Ontario COVID-19 Drugs and Biologics Clinical Practice Guidelines Working Group

Therapeutic Management of Adult Patients with COVID-19

Recommendations apply to patients > 18 years of age. Recommendations are based on the best available data and may change as additional data becomes available. Science Briefs can be found on the Ontario COVID-19 Science Advisory Table website.



SEVERITY OF ILLNESS

RECOMMENDATIONS

Critically III Patients

Patients requiring ventilatory and/or circulatory support, including high-flow nasal oxygen, non-invasive ventilation, invasive mechanical ventilation, or ECMO.

These patients are usually managed in an intensive care setting.

- <u>Dexamethasone</u> 6 mg PO/IV daily for 10 days (or until discharge if sooner) is recommended for critically ill patients with suspected or confirmed COVID-19.
- <u>Tocilizumab</u> (dosed according to body weight) is recommended for critically ill patients with suspected or confirmed COVID-19, who are on recommended doses of dexamethasone therapy (or a dose-equivalent corticosteroid) AND are within 14 days of hospital admission (or within 14 days of a new COVID-19 diagnosis if the infection was nosocomially acquired).
 - A second dose of tocilizumab may be considered after 24 hours if the patient is not improving.
 - The dose of intravenous tocilizumab may be determined by a weight-based dose strategy (8 mg/kg, maximum dose 800 mg) OR by a weight-based dose banding strategy (800 mg if weight >90 kg; 600 mg if weight >65 and ≤90 kg; 400 mg if weight >40 and ≤65 kg; and 8 mg/kg if weight ≤40 kg).
- Prophylactic dose low molecular weight or unfractionate by scin is recommended in critically ill patients hospitalized with COVID-19.
- These patients **should not receive therapeutic domain and an area of the should not receive therapeutic domain and area of the should not receive therapeutic domain and area of the should not receive therapeutic domain and area of the should not receive the should not receiv**
- Remdesivir is not recommended antically ill patie with COVID-1 ceiving mechanical ventilation.
- In critically ill patients recording her flow oxygen (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, or nor passive mechanism (i.e., oxygen by mas' paxygen by high-flow nasal cannula, oxygen by mas' paxygen by high-flow nasal cannula, oxygen by high-flow nasal
- Bacterial co-infection is common in C. D-19 per conia at presentation.

 Defined dempiric antial tics for bacter, one conia unless bacterial infection is strong poected. Conting empiric antible as for no more than 5 days and de-escalate on the basic microbiologic hisults and clinical judgment.

Moderately III Patients

Patients newly requiring low-flow supplemental oxygen.

These patients are usually managed in hospital wards.

- Dexamethasone 6 mg PO/IV daily for 10 days (or until discharge if soone frechended for moderately ill patients with suspected or confirmed COVID-19.
- If patients are discharged with home-based oxygen therapy, developing assone 6 mg until oxygen is no longer required (for a maximum of 10 day and based on side red.)
- Remdesivir 200 mg IV on day 1, then 100 mg IV daily days is recommended for moderately ill patients with suspected or confirmed CO 19.
- ▲ Therapeutic dose anticoagulation may be unsidered over pohylactic dose ticoagulation in moderately ill patients where felt to be low risk of bleeping.
- All other patients show the phylac lose anticoagulion.

- Tocili nab (dosed according to body weight) is recommended for moderately ill patients with a sected or confirmed COVID-19, who have evidence of systemic inflammation, effine is a serum CRP of 75 mg/L or higher, AND have evidence of disease progression i.e., increasing oxygen or ventilatory requirements) despite 24-48 hours of recommended doses of dexamethasone therapy (or a dose-equivalent corticosteroid), AND are within 14 days of hospital admission (or within 14 days of a new COVID-19 diagnosis if the infection was nosocomially acquired).
- A second dose of tocilizumab may be considered after 24 hours if the patient is not improving, with dosing strategies being the same as for critically ill patients.

Mildly III Patients

Patients who do not require new or additional supplemental oxygen from their baseline status, intravenous fluids, or other physiological support.

These patients are usually managed in an ambulatory/ outpatient setting.

- <u>Dexamethasone</u> is <u>recommended</u> finildly ill patients with suspected or confirmed COVID-19.
- Remdesivir is not recommended in the patients with suspected or confirmed COVID-19.
- <u>Tocilizumab</u> is not recommended outside of clinical trials for mildly ill patients with suspected or confirmed COVID-19.
- There is currently insufficient evidence to make a recommendation around anticoagulation for mildly ill patients.
- The panel was unable to reach a consensus on the use of inhaled budesonide based on the available evidence. At this time, a recommendation cannot be made for its use to change disease course or serious disease outcomes. In selected patients with increased risk of adverse COVID-19 outcomes (≥65 years, or ≥50 years with one or more of: immunosuppression; heart disease; hypertension; asthma; lung disease; diabetes; liver disease; stroke; neurologic disease; or obesity, inhaled budesonide 800 mcg twice daily for 14 days may reduce patient-reported symptoms and time to recovery.

CURRENTLY NOT RECOMMENDED

There is insufficient evidence to support the use of the following therapies in the treatment of COVID-19 outside of clinical trials or where other indications would justify its use:

- ♦ Anti-SARS-CoV-2 monoclonal antibodies
- Colchicine
- Interferon (with or without lopinavir-ritonavir and ribavirin)
- Vitamin D

RECOMMENDED AGAINST

The following therapies are not recommended for the treatment of COVID-19 due to lack of benefit, potential harm, and system implications of overuse:

- Antibiotics (azithromycin)
- Hydroxychloroquine or chloroquine
- Ivermectin
- Lopinavir/ritonavir

Click here for dosing and pharmacologic considerations for medications approved or under investigation for COVID-19