

6-1-2020

## COVID-19 clinical trials: A primer for the cardiovascular and cardio-oncology communities

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### Recommended Citation

Ky, Bonnie and Mann, Douglas L, "COVID-19 clinical trials: A primer for the cardiovascular and cardio-oncology communities." *JACC CardioOncology*. 2,2. . (2020).  
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## STATE-OF-THE-ART REVIEW

# COVID-19 Clinical Trials

## A Primer for the Cardiovascular and Cardio-Oncology Communities



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**ABSTRACT**

The coronavirus disease-2019 (COVID-19) pandemic has resulted in a proliferation of clinical trials designed to slow the spread of severe acute respiratory syndrome-coronavirus-2 (SARS-CoV-2). Many therapeutic agents that are being used to treat patients with COVID-19 are repurposed treatments for influenza, Ebola, or for malaria that were developed decades ago and are unlikely to be familiar to the cardiovascular and cardio-oncology communities. Here, we provide a foundation for cardiovascular and cardio-oncology physicians on the front line providing care to patients with COVID-19, so that they may better understand the emerging cardiovascular epidemiology and the biological rationale for the clinical trials that are ongoing for the treatment of patients with COVID-19. (J Am Coll Cardiol CardioOnc 2020;2:254-69)  
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**T**he coronavirus disease-2019 (COVID-19) pandemic has resulted in a proliferation of clinical trials that are designed to slow the spread of severe acute respiratory syndrome-coronavirus -2 (SARS-CoV-2), the virus that causes COVID-19. These therapies range from vaccines, to repurposed treatments for influenza, to drugs that were not effective in patients with Ebola, to treatments for malaria that were developed decades ago. Recognizing that patients with underlying cardiovascular risk factors, cardiovascular disease, or cancer have an increased risk for adverse outcomes with COVID-19, and recognizing that these vulnerable populations may be enrolled in COVID-19 clinical trials, here we present a critical review of the rationale for the different therapeutics that are currently being employed. As background, we first review the epidemiology of COVID-19, followed by the biology of CoV.

We then briefly define the complex interplay between the CoV and the renin-angiotensin system (RAS), which is directly relevant to the care of the majority of patients with cardiovascular disease or cancer who are receiving drugs that modulate this system. Finally, we review the mechanisms of action of the multiple therapies that are currently being studied in clinical trials. Given the breadth of information that is emerging, we will not discuss the role of vaccines.

**EPIDEMIOLOGY OF COVID-19**

The current impact of the novel CoV, SARS-CoV-2 is unquantifiable. The number of confirmed cases and deaths from the global COVID-19 pandemic increase daily (1,2). Although there is a great deal that still remains to be understood, initial reports from 552 hospitals in China describing 1,099 of the 7,736 patients infected with COVID-19 provide some insight

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This paper will co-publish in *JACC: Basic to Translational Science* and *JACC: CardioOncology*.

The authors attest they are in compliance with human studies committees and animal welfare regulations of the authors' institutions and Food and Drug Administration guidelines, including patient consent where appropriate. For more information, visit the *JACC: CardioOncology* [author instructions page](#).

Manuscript received April 13, 2020; accepted April 13, 2020.

## HIGHLIGHTS

- SARS-CoV-2, the virus that causes COVID-19, is a novel CoV that infects humans by binding to ACE2, which degrades angiotensin II, and hence plays a critical role in modulating the renin angiotensin system (RAS).
- The emerging epidemiology of COVID-19 suggests that patients with cardiovascular risk factors, including older age, cardiovascular disease, or cancer may be more susceptible to infection and suffer from worse clinical outcomes.
- Because of the limited understanding with respect to the interaction of RAS inhibitors and SARS-CoV-2 infectivity, we endorse current society recommendations to continue RAS antagonists for clinical indications for which these agents are known to be beneficial.
- Treatments for COVID-19 that are undergoing clinical trials range from therapies that block the entry of SARS-CoV-2 into host cells, to repurposed antiviral therapies such as protease inhibitors and nucleoside analogs that block viral replication by inhibiting viral RNA-dependent RNA polymerase.

into the disease (3). In this multicenter retrospective analysis, the majority were Wuhan residents or had contact with Wuhan residents, although 25.9% were neither. The median age of patients was 47 years (interquartile range [IQR]: 35 to 58 years), and 41.9% were female. Patients with more severe disease, compared with those with nonsevere disease, tended to be older and tended to suffer from at least 1 comorbidity. In this retrospective analysis, patients commonly received intravenous antibiotics (58.0%). Oseltamivir was administered in 35.8%, systemic steroids in 18.6%, and oxygen in 41.3% of patients. The median duration of hospitalization was 12.0 days (IQR: 10.0 to 14.0 days); however, the majority of the patients (93.6%) remained hospitalized at the time of data analysis and as such, the clinical course still largely remains to be defined.

## COVID-19 AND CARDIOVASCULAR COMPLICATIONS.

Epidemiologic data thus far suggest that patients with cardiovascular risk factors, including older age, cardiovascular disease, or cancer are more susceptible to

infection and suffer from worse clinical outcomes (4). COVID-19 can also directly result in a number of cardiovascular complications, including fulminant myocarditis, myocardial injury, heart failure, and arrhythmia (3,5,6). There have been a number of published case reports of clinically suspected myocarditis as suggested by: markedly elevated troponin levels, ST-segment elevation on electrocardiogram without obstructive coronary disease, severely decreased left ventricular systolic function, and shock (7), with cardiac magnetic resonance imaging evidence of diffuse myocardial edema and gadolinium enhancement (8). However, in another isolated autopsy report from a patient who suffered from SARS-CoV-2-related pneumonia and cardiac arrest, no obvious histological changes in the myocardium were observed with the exception of few interstitial mononuclear inflammatory infiltrates (9).

Elevated troponin levels have also been observed in those with worse clinical outcomes. In a retrospective, single-center analysis of 416 hospitalized patients with confirmed COVID-19, 19.7% displayed evidence of cardiac injury, as defined by elevated high-sensitivity troponin I levels greater than the 99th percentile upper limit. Those with confirmed cardiac injury tended to be older (median age of 74 vs. 60 years) and suffer from hypertension (59.8% vs. 23.4%), diabetes (24.4% vs. 12.0%), coronary heart disease (29.3% vs. 6.0%), heart failure (14.6% vs. 1.5%), or cancer (8.5% vs. 0.6%) (10).

## COVID-19 IN PATIENTS WITH CARDIOVASCULAR RISK FACTORS OR DISEASE.

Patients with cardiovascular risk factors or disease are at increased risk of suffering from worse clinical outcomes with COVID-19. In an analysis of 2 cohorts from Jinyintan Hospital and Wuhan Hospital of 191 patients, patients with hypertension, diabetes, or coronary heart disease were at increased risk of in-hospital mortality (11). The prevalence of hypertension among nonsurvivors was 48% as compared to 30% in survivors; 31% versus 19% for diabetes, and 13% versus 8% for cardiovascular disease. These comorbidities were also more likely to be present in patients who required intensive care unit admission (4). Other studies, including a recently published meta-analysis of 46,248 infected patients, have corroborated the observation that patients with cardiovascular risk factors or cardiovascular disease have worse clinical outcomes (12) and also suggest that hypertension (17 ± 7%; 95%

## ABBREVIATIONS AND ACRONYMS

**ACE** = angiotensin-converting enzyme

**AT1R** = angiotensin II type 1 receptor

**CI** = confidence interval

**CoV** = coronavirus

**COVID-19** = coronavirus disease-2019

**FDA** = Food and Drug Administration

**IFN** = interferon

**IL** = interleukin

**IQR** = interquartile range

**MERS** = Middle East respiratory syndrome

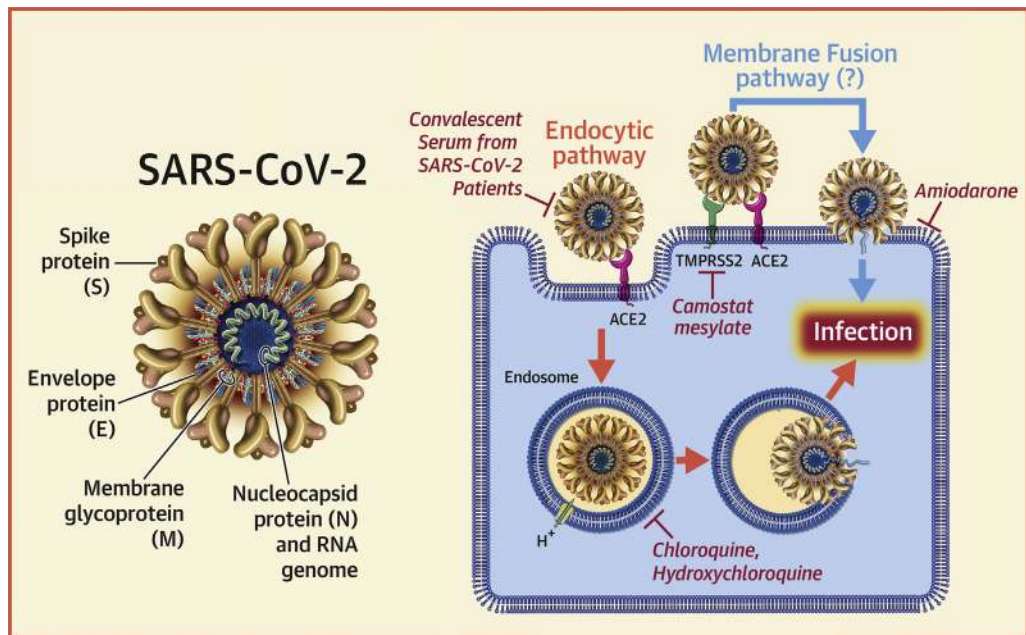
**RAS** = renin-angiotensin system

**RNA** = ribonucleic acid

**sACE2** = soluble angiotensin-converting enzyme 2

**SARS-CoV-2** = severe acute respiratory syndrome-coronavirus-2

**TMPRSS2** = transmembrane protease serine 2

**CENTRAL ILLUSTRATION SARS-CoV-2 Structure and Entry Into Cells**

Ky, B. et al. *J Am Coll Cardiol CardioOnc.* 2020;2(2):254-69.

The severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2) virus genome encodes 4 major structural proteins: the spike (S) protein; the nucleocapsid (N) protein; the membrane (M) protein; and the envelope (E) protein. The S protein is responsible for facilitating entry of the CoV into the target cell. The routes employed by SARS-CoV include endocytosis and membrane fusion. The route employed by SARS-CoV-2 is via endocytosis; whether SARS-CoV-2 enters cells by membrane fusion is not known. Binding of the S protein of SARS-CoV to angiotensin-converting enzyme 2 (ACE2) leads to the uptake of the virions into endosomes, where the viral S protein is activated by the pH-dependent cysteine protease cathepsin L. Activation of the S protein by cathepsin L can be blocked by bafilomycin A1 and ammonium chloride, which indirectly inhibit the activity of cathepsin L by interfering with endosomal acidification. Chloroquine and hydroxychloroquine are weak bases that diffuse into acidic cytoplasmic vesicles such as endosomes, lysosomes, or Golgi vesicles and thereby increases their pH. MDL28170 inhibits calpain and cathepsin L. SARS-CoV can also directly fuse with host cell membranes, after processing of the virus spike protein by transmembrane protease serine 2 (TMPRSS2), a type II cell membrane serine protease. Camostat mesylate is an orally active serine protease inhibitor. Modified from Simmons et al. (25). RNA = ribonucleic acid.

confidence interval [CI]: 14% to 22%), diabetes ( $8 \pm 6\%$ ; 95% CI: 6% to 11%), and cardiovascular disease ( $5 \pm 4\%$ ; 95% CI: 4% to 7%) were prevalent comorbidities among infected patients. Recent studies have also demonstrated that age and hypertension were predictors of an increased likelihood of cardiovascular complications, and cardiovascular complications were associated with a 4.26-fold increased risk of death (95% CI: 1.9 to 9.49) (10).

**COVID-19 IN PATIENTS WITH HEART TRANSPLANTATION.** There have been case series published on COVID-19 infection in heart transplant recipients. Two confirmed cases suggest similar presentations to nontransplant recipients and both patients demonstrated clinical improvement. A questionnaire of 87 heart transplant recipients in China, of which

importantly 96.6% undertook quarantine procedures, did not suggest a markedly elevated rate of SARS-CoV-2 infection in this population (13,14).

**COVID-19 IN PATIENTS WITH CANCER.** In a retrospective medical review of 1,524 patients with cancer who were admitted to the Department of Radiation and Medical Oncology in Zhongnan Hospital of Wuhan University from December 30, 2019, to February 17, 2020, the infection rate of SARS-CoV-2 in patients with cancer was 0.79% (95% CI: 0.3% to 1.2%) (15). In contrast, the estimated cumulative incidence of all COVID-19 cases in Wuhan was 0.37%. As a result, the odds of infection in patients with cancer were estimated to be 2.31 (95% CI: 1.89 to 3.02) greater. Patients with cancer who were infected had a median age of 66 years and were more likely to have

**TABLE 1 Select Treatment Trials Targeting RAS**

Drug Name	Mechanism of Action	NCT Number	Title	Study Population	Targeted Enrollment	Study Design	Primary Outcome Measure
Losartan	Anti-RAS	<a href="#">NCT04312009</a>	Losartan for Patients With COVID-19 Requiring Hospitalization	Age ≥18 yrs with presumptive positive laboratory test for SARS-CoV-2; admission to the hospital with a sequential organ failure assessment score ≥1 and increased oxygen requirement; randomization within 24 h of presentation	200	Randomized, double-blind, placebo controlled	Difference in oxygenation status at 7 days
Losartan	Anti-RAS	<a href="#">NCT04311177</a>	Losartan for Patients With COVID-19 Not Requiring Hospitalization	Age ≥18 yrs with presumptive positive laboratory test for SARS-CoV-2 or URI or fever	516	Randomized, double-blind, placebo controlled	Hospital admission up to 15 days

For an up-to-date listing of trials, search for "COVID-19" at the ClinicalTrials.gov website.  
 COVID-19 = coronavirus disease-2019; NCT = national clinical trial; OFA = organ failure assessment; RAS = renin-angiotensin system; SARS-CoV-2 = severe acute respiratory syndrome coronavirus-2; URI = upper respiratory infection.

non-small cell lung cancer (58.3%). Five of these patients were being treated with chemotherapy, immunotherapy, or radiation therapy. Three deaths were recorded.

In a multicenter, prospective cohort study of 2,007 cases from 575 hospitals, 1% of the 1,590 COVID-19 cases had a history of cancer (15). This in contrast to an incidence of cancer in the Chinese population of 0.29% per 100,000 people. Again, among those infected, lung cancer was most common, and patients tended to be older. Patients with cancer also suffered from an increased risk of adverse events that tended to occur earlier, including admission to the intensive care unit, need for invasive ventilation, or death, which occurred in 7 of 18 patients (39%), compared with 124 of 1,572 patients without cancer (8%). Patients with cancer who were recently treated with chemotherapy or surgery were also more likely to suffer from clinically severe adverse events. However, there is a critical need for additional studies to validate these early observations.

### THE CoV FAMILY

CoVs represent a large family of hundreds of enveloped, single-stranded, positive-sense ribonucleic acid (RNA) viruses that establish an infection primarily by targeting the mucosal surfaces of respiratory and intestinal tracts of a wide range of mammals and birds. There are 4 main subgroupings of CoVs: alpha, beta, gamma, and delta (16). The 7 CoVs that are capable of infecting humans include 229E (alpha CoV), NL63 (alpha CoV), OC43 (beta CoV), HKU1 (beta CoV), Middle East respiratory syndrome (MERS)-CoV (beta CoV), SARS-CoV (beta CoV), and SARS-CoV-2 (beta CoV). The prototype human CoV isolates 229E and OC43 have been causally linked to the common cold. SARS-CoV is the cause of the SARS, whereas

MERS-CoV was established as the cause of MERS. Identification and sequencing of the virus responsible for COVID-19 established that it was a novel CoV that shared 88% sequence identity with 2 bat-derived SARS-like CoVs (16). Subsequently, the 2019 novel CoV was shown to share a 79.5% sequence homology with SARS-CoV and was subsequently renamed SARS-CoV-2 (16). The genome of the CoVs encodes 4 major structural proteins: the spike (S) protein, nucleocapsid protein, membrane protein, and the envelope protein (Central Illustration). The S protein is responsible for facilitating entry of the CoV into the target cell (16,17) and is composed of a short intracellular tail, a transmembrane anchor, and a large ectodomain that consists of a receptor binding S1 subunit and a membrane-fusing S2 subunit (16).

### CoV VIROLOGY

Given that far more is known with respect to the virology of SARS-CoV than of SARS-CoV-2, and given that these 2 CoVs appear to have some overlapping biology and clinical presentations, we will discuss these 2 viruses together, with an emphasis on the most recent studies that have revealed unique biological aspects of SARS-CoV-2. We will review viral attachment, entry, and replication of SARS-CoV and SARS-CoV-2 in host cells. This discussion will be integrated with a review of the ongoing clinical trials that target these different aspects of the biology of SARS-CoV-2 (Tables 1 to 5).

Angiotensin-converting enzyme 2 (ACE2) is the entry receptor for SARS-CoV and SARS-CoV-2. Viruses enter cells by binding to host cell-encoded proteins that facilitate the entry of the virus into the cell, as well as allow the virus to survive and replicate within the cell. Some viruses, including certain strains of CoVs are capable of down-modulating the entry

**TABLE 2** Select Treatment and Prophylaxis Trials Targeting Viral Cell Entry

Drug Name	Mechanism of Action	NCT Number	Title	Study Population	Targeted Enrollment	Study Design	Primary Outcome Measure
<b>Treatment Trials</b>							
Camostat	Viral entry	<a href="#">NCT04321096</a>	The Impact of Camostat Mesylate on COVID-19 Infection (CamoCo-19)	Age 18-110 yrs, COVID-19-confirmed hospitalized patients (<48 h) or if hospital-acquired COVID-19 is suspected, <48 h since onset of symptoms	180	Randomized, double-blind placebo controlled, phase IIa trial	Time to clinical improvement at 30 days
Hydroxychloroquine	Viral entry	<a href="#">NCT04315896</a>	Hydroxychloroquine Treatment for Severe COVID-19 Pulmonary Infection (HYDRA Trial)	Age 18-80 yrs, COVID-19 confirmed by RT-PCR in any respiratory sample; severe disease defined by pulse O <sub>2</sub> < 91%, 3% decline from baseline pulse O <sub>2</sub> , or need for increased supplemental O <sub>2</sub> , mechanical ventilation, or sepsis	500	Randomized, double-blind, placebo controlled	All-cause hospital mortality at 120 days
Hydroxychloroquine	Viral entry	<a href="#">NCT04316377</a>	Norwegian Coronavirus Disease 2019 Study (NO COVID-19)	Age >18 yrs, hospitalized, moderately severe disease (NEWS score ≤6); SARS-CoV-2-positive test	202	Randomized, open, single arm	Rate of decline in SARS-CoV viral load at 96 h
<b>Prophylaxis Trials</b>							
Chloroquine phosphate	Viral entry	<a href="#">NCT04303507</a>	Chloroquine/Hydroxychloroquine Prevention of Coronavirus Disease (COVID-19) in the Healthcare Setting (COPCOV)	Age ≥16 yrs; health care worker or front-line participant with patient contact working in a health care facility; inpatient or relative of a patient and likely exposed to COVID-19; agree to not self-medicate with potential antivirals	40,000	Randomized, double-blind, placebo controlled	Number of symptomatic COVID-19 infections Severity of symptoms
Hydroxychloroquine	Viral entry	<a href="#">NCT04308668</a>	Post-exposure Prophylaxis/Pre-emptive Therapy for SARS-Coronavirus-2 (COVID-19 PEP)	Age >18 yrs; exposure to a COVID-19 case within 4 days as either a health care worker or household contact; symptomatic COVID-19 case with confirmed diagnosis within 4 days of symptom onset; or symptomatic health care worker with known COVID-19 contact and within 4 days of symptom onset	3,000	Randomized, double-blind, placebo controlled	Incidence of COVID-19 disease at 14 days Ordinal Scale of COVID-19 disease severity at 14 days
Hydroxychloroquine	Viral entry	<a href="#">NCT04318444</a>	Hydroxychloroquine Post Exposure Prophylaxis for Coronavirus Disease (COVID-19)	Age >18 yrs; household contact of index case: currently residing in the same household as an individual evaluated at NYP via outpatient, ED, or inpatient services who: 1) tests positive for COVID-19; or 2) is defined as suspected case, or PUI, by the treating physician	1,600	Randomized, double-blind, placebo controlled	Symptomatic, lab-confirmed COVID-19
Hydroxychloroquine	Viral entry	<a href="#">NCT04318015</a>	Hydroxychloroquine Chemoprophylaxis in Healthcare Personnel in Contact With COVID-19 Patients (PHYDRA Trial)	Age >18 yrs; health care personnel exposed to patients with COVID-19 respiratory disease (physicians, nurses, chemists, pharmacists, janitors, stretcher-bearer, administrative, and respiratory therapists)	400	Randomized, double-blind, placebo controlled	Symptomatic COVID-19 infection rate at 60 days
<p>For an up-to-date listing of trials, search for "COVID-19" at the <a href="https://clinicaltrials.gov">ClinicalTrials.gov</a> website.</p> <p>ED = emergency department; NEWS = National Early Warning Score; NYP = New York Presbyterian; PUI = person under investigation; RT-PCR = reverse transcriptase-polymerase chain reaction; other abbreviations as in <a href="#">Table 1</a>.</p>							

**TABLE 3 Select Treatment Trials Targeting Viral Replication**

Drug Name	Mechanism of Action	NCT Number	Title	Study Population	Targeted Enrollment	Study Design	Primary Outcome Measure
Umifenovir	Antiretroviral	NCT04260594	Clinical Study of Arbidol Hydrochloride Tablets in the Treatment of Pneumonia Caused by Novel Coronavirus	Age ≥18 yrs; subjects with pneumonia diagnosed as 2019-nCoV infection; detection of 2019-nCoV nucleic acid-positive by RT-PCR in respiratory tract or blood samples; virus gene sequence of respiratory tract or blood samples is highly homologous to the known 2019-nCoV	380	Randomized, single-arm, open-label umifenovir	Negative viral conversion rate at 7 days
ASCO9 + ritonavir; lopinavir + ritonavir	Antiretroviral	NCT04261907	Evaluating and Comparing the Safety and Efficiency of ASCO9/Ritonavir and Lopinavir/Ritonavir for Novel Coronavirus Infection	Age between 18 to 75 yrs; lab (RT-PCR) and clinically confirmed case of 2019-nCoV pneumonia; hospitalized with a new onset respiratory illness (≤7 days since illness onset)	160	Randomized, open-label ASCO9/ritonavir or lopinavir/ritonavir	The incidence of adverse outcomes, defined by at least 1 of the following: pulse O <sub>2</sub> ≤93% without O <sub>2</sub> supplementation, PaO <sub>2</sub> -to-FiO <sub>2</sub> ratio ≤300 or RR ≥30 breaths/min assessed at 14 days
Darunavir + cobicistat	Antiretroviral	NCT04252274	Efficacy and Safety of Darunavir and Cobicistat for Treatment of COVID-19 (DC-COVID-19)	Pneumonia caused by 2019-nCoV	30	Randomized, open-label, single-arm	The viral clearance rate of throat swabs, sputum, or lower respiratory tract secretions at day 7
Lopinavir + ritonavir; umifenovir	Antiretroviral	NCT04252885	The Efficacy of Lopinavir Plus Ritonavir and Arbidol Against Novel Coronavirus Infection (ELACOI)	Age 18-80 yrs; confirmation of SARS-CoV-2 infection by RT-PCR with normal kidney and liver function	125	Randomized, open-label (1:1:1) to lopinavir + ritonavir; or umifenovir; or standard care	The rate of viral inhibition, as determined by RT-PCR at days 2, 4, 7, 10, 14, and 21
Lopinavir + ritonavir	Antiretroviral	NCT04330690	Treatments for COVID-19: Canadian Arm of the SOLIDARITY Trial (CATCO)	Age >6 months with confirmed SARS-CoV-2 by RT-PCR, admitted to hospital	440	Randomized, open-label (1:1) of lopinavir + ritonavir or standard care	Efficacy of intervention at 29 days as determined by 10-point ordinal scale of clinical status
Remdesivir	Antiretroviral	NCT04280705	Adaptive COVID-19 Treatment Trial (ACTT)	Age 18-99 yrs, PCR-confirmed novel coronavirus infection by lab assay; illness as defined by abnormal radiographic imaging, clinical assessment, and pulse O <sub>2</sub> ≤94%, requiring O <sub>2</sub> , or requiring mechanical ventilation	572	Adaptive, randomized, double-blind placebo controlled	Time to recovery at day 29
Remdesivir	Antiretroviral	NCT04292899	Study to Evaluate the Safety and Antiviral Activity of Remdesivir (GS-5734) in Participants With Severe Coronavirus Disease (COVID-19)	Age ≥18 yrs; confirmation of SARS-CoV-2 infection by RT-PCR ≤4 days before randomization; current hospitalization with pulse O <sub>2</sub> ≤ 94%	6,000	Randomized, open-label study of remdesivir 5 days; or remdesivir 10 days	Odds of clinical improvement on a 7-point ordinal scale by day 11
Remdesivir	Antiretroviral	NCT04292730	Study to Evaluate the Safety and Antiviral Activity of Remdesivir (GS-5734) in Participants With Moderate Coronavirus Disease (COVID-19) Compared to Standard of Care Treatment	Age ≥18 yrs; confirmation of SARS-CoV-2 infection by RT-PCR ≤4 days before randomization; current hospitalization with fever, pulse O <sub>2</sub> >94%, radiographic evidence of pulmonary infiltrates	1,600	Randomized, open-label study of remdesivir 5 days; or remdesivir 10 days; or standard of care	Odds of clinical improvement on a 7-point ordinal scale by day 11

For an up-to-date listing of trials, search for "COVID-19" at the ClinicalTrials.gov website.

ECMO = extracorporeal membrane oxygenation; FiO<sub>2</sub> = fraction of inspired O<sub>2</sub>; nCoV = novel coronavirus; PaO<sub>2</sub> = partial arterial O<sub>2</sub> pressure; PCR = polymerase chain reaction; RR = respiratory rate; other abbreviations as in Tables 1 and 2.

**TABLE 4** Select Treatment and Prophylaxis Trials Targeting the Immune System

Drug Name	Mechanism of Action	NCT Number	Title	Study Population	Targeted Enrollment	Study Design	Primary Outcome Measure
<b>Treatment Trials</b>							
IFN- $\alpha$ 1 $\beta$	Immunomodulatory	<a href="#">NCT04293887</a>	Efficacy and Safety of IFN- $\alpha$ 1 $\beta$ in the Treatment of Novel Coronavirus Patients	Age $\geq$ 18 yrs with clinically diagnosed coronavirus pneumonia within 7 days, including RT-PCR evidence of coronavirus and symptoms	328	Randomized, open-label, single-arm	Incidence of side effects within 14 days including dyspnea, pulse O <sub>2</sub> $\leq$ 94%, and RR $\geq$ 24 breaths/min
Methylprednisolone	Immunomodulatory	<a href="#">NCT04273321</a>	Efficacy and Safety of Corticosteroids in COVID-19	Age >18 yrs, diagnosis of novel coronavirus pneumonia (COVID-19)	400	Randomized, open-label, single-arm	Incidence of treatment failure in 14 days
Methylprednisolone	Immunomodulatory	<a href="#">NCT04244591</a>	Glucocorticoid Therapy for COVID-19 Critically Ill Patients With Severe Acute Respiratory Failure	Age >18 yrs, RT-PCR-confirmed infection, symptoms for >7 days, PaO <sub>2</sub> /FiO <sub>2</sub> <200, positive pressure ventilation or HFNC higher than 45 l/min for <48 h, requiring ICU admission	80	Randomized, open-label of glucocorticoid therapy or standard of care	Murray lung injury score at 7 days
Sarilumab	Immunomodulatory	<a href="#">NCT04315298</a>	Evaluation of the Efficacy and Safety of Sarilumab in Hospitalized Patients With COVID-19	Age $\geq$ 18 yrs; confirmation of SARS-CoV-2 infection by RT-PCR; current hospitalization with evidence of pneumonia and severe disease, critical disease, or multiorgan system dysfunction	400	Adaptive, randomized, double-blind, placebo-controlled with high and low doses	Percent change in C-reactive protein levels at 4 days Percentage of patients reporting clinical severity rated on a 7-point ordinal scale
Siltuximab	Immunomodulatory	<a href="#">NCT04329650</a>	Efficacy and Safety of Siltuximab vs. Corticosteroids in Hospitalized Patients With COVID-19 Pneumonia	Age $\geq$ 18 yrs; confirmation of SARS-CoV-2 infection by RT-PCR; current hospitalization with evidence of pneumonia; maximum O <sub>2</sub> support of 35%	100	Randomized, open-label of siltuximab or methylprednisolone	Proportion of patients requiring ICU admission at 29 days
Tocilizumab	Immunomodulatory	<a href="#">NCT04317092</a>	Tocilizumab in COVID-19 Pneumonia (TOCOVID-19)	No age or sex limit; SARS-CoV-2 infection by RT-PCR, current hospitalization secondary to pneumonia; pulse O <sub>2</sub> $\leq$ 93%, requiring O <sub>2</sub> , or requiring mechanical ventilation (invasive or noninvasive)	400	Open-label, single-arm	Mortality at 1 month
Tocilizumab	Immunomodulatory	<a href="#">NCT04320615</a>	A Study to Evaluate the Safety and Efficacy of Tocilizumab in Patients With Severe COVID-19 Pneumonia (COVACTA)	Age $\geq$ 18 yrs; hospitalized with COVID-19 pneumonia per WHO criteria; pulse O <sub>2</sub> $\leq$ 93% or PaO <sub>2</sub> /FiO <sub>2</sub> <300	330	Randomized, double-blind placebo controlled	Clinical status using a 7-category ordinal scale at 28 days
Anakinra, siltuximab, or tocilizumab	Immunomodulatory	<a href="#">NCT04330638</a>	Treatment of COVID-19 Patients With Anti-interleukin Drugs (COV-AID)	Age $\geq$ 18 yrs; hospitalized with confirmed COVID-19 diagnosis by RT-PCR or other laboratory test; hypoxia defined by PaO <sub>2</sub> /FiO <sub>2</sub> ; CXR or CT scan with bilateral infiltrates	342	Randomized, open-label (1:1:1) to anakinra, or siltuximab, or anakinra + siltuximab, or tocilizumab, or anakinra + tocilizumab	Time to clinical improvement at 15 days
<b>Prophylaxis Trial</b>							
Recombinant human IFN- $\alpha$ 1 $\beta$ and thymosin $\alpha$ 1	Immunomodulatory	<a href="#">NCT04320238</a>	Experimental Trial of rhIFN $\alpha$ Nasal Drops to Prevent 2019-nCoV in Medical Staff	Age 18 to 65 yrs, formally serving as medical staff in Taihe Hospital	2,944	2-arm, open-label to IFN- $\alpha$ 1 $\beta$ in a low-risk group and IFN- $\alpha$ 1 $\beta$ and thymosin $\alpha$ 1 in a high-risk group	New COVID-19 diagnosis at 28 days

For an up-to-date listing of trials, search for "COVID-19" at the ClinicalTrials.gov website.

CT = computed tomography; CXR = chest x-ray; HFNC = high flow nasal cannula; ICU = intensive care unit; IFN = interferon; rhIFN = recombinant human interferon; WHO = World Health Organization; other abbreviations as in [Tables 1 to 3](#).



receptor once they gain access to the cell. Receptor down-modulation is a strategy broadly used by many viruses to escape the immune system, as well as establish the best environment for viral replication and spread (18). Receptor down-modulation may also disrupt many of the natural physiologic functions of the host cell, resulting in cell death leading to organ level dysfunction.

The entry receptor utilized by both SARS-CoV and SARS-CoV-2 is ACE2 (**Central Illustration**), which is a type I transmembrane carboxypeptidase with 40% homology to ACE. ACE plays a critical role in activation of the RAS, by processing angiotensin I (angiotensin 1-10) to angiotensin II (angiotensin 1-8), the major effector peptide of RAS, which mediates its effects through selective interactions with G-protein-coupled angiotensin II type 1 (AT1) and type 2 (AT2) receptors (19). ACE, however, has not been implicated in the entry of human CoVs into cells.

ACE2 is highly expressed in the mouth, tongue, and types I and II alveolar epithelial cells in the lungs. ACE2 is also abundantly expressed by cardiovascular endothelium, cardiac myocytes, cardiac fibroblasts, as well as epithelial cells of the kidney and testis. The major substrate of ACE2 is angiotensin II, which is cleaved to angiotensin 1-7 (**Figure 1**) and functions through association with the G-protein-coupled receptor Mas receptor. The ACE2-angiotensin (1-7)-Mas receptor axis is regarded as the counter-regulatory arm of the RAS by opposing the effects of the ACE-angiotensin II axis-AT1. Although the precise role of ACE2 is still being evaluated, studies have shown that ACE2 exerts protective effects in the pulmonary and the cardiovascular systems, where it serves to oppose the deleterious effects of RAS activation (20-22).

Infection with SARS-CoV and SARS-CoV-2 is triggered by binding of the S protein on the surface of the CoV to ACE2 that is expressed on the cell surface. The receptor binding domain of the S protein of SARS-CoV-2 is located on the S1 subunit, which undergoes a conformational change when it binds to ACE2, which facilitates viral attachment to the surface of target cells (17). Binding of SARS-CoV-2 to ACE2 can result in uptake of virions into endosomes (**Central Illustration**). Viral entry into the cell requires priming of the S protein by the serine protease transmembrane protease serine 2 (TMPRSS2), which cleaves the viral S protein at the S1/S2 and the S2' site and allows fusion of viral and cellular membranes (23). The S proteins of SARS-CoV-2 can also use pH-sensitive endosomal proteases (cathepsin B and L) for priming and entry into cells. Interestingly, the binding affinity of the SARS-CoV-2 S ectodomain to

ACE2 is 10- to 20-fold higher than the binding of the SARS-CoV ectodomain to ACE2 (17). The increase in stickiness of the SARS-CoV-2 capsid S protein makes disease transmission more likely and might explain the increased person-to-person transmission with SARS-CoV-2 compared with that of SARS-CoV. Insofar as the viral S proteins are the part of the virus that interacts with the immune system, they may serve as a promising target for vaccines. Relevant to this discussion, convalescent sera from patients with SARS have been shown to block the entry of SARS-CoV-2 entry into cultured cells, albeit with less efficiency than SARS-CoV (23). However, monoclonal antibodies raised against the receptor binding domain of the S1 protein of SARS-CoV do not bind to the receptor binding domain of the S1 protein of SARS-CoV-2, suggesting that SARS-directed antibodies are not cross reactive and that SARS-CoV-2 proteins are necessary to develop effective antibodies. Although ACE inhibitors do not inhibit ACE2, Hoffman et al. (23) demonstrated that anti-ACE2 antibody prevented entry of viral vectors into cell lines expressing the SARS-CoV-2 S protein.

**INTERACTION OF CoV WITH THE RAS.** An additional layer of complexity to understanding the pathophysiology of the SARS-CoV-2 in humans stems from the complexity of the interactions of CoVs with the RAS (**Figure 1**), as well as the widespread use of drugs that interfere with the RAS, including ACE inhibitors, angiotensin-receptor antagonists, or angiotensin receptor-neprilysin inhibitors. Each of these drugs has different effects on the expression of the various components of the RAS in different tissue beds. Here we will briefly discuss these important interactions, as well as their implications for the treatment of patients with COVID-19.

Previous studies have shown that SARS-CoV S proteins induce the expression of a cell surface metalloenzyme termed a disintegrin and metalloproteinase-17, which was originally described as the enzyme that cleaves membrane-bound tumor necrosis factor- $\alpha$  from the cell surface and allows it to circulate in the soluble form of tumor necrosis factor- $\alpha$  (24). As shown in **Figure 1**, activation of a disintegrin and metalloproteinase-17 results in the proteolytic cleavage of ACE2 (referred to as shedding) from the cell surface, with the release of the catalytically active soluble angiotensin-converting enzyme 2 (sACE2) ectodomains into the circulation (22,25). A decrease in ACE2 levels on the cell surface would be expected to result in a decrease in the levels of angiotensin 1-7 (cytoprotective) and a corresponding increase in tissue levels of angiotensin II (proinflammatory and profibrotic). The importance of SARS-CoV2-induced

**TABLE 5 Select Treatment Trials With Multiple Targets**

Drug Name	Mechanism of Action	NCT Number	Title
Lopinavir + ritonavir; ribavirin; IFN- $\beta$ 1b	Antiretroviral and immunomodulatory	<a href="#">NCT04276688</a>	Lopinavir/Ritonavir, Ribavirin and IFN-beta Combination for nCoV Treatment
Lopinavir + ritonavir; hydroxychloroquine	Antiretroviral and viral entry	<a href="#">NCT04307693</a>	Comparison of Lopinavir/Ritonavir or Hydroxychloroquine in Patients With Mild Coronavirus Disease (COVID-19)
Remdesivir + hydroxychloroquine; remdesivir; hydroxychloroquine	Antiretroviral and viral entry	<a href="#">NCT04321616</a>	Efficacy of Different Anti-viral Drugs in COVID-19 Infected Patients
Combinations of oseltamivir, chloroquine, darunavir, ritonavir, lopinavir, oseltamivir, favipiravir	Antiretroviral and viral entry	<a href="#">NCT04303299</a>	Various Combination of Protease Inhibitors, Oseltamivir, Favipiravir, and Hydroxychloroquine for Treatment of COVID19: A Randomized Control Trial (THDMS-COVID-19)
Favipiravir; chloroquine phosphate	Antiretroviral and viral entry	<a href="#">NCT04319900</a>	Clinical Trial of Favipiravir Tablets Combined With Chloroquine Phosphate in the Treatment of Novel Coronavirus Pneumonia
Remdesivir; lopinavir + ritonavir; IFN- $\beta$ 1a; hydroxychloroquine	Antiretroviral, viral entry, and immunomodulatory	<a href="#">NCT04315948</a>	Trial of Treatments for COVID-19 in Hospitalized Adults (DisCoVeRy)
Favipiravir + tocilizumab	Antiretroviral and immunomodulatory	<a href="#">NCT04310228</a>	Favipiravir Combined With Tocilizumab in the Treatment of Corona Virus Disease 2019

For an up-to-date listing of trials, search for "COVID-19" at the [ClinicalTrials.gov](#) website.  
IL = interleukin; other abbreviations as in [Tables 1 to 4](#).

*Continued on the next page*

down-regulation of cell surface ACE2 was demonstrated in experimental studies, wherein administration of recombinant human ACE2 protein, genetic deletion of the AT1 receptor, or administration of an AT1 receptor antagonist were shown to be protective in acute lung injury models (21,22). These and other observations have suggested that the use of AT1 receptor antagonists may be beneficial in patients with COVID-19 (26), and consistent with this, losartan is currently being tested in randomized, double-blind placebo controlled studies as a potential therapy in hospitalized infected patients (Table 1). Relevant to this discussion, the ACE inhibitors in clinical use do not directly affect ACE2 activity (27). The biological significance of circulating sACE2 is not known. Of note, sACE2 retains its ability to bind the S protein of SARS-CoV and was shown to prevent entry of SARS-CoV into cells in vitro (28). Thus, sACE2 may act as a decoy receptor that prevents SARS-CoV-2 from binding to ACE2 on the cell surface. APN01 is a human recombinant sACE2 that has been shown to block the

early stages of SARS-CoV-2 infections in cell culture and human tissue organoid cultures (29). APN01 has already undergone safety and tolerability testing in a phase II trial of healthy volunteers (NCT00886353), but at the time of this writing is not being tested clinically in patients with COVID-19.

The recognition that many patients with COVID-19 have underlying medical conditions that are treated with ACE inhibitors and AT1 receptor antagonists (30), coupled with the knowledge that higher urinary ACE2 levels have been observed in patients treated with AT1 receptor antagonists (26), has given rise to the concern that pharmacologic up-regulation of ACE2 by RAS inhibitors may influence the infectivity of SARS-CoV-2 in a patient population that is already at high risk for severe COVID-19 infection (31). However, as noted in a recent review (32) on this topic, the experimental and clinical data often yield conflicting results with respect to the role of ACE inhibitors and AT1 receptor antagonists on ACE2 levels in different pathophysiological contexts. These conflicting results

**TABLE 5 Continued**

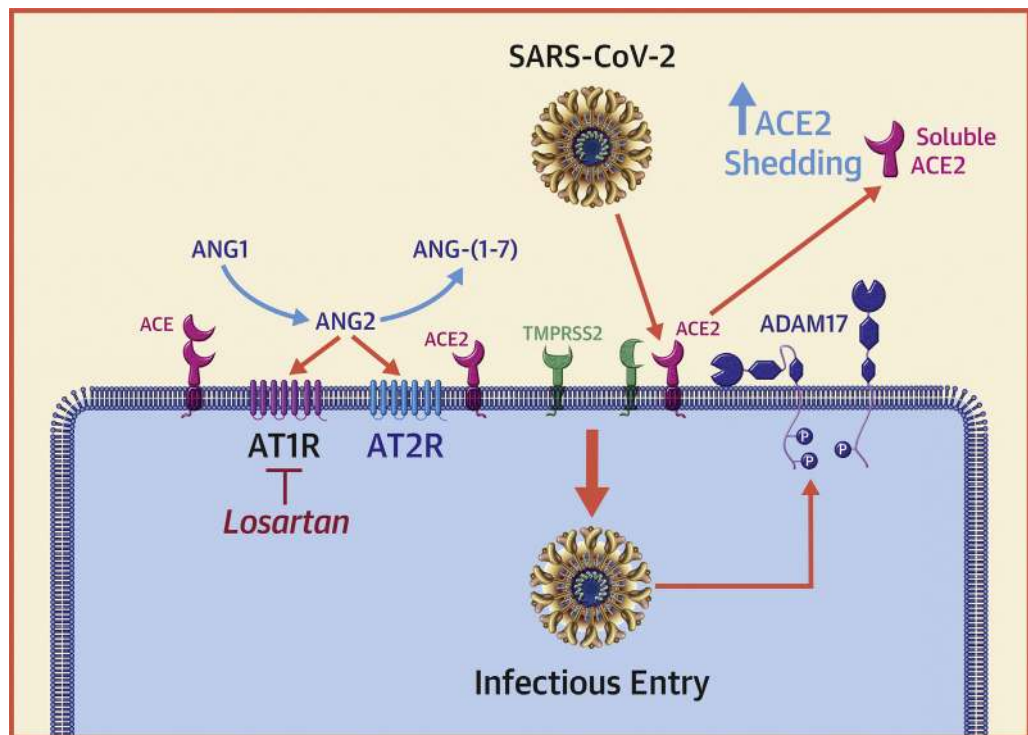
Study Population	Targeted Enrollment	Study Design	Primary Outcome Measure
Age ≥18 yrs hospitalized for virologically confirmed 2019-nCoV infection with NEWS ≥1 on recruitment; febrile with symptoms and duration of symptoms ≤10 days	127	Randomized, open-label (1:1:1) to lopinavir + ritonavir; or ribavirin; or IFN-β1b	Time to negative nasopharyngeal viral RT-PCR assessed up to 1 month
Age 16-99 yrs with confirmed mild COVID-19 (NEWS 0-4)	150	Randomized, open-label (1:1:1) lopinavir + ritonavir; or hydroxychloroquine; or standard care	Viral load at hospital days 3, 5, 7, 10, 14, 18
Age ≥18 yrs with confirmed SARS-CoV-2 by RT-PCR, admitted to hospital or ICU	700	Randomized, Open-Label (1:1:1) remdesivir; or hydroxychloroquine; or remdesivir + hydroxychloroquine adaptive controlled design; comparison with standard of care	In-hospital mortality at 3 weeks
Age 16-100 yrs with COVID-19 diagnosis	320	Randomized, open-label, oseltamivir + chloroquine; or darunavir + ritonavir + oseltamivir; or lopinavir + ritonavir + oseltamivir; or favipiravir + lopinavir + ritonavir; or darunavir + ritonavir + oseltamivir + chloroquine; or darunavir + ritonavir + favipiravir + chloroquine	Time to negative detection of SARS-CoV-2 in nasopharyngeal swab at 24 weeks
Age 18-75 yrs; diagnosed with nCoV pneumonia with a course of illness no more than 14 days; if the course is >14 days, no progression by chest radiograph within 7 days; respiratory symptoms; positive COVID-19 RT-PCR within 3 days	150	Randomized, double-blind 3-arm study of favipiravir + chloroquine; or favipiravir; or placebo	Time to improvement of respiratory symptoms Number of days of viral shedding Frequency of improvement of respiratory symptoms
Age ≥18 yrs, laboratory-confirmed SARS-CoV-2 infection as determined by RT-PCR or other assay <72 h prior to randomization, hospitalized patients with illness of any duration, and pulmonary exam abnormalities and pulse O <sub>2</sub> ≤ 94%, requiring O <sub>2</sub> , or requiring mechanical ventilation, or acute respiratory failure requiring mechanical ventilation and/or supplemental O <sub>2</sub>	3,100	Adaptive, randomized, open-label 1:1:1:1 to remdesivir; or lopinavir/ritonavir; or lopinavir/ritonavir + INF-β1α; or hydroxychloroquine; or standard of care	Percentage of subjects reporting disease severity on a 7-point ordinal clinical scale reflective of hospitalization and oxygenation status; or death at 15 days
Age 18-65 yrs, COVID-19 diagnosis, increased IL-6	150	Randomized, open-label (1:1:1) of favipiravir + tocilizumab; or favipiravir; or tocilizumab	Clinical cure rate at 3 months

suggest that the effects on RAS inhibitors on ACE2 are complex and nuanced and should not be assumed to be the same for all RAS inhibitors, nor should it be assumed that changes in ACE2 levels in the heart or other tissues necessarily reflect changes in ACE2 levels in the lung, which is the portal of entry for SARS-CoV-2. Given that we have limited understanding with respect to the interaction of RAS inhibitors, ACE2 levels, and SARS-CoV-2 infectivity in humans, we do not believe that it is possible to make definitive statements that go beyond the joint statement issued on March 17, 2020, by the Heart Failure Society of America, American College of Cardiology, and American Heart Association, that recommended “continuing renin-angiotensin-aldosterone system antagonists for those patients who are currently prescribed such agents for indications for which these agents are known to be beneficial” (33).

**ENTRY OF SARS-CoV AND SARS-CoV-2 INTO CELLS.** The entry of enveloped viruses into host cells occurs through 2 primary mechanisms: the first is direct fusion of the viral membrane with the plasma membrane of the host cells, which allows the virus to directly deliver its genomic material into the cytosol; and the second is that the virus hijacks the cell’s

endocytic machinery, by binding to a cell surface receptor, which then triggers endocytosis of the virus-receptor complex (**Central Illustration**). In the endocytic pathway, the endocytosed virions are subjected to an activation step within the endosome, which is typically mediated by the acidic environment of the endosome, resulting in fusion of the viral and endosomal membranes, which allows for the release of the viral genome into the cytosol. Several viruses, including human immunodeficiency virus and SARS-CoV use direct membrane fusions at the cell surface or endocytosis to enter cells. As noted, recent studies suggest that SARS-CoV-2 binds to ACE2, which leads to endocytosis of the receptor-virus complex (23). What is not known at this time is whether SARS-CoV-2 is also capable of directly fusing with the lipid membrane of cells. However, based on the similarities of how SARS-CoV and SARS-CoV-2 behave, it is likely that their modes of entry into cells will be similar. Understanding these differences in cell entry has implications for developing novel therapeutics.

**THERAPEUTICS TARGETING ENDOCYTOSIS.** The entry of SARS-CoV into cells was shown to occur by

**FIGURE 1** Interaction of CoVs With the RAS

Angiotensin-converting enzyme (ACE) converts angiotensin I (ANG1) (angiotensin 1-10) to angiotensin II (ANG2) (angiotensin 1-8), which is the major effector peptide of the renin-angiotensin system (RAS). ANG2 mediates its effects through selective interactions with G-protein-coupled angiotensin II type 1 receptor (AT1R) and G-protein-coupled angiotensin II type 2 receptor (AT2R). ANG2 is degraded to ANG-(1-7) by angiotensin-converting enzyme 2 (ACE2), ANG-(1-7) binds to the Mas receptor (not shown). The ACE2-ANG-(1-7)-Mas receptor axis opposes the effects of ACE-ANG2-AT1 axis. The binding of the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) spike protein to ACE2 induces ACE2 shedding by activating a disintegrin and metalloproteinase-17 (ADAM-17). A decrease in ACE2 levels would be expected to result in a decrease in the levels ANG-(1-7) levels (cytoprotective) and a corresponding increase in tissue levels of ANG2 (pro-inflammatory and profibrotic). Transmembrane serine protease 2 (TMPRSS2), a type II cell membrane serine protease that activates the spike protein of SARS-CoV-2 and allows it to bind to ACE2. Modified from Simmons et al. (25). CoVs = coronaviruses; P = phosphorylation.

direct fusion of the viral membranes with the plasma membrane of the host cell (**Central Illustration**), through a process that requires processing of the viral S protein by TMPRSS2 at or near the cell surface. Processing of the S protein exposes the fusion peptide of the S protein that inserts into the cell membrane, which brings the envelope of the viral membrane into closer approximation with the membrane of the host cell, thereby facilitating fusion (34). At the time of this writing, the uptake of SARS-CoV-2 into cells has been shown to occur through endocytosis of the SARS-CoV-2-ACE2 complex, which also requires priming of the S protein by TMPRSS2. It is not known whether SARS-CoV-2 also enters through direct fusion.

Based on the evidence linking TMPRSS2-mediated SARS-CoV-2 activation to SARS-CoV-2 infectivity (23,35), the small molecule serine protease inhibitor camostat mesylate may also be an attractive target for

clinical trials with SARS-CoV-2 (**Table 2**). Camostat mesylate has already been shown to inhibit replication of influenza and parainfluenza viruses and to prevent the development of pneumonia and viral myocarditis in infected mice (36). Given that the SARS-CoV-2 S protein is activated by the pH-dependent cysteine protease cathepsin L, this processing step may be sensitive to inhibition with drugs that indirectly inhibit cathepsin L activity by interfering with endosomal acidification (e.g., bafilomycin A1) or by compounds that directly block the proteolytic activity of cathepsin L.

At the time of this writing, it has also been suggested that the antimalarial drugs chloroquine and hydroxychloroquine might exert a potent antiviral effect by virtue of their ability to increase endosomal pH. Inside cells, chloroquine and hydroxychloroquine are rapidly protonated and concentrated in

endosomes. The positive charge of the chloroquine increases the pH of the endosome, which prevents cathepsin-induced priming of the viral S protein. Both chloroquine and hydroxychloroquine decrease SARS-CoV-2 replication in cultured cells; however, hydroxychloroquine is more potent than chloroquine (37). In a small single-arm study of patients with confirmed COVID-19, treatment with hydroxychloroquine was associated with a significant difference in clearing of viral nasopharyngeal carriage of SARS-CoV-2 within 3 to 6 days when compared with that of untreated control subjects that were studied at 3 to 6 days. Azithromycin when added to hydroxychloroquine was significantly more efficient for virus elimination (38). However, both therapies can result in QT prolongation, and as such, caution needs to be exercised when using these therapies together. Chloroquine and hydroxychloroquine can also manifest in cardiotoxicity, including cardiomyopathy, both systolic and diastolic, atrioventricular block, and bundle branch block (39). Hydroxychloroquine will be used as one of the treatment arms in the World Health Organization (WHO) multinational SOLIDARITY (Efficacy of Different Antiviral Drugs in SARS-CoV-2) trial (40) and is also currently being investigated in a number of other studies (Tables 2 and 5). Interestingly, amiodarone, which is a cationic amphiphile, was shown to inhibit Ebola virus infection in vitro in target cells, using concentrations of amiodarone that overlapped those detected in the sera of patients treated for arrhythmias. Both amiodarone and its main metabolite, monodesethyl amiodarone, were shown to interfere with the fusion of the viral envelope with the endosomal membrane, thus blocking viral replication (41). Amiodarone has also been shown to inhibit SARS-CoV infection and spreading in vitro by altering the late compartments of the endocytic pathway by acting after the transit of the virus through endosomes (42).

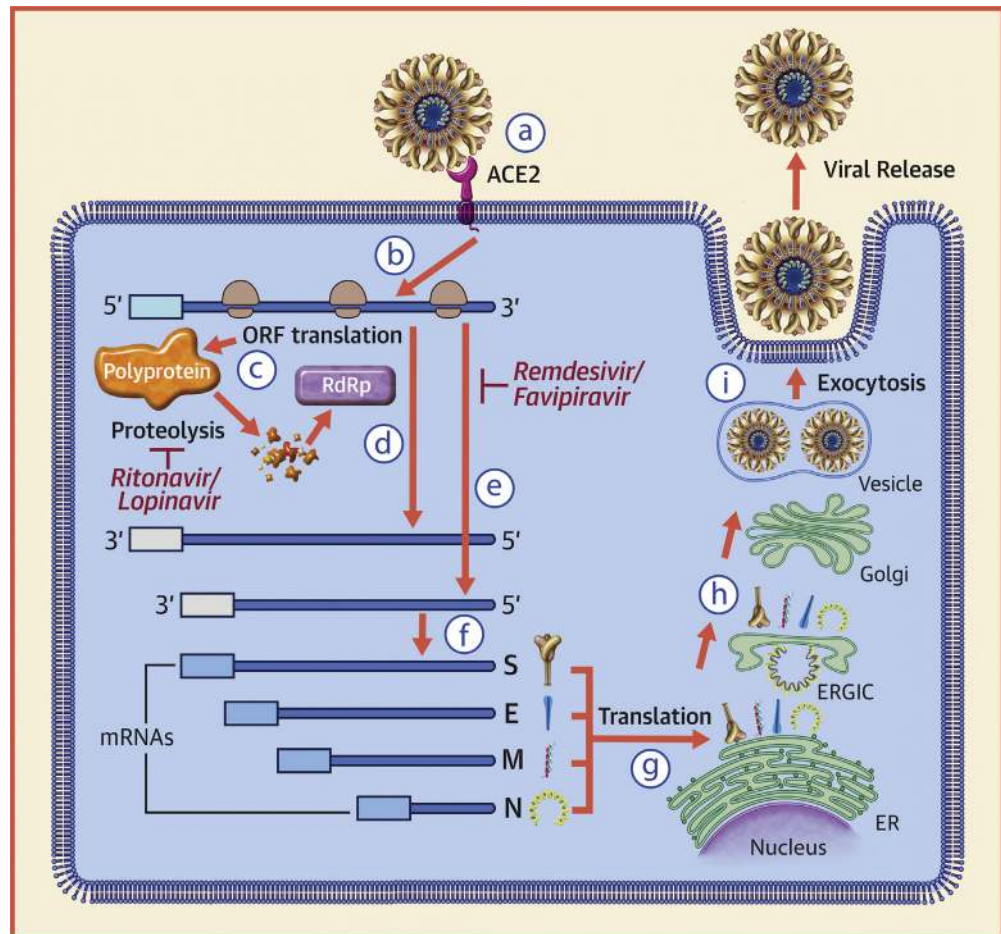
#### REPLICATION OF SARS-CoV IN HOST CELLS.

Because of the exceptionally large size of the CoV RNA genome (~30 kb) and the complexity of CoV-host cell interactions, coupled with the novelty of the SARS-CoV-2 genome, very little is known regarding SARS-CoV-2 replication in cells, let alone how the virus interacts with the host. Given that antiviral strategies are being considered for treatment of patients with COVID-19, here we will review what is generally understood about SARS-CoV replication in mammalian cells, recognizing that this information may change as we learn more about SARS-CoV-2 (see Figure 2).

Once the genomic RNA of SARS-CoV is released into the cytoplasm of the host cell, the positive-strand viral RNA is translated on host ribosomes into a large polypeptide termed the replicase, which undergoes proteolytic cleavage to yield proteins that are required from genome replication, including a viral RNA-dependent RNA polymerase. The viral RNA-dependent RNA polymerase generates a full-length, antisense negative-strand viral RNA template, which is used for replicating positive strand viral genomic RNA, as well as shorter subgenomic negative strand RNAs that serve as templates for synthesizing messenger RNAs that code for structural proteins of the virus, including the S, membrane, envelope, and nucleocapsid proteins. Translation of viral messenger RNAs occurs using the host endoplasmic reticulum. Once the viral structural proteins, S, envelope, and membrane, are translated in the endoplasmic reticulum, they move along the secretory pathway to the endoplasmic reticulum-Golgi intermediate compartment. There, the viral proteins become encapsulated and bud into membranes containing viral structural proteins. Following assembly and maturation, virions are transported to the cell surface in vesicles and released by exocytosis (43,44).

**THERAPEUTICS FOR VIRAL REPLICATION.** There are a number of antiviral drugs that are being repurposed for the treatment of SARS-CoV-2. A partial list of these antiviral drugs are discussed next.

**Nucleoside analogs.** Remdesivir (GS-5734, Gilead Sciences, Inc., Foster City, California) is a nucleoside analog that exhibits broad antiviral activity. Remdesivir is a prodrug that is metabolized to its active form GS-441524, which interferes with the action of viral RNA-dependent RNA polymerase, resulting in a decrease in viral RNA production. It is not known, however, whether remdesivir terminates RNA chains or causes mutations in them. Remdesivir was effective against multiple types of CoVs in cell culture and a mouse model of SARS (45); however, it did not show an effect in patients with Ebola. Remdesivir is currently being tested in several clinical trials for hospitalized patients with COVID-19 and pneumonia (Tables 3 and 5). Remdesivir is also 1 of the 4 treatment arms in the multinational SOLIDARITY trial, which is the World Health Organization's sponsored multinational randomized, open clinical trial to evaluate the safety and comparative efficacy of hydroxychloroquine, remdesivir, the combination of lopinavir and ritonavir, and the combination lopinavir and ritonavir plus interferon-beta (40). SOLIDARITY will use an adaptive design, which will allow

**FIGURE 2** The Replication Strategy of SARS-CoV

**(a)** The severe acute respiratory syndrome coronavirus (SARS-CoV) spike (S) glycoprotein attaches to the angiotensin-converting enzyme 2 (ACE2) receptor on the cell surface. On entering the cytoplasm, the viral core particle, which contains the positive (5' to 3') strand genomic ribonucleic acid (RNA), is released into the cytoplasm of the cell **(b)**. The positive-strand viral RNA is translated on host ribosomes to generate a large polyprotein **(c)** that undergoes proteolytic processing to generate multiple viral proteins, including an RNA-dependent RNA polymerase (RdRp). The RNA-dependent RNA polymerase generates a full-length, antisense negative-strand (3' to 5') viral RNA strand **(d)** that serves as template for replicating positive-strand viral genomic RNA, as well as shorter negative-strand RNAs **(e)** that serve as templates for synthesizing messenger ribonucleic acids (mRNAs) that code for structural proteins of the virus **(f)**, including the S, membrane (M), envelope (E), and nucleocapsid (N) proteins. Translation of viral mRNAs occurs using the host endoplasmic reticulum (ER) **(g)**. Once the viral structural proteins, S, E, and M, are translated and inserted into the ER, they move along the secretory pathway to the endoplasmic reticulum-Golgi intermediate compartment (ERGIC) **(h)**. The viral proteins become encapsulated and bud into membranes containing viral structural proteins, where mature virions are assembled. **(i)** Following assembly, virions are transported to the cell surface in vesicles and released by exocytosis. Modified from Turner et al. (43). ORF = open reading frame.

for discontinuation of drugs that lack effectiveness, as well as adding new drugs that appear promising. This type of trial design offers flexibility and efficiency, particularly in the identification of early signals related to either efficacy or toxicity, while maintaining study validity (46).

Favipiravir (Avigan, Fujifilm Toyama Chemical, Tokyo, Japan) is another nucleoside analog antiviral

drug that inhibits viral RNA-dependent RNA polymerase. Like remdesivir, it is a prodrug that is metabolized to its active form, favipiravir-ribofuranosyl-5'-triphosphate. Although favipiravir has undergone phase III clinical trials for the treatment of influenza, it is not yet approved by the U.S. Food and Drug Administration (FDA). Japan has granted approval for favipiravir for treating viral

strains unresponsive to current antivirals. In preliminary studies, favipiravir was shown to have more potent antiviral activity than lopinavir/ritonavir (47).

Ribavirin (Copegus, Genentech Inc., San Francisco, California) is a prodrug that acts as nucleoside inhibitor. The metabolites of ribavirin resemble adenosine or guanosine nucleosides that then become incorporated into viral RNA and inhibit RNA-dependent replication in RNA viruses. Ribavirin is currently FDA-approved for the treatment of chronic hepatitis C virus infection in combination with peginterferon alfa-2a (Pegasys, Genentech).

**Protease inhibitors.** Lopinavir is a protease inhibitor class that is used in fixed-dose combination with another protease inhibitor, ritonavir (lopinavir/ritonavir [Kaletra, AbbVie Inc., North Chicago, Illinois]) for the treatment of human immunodeficiency virus. Results from a randomized, open-label study of 199 hospitalized adult patients with confirmed SARS-CoV-2 infection assigned 1:1 to lopinavir 400 mg-ritonavir 100 mg twice daily for 14 days with standard of care or standard of care alone were recently published (48). All patients had an oxygen saturation of 94% on room air or a ratio of partial pressure of oxygen to the fraction of inspired oxygen <300 mg Hg. The primary endpoint was time to clinical improvement, where clinical improvement was defined based on an ordinal scale or survival from the hospital. The study was designed for 80% power with a 2-sided significance level of  $\alpha$  of 0.05 to detect an 8-day difference in median time to clinical improvement. Here, the median time to clinical improvement was 16.0 days (IQR: 13.0 to 17.0 days) in the lopinavir/ritonavir group compared with 16.0 days (IQR: 15.0 to 18.0 days) with standard care. The mortality at 28 days in the treatment group was similar to that observed in the standard care group (19.2% vs. 25%; difference: -5.8%; 95% CI: -17.3% to 5.7%), as was the detectable viral load. However, there were some suggestions of potential benefit with lopinavir/ritonavir with a shorter intensive care unit stay (median: 6 days vs. 11 days) and a shorter time to hospital discharge (median: 12 days vs. 14 days). As noted, the fixed-dose combination of lopinavir/ritonavir is 1 treatment arm in the SOLIDARITY trial (Table 3) (40).

**IMMUNOMODULATORY THERAPIES.** Interferons (IFNs) are cytokines that activate the innate immune system in response to viral infection. Type I interferons (IFN- $\alpha/\beta$ ) are synthesized by most cell types in the body response to a viral infection, whereas type II interferon (IFN- $\gamma$ ) is produced by

immune cells following antigen stimulation. Both type I and type II IFNs provoke the synthesis of proteins that have antiviral and immunomodulatory effects. Recombinant IFN- $\beta$  has been shown to inhibit SARS-CoV replication in vitro more effectively than either IFN- $\alpha$  or IFN- $\gamma$  (49,50). Interestingly, IFN- $\gamma$  down-regulates the expression of ACE2 on the cell surface and protects type I pneumocytes from SARS-CoV infection (51). The combination of lopinavir/ritonavir and IFN- $\beta$ 1b is being evaluated in the treatment of laboratory-confirmed MERS requiring hospitalization (52) and will also be evaluated in the SOLIDARITY trial (40).

A number of additional immunomodulatory agents are also currently being evaluated, including the interleukin (IL)-6 inhibitor, tocilizumab, and glucocorticosteroids (Tables 4 and 5), given the cytokine storm syndrome that has been observed in subgroups with severe COVID-19 (53) with increased levels of IL-2, IL-6, IL-7, and additional inflammatory cytokines (54). One meta-analysis suggested that the mean IL-6 levels were 2.9-fold (95% CI: 1.17 to 7.19-fold) greater in patients with complicated compared with noncomplicated COVID-19 (54). Tocilizumab (Actemra, Genentech) is FDA-approved for the treatment of severe cytokine release syndrome in patients treated with chimeric antigen receptor T-cell therapy and is also approved for the treatment of rheumatoid arthritis (55-58). Tocilizumab is a monoclonal antibody that binds the IL-6 receptor, both the membrane-bound and soluble forms, thus inhibiting both classic and trans-IL-6 downstream signaling. Similarly, the IL-6 humanized murine chimeric monoclonal antibody siltuximab, although not FDA-approved for the treatment of cytokine release syndrome, has also been used in the treatment of cytokine release syndrome and is also being studied as a potential therapy in severe COVID-19 infections. Siltuximab (Sylvant, Janssen Biotech Inc., Horsham, Pennsylvania) binds directly to IL-6 and prevents the activation of immune effector cells. Sarilumab (Kevzara, Sanofi US, Bridgewater, New Jersey) is a human monoclonal antibody against the IL-6 receptor that was developed for the treatment of rheumatoid arthritis that is also being evaluated for severe COVID-19.

There are no systematically obtained clinical data yet that support a benefit to the use of steroids, and some reports have suggested a possible detriment with delayed viral clearance and increased risk of infection with MERS and SARS, although the role of steroids in COVID-19 is an area of active investigation (Table 4) (59).

## COVID-19 AND CARDIOVASCULAR DISEASE

The COVID-19 pandemic has presented innumerable challenges to health care organizations and health care providers. Given that the vast majority of patients with cardiovascular disease are at high risk for SARS-CoV-2 infection, the cardiovascular and cardio-oncology communities will play a major role in caring for patients with COVID-19 now and for the foreseeable future. As a community, we have a long tradition of enrolling patients into clinical trials that evaluate therapeutic agents whose mechanisms of action are familiar, which facilitates reaching clinical equipoise when enrolling patients in clinical trials. In the coming months, our communities will be asked to contribute patients to clinical trials where the mechanisms of action of the therapeutic agents are less familiar and the knowledge base required for providing care for COVID-19 is accelerating at a dizzying pace. Here we have tried to provide a foundation for physicians who are on the front line of

providing care to patients with COVID-19, so that they can better understand the emerging cardiovascular epidemiology of COVID-19, as well as the biological rationale for the plethora of clinical trials that are either being designed or are currently recruiting patients.

**ACKNOWLEDGMENT** The authors would like to acknowledge Elizabeth Thompson for her assistance in reviewing the ClinicalTrials.gov website.

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**KEY WORDS** ACE2, clinical trials, COVID-19, renin angiotensin system, SARS-CoV-2