

Infection Control and Testing:
For healthcare workers performing aerosol-generating procedures* on patients with COVID-19 in the ICU, we recommend using fitted respirator masks (N95 respirators, FFP2, or equivalent) , as opposed to surgical/medical masks, in addition to other personal protective equipment (i.e., gloves, gown, and eye protection, such as a face shield or safety goggles)
We recommend performing aerosol-generating procedures on ICU patients with COVID-19 in a negative pressure room.
For intubated and mechanically ventilated adults with suspicion of COVID-19: For diagnostic testing, we suggest obtaining lower respiratory tract samples in preference to upper respiratory tract (nasopharyngeal or oropharyngeal) samples.
For intubated and mechanically ventilated adults with suspicion of COVID-19: With regard to lower respiratory samples, we suggest obtaining endotracheal aspirates in preference to bronchial wash or bronchoalveolar lavage samples.
Hemodynamics:
In adults with COVID-19 and shock , we suggest using dynamic parameters skin temperature, capillary refilling time, and/or serum lactate measurement over static parameters in order to assess fluid responsiveness.
For the acute resuscitation of adults with COVID-19 and shock , we suggest using a conservative over a liberal fluid strategy.
For the acute resuscitation of adults with COVID-19 and shock , we recommend using crystalloids over colloids.
For the acute resuscitation of adults with COVID-19 and shock , we suggest using buffered/balanced crystalloids over unbalanced crystalloids.
For adults with COVID-19 and shock , we suggest using norepinephrine as the first-line vasoactive agent, over other agents.
For adults with COVID-19 and shock , we suggest adding vasopressin as a second-line
For adults with COVID-19 and shock , we suggest titrating vasoactive agents to target a MAP of 60-65 mmHg, rather than higher MAP targets.
For adults with COVID-19 and shock with evidence of cardiac dysfunction and persistent hypoperfusion despite fluid resuscitation and norepinephrine , we suggest adding dobutamine, over increasing norepinephrine dose.

Ventilation
In adults with COVID-19, we suggest starting supplemental oxygen if the peripheral oxygen saturation (SPO ₂) is < 92%, and recommend starting supplemental oxygen if SPO ₂ is < 90%
In adults with COVID-19 and acute hypoxemic respiratory failure on oxygen , we recommend that SPO ₂ be maintained no higher than 96%.
For adults with COVID-19 and acute hypoxemic respiratory failure despite conventional oxygen therapy, we suggest using HFNC over conventional oxygen therapy.
In adults with COVID-19 and acute hypoxemic respiratory failure , we suggest using HFNC over NIPPV.
In adults with COVID-19 and acute hypoxemic respiratory failure , if HFNC is not available and there is no urgent indication for endotracheal intubation, we suggest a trial of NIPPV with close monitoring and short-interval assessment for worsening of respiratory failure.
In adults with COVID-19 receiving NIPPV or HFNC, we recommend close monitoring for worsening of respiratory status, and early intubation in a controlled setting if worsening occurs.
In mechanically ventilated adults with COVID-19 and ARDS, we recommend using low tidal volume (Vt) ventilation (Vt 4-8 mL/kg of predicted body weight), over higher tidal volumes (Vt>8 mL/kg).
For mechanically ventilated adults with COVID-19 and ARDS , we recommend targeting plateau pressures (Pplat) of < 30 cm H ₂ O.
For mechanically ventilated adults with COVID-19 and moderate to severe ARDS, we suggest using a higher PEEP strategy, over a lower PEEP strategy.
For mechanically ventilated adults with COVID-19 and moderate to severe ARDS , we suggest prone ventilation for 12 to 16 hours , over no prone ventilation.
For mechanically ventilated adults with COVID-19 and moderate to severe ARDS : We suggest using, as needed, intermittent boluses of neuromuscular blocking agents (NMBA), over continuous NMBA infusion, to facilitate protective lung ventilation.
For mechanically ventilated adults with COVID-19 and hypoxemia despite optimizing ventilation, we suggest using recruitment maneuvers, over not using recruitment maneuvers.

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Spread: Droplet spread, survives 2-3 hours on most surfaces, 2 days on smooth metal/plastic

Incubation: 2-14 days

1st week: Fever, cough, headache, fatigue, myalgias, pharyngitis

2nd week: Resolves in 80%, Viral pneumonia 20%

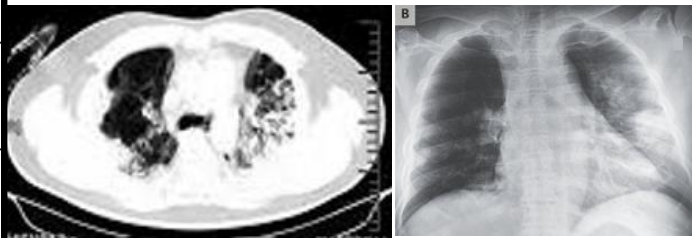
Risk increased: Heart/lung disease, immunosuppression, poorly controlled DM

Exam: Non specific

Labs: Lymphopenia with normal WBC count or relative leukopenia, Elevated Ferritin/CRP/D-Dimer is negative prognostic indicator

Mortality: Due to oxygenation failure or septic shock/multiorgan failure

Imaging:



Testing: CBC, CMP, ABG, Troponin, G6PD, Rapid flu testing and bacterial sputum and blood cultures (coinfection with BACTERIAL respiratory pathogens unlikely), Coronavirus PCR testing, CRP, Ferritin, D-Dimer

Treatment: Symptomatic support for stable patients otherwise refer to guidelines for critical care support -Currently under investigation (Plaquenil, Azithromycin and Remdesivir)