MANAGEMENT OF ADULTS WITH MODERATE TO SEVERE COVID-19

NATIONAL COVID-19 CLINICAL EVIDENCE TASKFORCE

LEGEND

EBR: Evidence-Based Recommendation
CBR: Consensus-Based Recommendation
PP: Practice Point

Living guidance

Not prioritised for review

VERSION 13.0
PUBLISHED
10 SEPTEMBER 2020

Setting of care

ADMISSIONS

Manage people with likely or confirmed COVID-19 out of hospital where possible. **PP** [Taskforce]

Consider admission of people with likely or confirmed COVID-19 if they are haemodynamically unstable, hypoxaemic (SaO_2 on room air ≤ 92%), have comorbidities, or an unsuitable home environment. **PP** [Taskforce]

MANAGING RISK OF INFECTION

As per the current national guidance on the use of personal protective equipment (PPE) in hospitals during the COVID-19 outbreak:

- follow contact and droplet precautions for routine patient care of people with suspected or confirmed COVID-19
- add contact and airborne precautions when aerosol-generating procedures are required.

PP [Taskforce/AHPPC]

Definition of disease severity

Moderate illness
Stable adult patient presenting with respiratory and/or systemic symptoms or signs. Able to maintain oxygen saturation above 92% (or above 90% for patients with chronic lung disease) with up to 4 L/min oxygen via nasal prongs.

Characteristics:
- prostration, severe asthenia, fever > 38° C or persistent cough
- clinical or radiological signs of lung involvement
- no clinical or laboratory indicators of clinical severity or respiratory impairment

Severe illness
Adult patients meeting any of the following criteria:
- respiratory rate ≥ 30 breaths/min
- oxygen saturation ≤ 92% at a rest state
- arterial partial pressure of oxygen (PaO_2)/inspired oxygen fraction (FiO_2) ≤ 300

Testing and monitoring of inpatients

BASELINE TESTING AND DIAGNOSTIC WORK UP

In all people with suspected or confirmed COVID-19, perform haematology, biochemistry laboratory testing, a CXR and an ECG on admission. **PP** [Taskforce]

Investigate people with suspected or confirmed COVID-19 for influenza, CAP and other differential diagnoses as per usual practice. **PP** [Taskforce]

In cases of suspected COVID-19 that have not been confirmed by positive PCR, collect serum during the acute phase of the illness (preferably within the first 7 days of symptom onset); store and test the serum in parallel with convalescent sera collected 2 or more weeks after the onset of illness. **PP** [Taskforce/CDNA]

In cases where a strong clinical suspicion of COVID-19 remains after a negative SARS-CoV-2 PCR:
- continue isolation and treatment as for a provisional COVID-19 diagnosis;
- repeat SARS-CoV-2 PCR as soon as possible, adding a stool PCR if loose stool.

PP [Taskforce/ASID]

MONITORING AND MARKERS OF CLINICAL DETERIORATION

Monitoring — CONSENSUS RECOMMENDATION
For people with COVID-19, monitor markers of clinical progression, such as rapidly progressive respiratory failure and sepsis, especially on days 5 to 10 after onset of symptoms. **CBR** [Taskforce]

In all people with suspected or confirmed COVID-19, perform ECG and haematology and biochemistry laboratory tests as clinically indicated to monitor for complications, such as acute liver injury, acute kidney injury, acute cardiac injury or shock. **PP** [Taskforce]

Only repeat CXR in people with suspected or confirmed COVID-19 if clinically indicated (e.g. in cases of clinical deterioration or recent intubation). **PP** [Taskforce/ASID]

Do not routinely perform CT scanning in people with suspected or confirmed COVID-19. **PP** [Taskforce]

Supportive care in hospital

GENERAL

In all people with suspected or confirmed COVID-19, anticipate complications such as arrhythmias, cardiac impairment, sepsis and multi-organ dysfunction, and address using existing standards of care.
PP [Taskforce/ACEM]

Dexamethasone — RECOMMENDED
Use dexamethasone 6 mg daily intravenously or orally for up to 10 days in **adults with COVID-19 who are receiving oxygen** (including mechanically ventilated patients). **EBR** [Taskforce]

CONDITIONAL RECOMMENDATION AGAINST
Do not routinely use dexamethasone to treat **COVID-19 in adults who do not require oxygen**. **EBR** [Taskforce]

Dexamethasone or other corticosteroids may still be considered for other evidence-based indications in people who have COVID-19. **PP** [Taskforce]

SUPPORTIVE ANTI-INFECTIOUS THERAPY

In people with suspected or confirmed COVID-19 who are hypoxaemic (SaO_2 on room air ≤ 92%) or have pleural effusion or purulent sputum, prescribe antibiotics according to local pneumonia guidelines.

If the onset of bacterial pneumonia symptoms occurs within 72 hours of hospital admission, choose antibiotics according to local CAP guidelines.

If the onset of bacterial pneumonia symptoms occurs more than 72 hours after admission, choose antibiotics according to local HAP guidelines.
PP [Taskforce/ASID]

In people with suspected or confirmed COVID-19 with onset of symptoms < 48 hours, request an influenza PCR test.

If disease is severe, consider prescribing oseltamivir 75 mg BD (or a renally adjusted dose). If the influenza PCR is negative, cease oseltamivir.
PP [Taskforce/ASID]

Supportive care in hospital (cont.)

Remdesivir
CONDITIONAL RECOMMENDATION FOR

Whenever possible remdesivir should be administered in the context of a randomised trial with appropriate ethical approval. Use of remdesivir for adults with moderate, severe or critical COVID-19 outside of a trial setting may be considered. **EBR** [Taskforce]

Hydroxychloroquine
NOT RECOMMENDED

Do not use hydroxychloroquine for the treatment of COVID-19. **EBR** [Taskforce]

Disease-modifying treatments not recommended outside of clinical trials
NOT RECOMMENDED

For people with COVID-19, do not use the following disease-modifying treatments outside of randomised trials with appropriate ethical approval **EBR** [Taskforce]:

- Aprepitant
- Baloxavir marboxil
- Chloroquine
- Colchicine
- Convalescent plasma
- Darunavir-cobicistat
- Favipiravir
- Immunoglobulin plus methylprednisolone
- Interferon β-1a
- Interferon β-1b
- Interferon gamma
- Lopinavir-ritonavir
- Ruxolitinib
- Sofosbuvir-daclatasvir
- Telmisartan
- Umifenovir
- Other disease-modifying treatments

Trials are needed in special populations, including children and adolescents, pregnant and breastfeeding women, older people living with frailty and those receiving palliative care. Until further evidence is available, do not use other disease-modifying treatments in these populations unless they are eligible to be enrolled in trials. **PP** [Taskforce]

Escalation of care

HOSPITALS WITH ICU

Urgently refer people with suspected or confirmed COVID-19 to intensive care if they are haemodynamically unstable, have rapidly worsening tachypnoea or hypoxaemia, or require ≥ 40% FiO_2 to maintain SaO_2 ≥ 92% (or acceptable saturations in those with lower baselines). **PP** [Taskforce/ASID]

HOSPITALS WITHOUT ICU

Consider the need for early transfer of people with suspected or confirmed COVID-19 to a higher-level facility with an ICU. **PP** [Taskforce/ASID]

When preparing for transfer of people with suspected or confirmed COVID-19, consider infection control implications and whether intubation is required prior to transfer, as per local retrieval team policies. **PP** [Taskforce/ASID]

Discharge planning

People with suspected or confirmed COVID-19 who are clinically ready for hospital discharge should stay in home isolation after discharge until:

- at least 14 days have passed since onset of symptoms; AND
- there has been resolution of fever and respiratory symptoms of the acute illness for the previous 72 hours. **PP** [Taskforce/CDNA]

People with suspected or confirmed COVID-19 with an incomplete resolution of symptoms but who are clinically ready for hospital discharge should stay in home isolation after discharge until:

- at least 14 days have passed since onset of symptoms; AND
- there has been substantial improvement in symptoms of the acute illness (including resolution of fever for the previous 72 hours); AND
- the person has had two consecutive respiratory specimens negative for SARS-CoV-2 by PCR taken at least 24 hours. **PP** [Taskforce/CDNA]

OTHER TREATMENTS

VTE prophylaxis
CONSENSUS RECOMMENDATION

Use prophylactic doses of anticoagulants, preferably LMWH (e.g. enoxaparin 40 mg once daily or dalteparin 5000 IU once daily) in **adults with moderate COVID-19 or other indications**, unless there is a contraindication, such as risk for major bleeding. Where eGFR (see below) is less than 30 mL/min/1.73m², unfractionated heparin or clearance-adjusted doses of LMWH may be used (e.g. enoxaparin 20 mg once daily or dalteparin 2500 IU once daily). **CBR** [Taskforce]

For body weights outside 50-90 kg or heights outside 150-180 cm, calculate the BSA and multiply the eGFR by BSA/1.73. **PP** [Taskforce]

Increased-dose VTE prophylaxis
CONSENSUS RECOMMENDATION

Consider using increased prophylactic dosing of anticoagulants, preferably LMWH (e.g. enoxaparin 40 mg twice daily or dalteparin 5000 IU twice daily) in **adults with severe or critical COVID-19 or other indications,** unless there is a contraindication, such as risk for major bleeding or platelet count < 30 x 10^9/L. Where eGFR (see below) is less than 30 mL/min/1.73m², unfractionated heparin or clearance-adjusted doses of LMWH may be used (e.g. enoxaparin 40 mg once daily or dalteparin 5000 IU once daily). **CBR** [Taskforce]

For body weights outside 50-90 kg or heights outside 150-180 cm, calculate the BSA and multiply the eGFR by BSA/1.73. **PP** [Taskforce]

In all people with suspected or confirmed COVID-19, switch nebulisers to metered aerosols with spacers if possible. **PP** [Taskforce/ANZICS/ASID]

In people with suspected or confirmed COVID-19, consider alternative routes of administration for intranasal medicines, recognising that in some situations administration via the intranasal route may be a safer option for affected individuals and healthcare workers. **PP** [Taskforce/ACSQHC]

FLUID MANAGEMENT

In all patients with suspected or confirmed moderate to severe COVID-19, use a restrictive fluid management strategy, avoiding the use of 'maintenance' intravenous fluids, high volume enteral nutrition, and fluid bolus for hypotension. **PP** [Taskforce/ANZICS]

RESPIRATORY SUPPORT

In people with suspected or confirmed COVID-19 and a SaO_2 ≤ 92% or significantly below baseline, initiate supplemental oxygen (1-4 L/min) via nasal prongs. **PP** [Taskforce/ASID]

For details of high level respiratory support see the **RESPIRATORY SUPPORT FOR SEVERE TO CRITICAL COVID-19** Clinical Flowchart

Sources

ACEM – Australasian College for Emergency Medicine Clinical guidelines for the management of COVID-19 in Australasian emergency departments. V1.0, 26 March 2020

ACSQHC – Australian Commission on Safety and Quality in Health Care. COVID-19 Position Statement - Managing fever associated with COVID-19 (Revised 29 April 2020). Managing intranasal administration of medicines for patients during COVID-19 (Revised 19 May 2020)

AHPPC – Australian Health Protection Principal Committee (AHPPC). Guidance on the use of personal protective equipment (PPE) in hospitals during the COVID-19 outbreak. Updated 19 June 2020

ANZICS – The Australian and New Zealand Intensive Care Society (ANZICS) COVID-19 Guidelines. V1.0, 16 March 2020

ASID – Interim guidelines for the clinical management of COVID-19 in adults. Australasian Society for Infectious Diseases (ASID). V1.0, 20 March 2020

CDNA – Coronavirus Disease 2019 (COVID-19) Communicable Diseases Network Australia (CDNA) National Guidelines for Public Health Units. V.3.8 23 August 2020

Taskforce – Current guidance from the National COVID-19 Clinical Evidence Taskforce