-19 cases [4]. Similarly, Anakinra (anti-IL-1R) is also important in decreasing the inflammation [136].

Corticosteroids are well known anti-inflammatory molecules are effective for the treatment of a variety of inflammatory diseases [137]. They behaved like a double edged sword as on one hand they reduce inflammation while on the other hand they inhibit immune response thereby delaying the infection clearance [137]. In COVID-19, various clinical trials have outlined the role of corticosteroids such as one retrospective observational study revealed that 72% of ICU patients with COVID-19 are receiving glucocorticoids treatment [48]. Further, another study revealed that treatment of COVID-19 patients having ARDS, with steroids significantly decreased the mortality rate compared to those that did not receive steroids (46% vs 68%) [100].

Mesenchymal stem cells (MSCs) are known to play important roles in immune modulation either by secreting cytokines or directly interacting with target cells. MSCs are available at multiple sources including blood, bone, adipose tissue, placenta, umbilical cord etc. [138]. A clinical trial in COVID-19 patients (n=17) revealed that the treatment cleared inflammatory cytokines secreting T cells population whereas dendritic cell percentage and IL-10 levels were increased. Further, absence of ACE2 and TMPRSS2 in the MSCs further highlights their importance in the COVID-19 treatment [139].

Recently, studies are focussing on curbing the complement system pathway, a component of innate immune system, in reducing hyperinflammatory and

### Table 4 Ongoing clinical trials of therapeutic drugs for SARS-CoV-2

| Drugs                                      | Role                                                                 | References |
|--------------------------------------------|---------------------------------------------------------------------|------------|
| **Antiviral Medications (Blockage of the virus entry)** |                                                                     |            |
| Chloroquine & Hydroxychloroquine           | Block virus infection by increasing endosomal pH                     | [144]      |
|                                            | Reduce the viral copy number of SARS-CoV-2                           | [132]      |
|                                            | Combination of hydroxychloroquine and azithromycin demonstrated positive effect |            |
| **Viral RNA synthesis inhibitors**        |                                                                     |            |
| Remdesivir                                 | Antiviral activities against the single stranded RNA viruses         | [145]      |
|                                            | Inhibit the infection and pathology of MERS-CoV and SARS-CoV-II      | [146]      |
| Favipiravir                                 | Functions as an inhibitor of the RNA-dependent RNA polymerase by structurally resembling the endogenous guanine | [147]      |
| Lopinavir—ritonavir combination            | Inhibiting HIV-1 protease for protein cleavage, resulting in non-infectious, immature viral particles | [148]      |
|                                            |                                                                     | [149]      |
|                                            |                                                                     | [150]      |
| Ivermectin                                 | Anti-parasitic agent                                                 | [151]      |
|                                            | Reduce viral RNA up to 5000-fold after 48 h of infection with SARS-CoV-2 | [152]      |
|                                            |                                                                     | [153]      |
| Darunavir                                   | Anti-retroviral protease inhibitor uses in combination drugs such as ritonavir or cobicistat | [154]      |
|                                            |                                                                     | [155]      |
| **Blocking Virus–Cell Membrane Fusion**    |                                                                     |            |
| Recombinant Human Angiotensin-converting Enzyme 2 (rhACE2) | Blocking the S protein of SARS-CoV2 from interacting with the cellular ACE2 | [156]      |
|                                            | rhACE2 could inhibit SARS-CoV-2 replication in cellular and embryonic stem |            |
|                                            | decreased serum level of angiotensin II                             |            |
| Arbidol Hydrochloride (Umifenovir)         | Entry inhibitor against influenza viruses and arboviruses           | [157]      |
|                                            |                                                                     | [158]      |
hypercoagulation stage in severe COVID-19 patients [140]. Furthermore, Ibrahim et al. (2020) reported that intravenous injection of N-acetyl cysteine in small cases of COVID-19, demonstrated significantly decreased inflammation and clinical improvement along with markedly reduced CRP levels in all patients and ferritin in 9/10 patients [141]. Lastly, other options such as Imatinib a tyrosine kinase inhibitor and colchicine, an anti-inflammatory drug have also been explored in few COVID-19 cases [142, 143].

Present and Future Prospects of Ongoing Clinical Trial

Currently, various clinical trials are ongoing to test specificity and efficacy of vaccines and antibodies precisely targeting SARS-CoV-2. Multiple pharmaceutical drugs that have kept therapeutic promises in the treatment of COVID-19 includes human immunoglobulin, chloroquine, hydroxychloroquine, remdesivir, favipiravir, ritonavir, lopinavir, arbidol etc. The role of these therapeutic drugs shown in Table 4.

Conclusion

Current review addressed relevant information of SARS-CoV-2, incidence, etiopathogenesis, clinical characteristics with multiorgan involvements in associated comorbid diseases, present and future prospects of clinical trials to prevent, manage and control COVID-19 pandemic infection. Information depicted in present review may serve to fill existing lacunae in knowledge on pathogenesis of COVID-19. Moreover, aspects of SARS-CoV-2 etiologic-pathogenesis and replication depicted inside infected cells might help in development of present and future prospects of ongoing clinical trials. New updates poured reticent on deadly virus from all over world are considered in present review to enhance knowledge and clear out concepts for present future prospects of clinico-trials to ascertain success in the possible drugs development for COVID-19 treatment. Extensive research to recognize novel pathways and their cross talk to combat this virus in precarious settings is our future positive hope.

Compliance with Ethical Standards

Conflict of Interests Authors declare that they have no conflict of interest to disclose.

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