Pre-assessment action plan for respiratory infections



This pre-assessment supports prevention and testing of respiratory pathogens and access to antiviral medication for adults who are at higher risk of severe disease from COVID-19 and influenza. The pre-assessment should be completed by the person's regular doctor before the person tests positive for COVID-19 or influenza. It may also be used for people who are travelling interstate, internationally or on cruise vessels to support decision making for antiviral medicines.

to support decision making for antiviral medicines.						
An assessment and prescription by a doctor will still be required at the time of illness.						
Date completed: / / (dd/mm/yyyy)						
Patient details						
Patient's full name:						
DOB: / / (dd/mm/yyyy) Gender: Male Female Prefer not to say						
Prevention						
VACCINATION						
COVID-19 vaccine:						
Last dose received: / / (dd/mm/yyyy)						
Last COVID-19 infection date: / / (dd/mm/yyyy)						
NEXT COVID-19 BOOSTER DOSE DUE*: / / (dd/mm/yyyy)						
* If the patient has a confirmed COVID-19 infection after this checklist has been completed, the next vaccine due date may change (see ATAGI recommendations)						
Patient is up to date with COVID-19 vaccination COVID-19 vaccination declined						
Influenza vaccine:						
Last dose received: / / (dd/mm/yyyy)						
Patient is up to date with seasonal influenza vaccination Influenza vaccination declined						

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Testing

TESTING ADVICE FOR COVID-19, INFLUENZA AND OTHER RESPIRATORY PATHOGENS

People at higher risk of severe disease should discuss with their doctor what test should be done if they develop symptoms (runny nose, sore throat, cough, fever).

See below for considerations:

- Patients should do a RAT first as this will provide a quick result. If negative, they should get a PCR test immediately if they are at higher risk of severe disease.
- Consider providing them with a pre-filled pathology form for a COVID-19, and influenza and RSV PCR test in case they do get symptoms.
- If the patient's RAT or PCR is **positive** for COVID-19 or influenza, they should contact a doctor for a review and prescription (if indicated) as soon as possible, and within 5 days for COVID-19 antivirals, and within 48 hours for influenza antivirals, since symptom onset or positive test.

Treatment

TREATMENT FOR COVID-19

Nirmatrelvir plus ritonavir (Paxlovid™) and **molnupiravir** (Lagevrio®) are available in Australia for the treatment of confirmed COVID-19. Refer to the <u>National Clinical Evidence Taskforce COVID-19</u> living guidelines for the latest treatment recommendations.

Nirmatrelvir plus ritonavir (Paxlovid™) is preferred to molnupiravir (Lagevrio®) for the treatment of COVID-19 in high-risk individuals, however its use is limited by contraindications and drug interactions. In the case of a contraindication, molnupiravir is recommended. The earliest possible antiviral treatment is associated with the most favourable outcomes and should be an aim of treatment.

See Flowchart - Prescribing considerations for nirmatrelvir plus ritonavir (Paxlovid™).

<u>PBS eligibility</u> should also be confirmed at the time of prescribing. Patients who do not meet PBS criteria but meet the <u>National Clinical Evidence Taskforce COVID-19</u> for oral treatment may be able to access antivirals through NSW Health pharmacy departments with a <u>Prescription and declaration form – oral antiviral medicines</u> for COVID-19.

Nirmatrelvir plus ritonavir (Paxlovid™):

Refer to the CEC drug guideline and TGA approved Product Information:

And drug interaction checker: e.g., https://www.covid19-druginteractions.org/checker

Is the patient suitable for nirmatrelvir plus ritonavir? Yes No

Recommended dosing

Note: Nirmatrelvir plus ritonavir dosing requires adjustment in renal impairment. If there is a concern or a change in the patient's condition, then renal function should be checked prior to prescribing.

Most recent eGFR: / / (dd/mm/yyyy)

Adequate renal function (eGFR ≥ 60mL/min)	Nirmatrelvir 300 mg + ritonavir 100 mg every 12 hours for 5 days
Moderate renal impairment (eGFR ≥ 30 to < 60 mL/min)	Nirmatrelvir 150 mg + ritonavir 100 mg every 12 hours for 5 days
Severe renal impairment (eGFR < 30 mL/min)	USE IS CONTRAINDICATED

Source: TGA approved Product Information

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Treatment (cont.)

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Refer to the <u>CEC drug guideline</u> and <u>TGA approved Product Information</u>:

Is the patient suitable for molnupiravir? Yes No

Recommended dosing

Molnupiravir 800 mg (4 x 200 mg capsules) every 12 hours for 5 days.

TREATMENT FOR INFLUENZA

Oseltamivir (Tamiflu®):

Can be used for treatment for confirmed influenza or prophylaxis to confirmed exposure. Refer to the <u>eTG Therapeutic Guidelines</u> and <u>TGA approved Product Information</u> for the latest treatment recommendations.

Is the patient suitable for treatment and/or prophylaxis with oseltamivir? Yes No

Recommended dosing

Normal renal function	For treatment of confirmed influenza	Oseltamivir 75 mg twice daily for 5 days	
	For prophylaxis after confirmed exposure	Oseltamivir 75 mg once daily for 10 days	
Impaired renal function	Refer to the <u>eTG Therapeutic Guidelines</u> and <u>TGA approved Product Information</u> for dose adjustment		

For advice on preparing and administering oseltamivir in patients with swallowing difficulties or enteral feeding tubes, see Information for clinicians.

Additional documents

The following supporting documentation has been attached to the patient's file (if applicable):

A pre-prepared pathology form, after discussion on how and when it should be used

Updated health summary, medication list, and any relevant pathology results (e.g., renal function)

A copy of the patient's drug interaction summary

Medical	practitioner
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Doctor's name (print):				
Contact number:				

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