NSW GOVERNMENT | Health

POISONS AND THERAPEUTIC GOODS ACT 1966

AUTHORITY

I, JUDITH MACKSON, Chief Pharmacist, a duly appointed delegate of the Secretary, NSW Health, make this instrument pursuant to clauses 170 and 171 of the *Poisons and Therapeutic Goods Regulation 2008*, for the purpose of section 10(4)(d) of the *Poisons and Therapeutic Goods Act 1966* and clause 53 of the *Poisons and Therapeutic Goods Regulation 2008.*

Judith Mackson
Chief Pharmacist
(Delegation Numbers: PH427, PH380, PH381)

2 September 2022

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1) Authority

1. This instrument authorises a registered nurse at a public hospital to supply the following restricted substances:
   a) molnupiravir
   b) nirmatrelvir plus ritonavir.

2. The authority signed on 24 February 2022 and annexed to this authority is cancelled.

2) Conditions

1. The registered nurse must only supply the restricted substance:
   a) at a public hospital that does not have an onsite Pharmacy Department, and
   b) at a time when a medical practitioner is not onsite at the public hospital.

2. The restricted substance must only be supplied on a prescription issued by a medical practitioner sent to the registered nurse by email or facsimile.

3. The registered nurse must supply the restricted supply in the unopened pack supplied by a public hospital Pharmacy Department.

4. The registered nurse must label the pack containing the restricted substance with the following details:
   a) the patient's name
   b) the name, strength, form and quantity of the restricted substance supplied
   c) the directions for use on the prescription
   d) the date of supply
   e) the name, address and telephone number of the public hospital from where the restricted substance is supplied
   f) the words 'KEEP OUT OF REACH OF CHILDREN' in red on a white background.

5. The registered nurse must record the supply in the medical record of the person for who the prescription is issued with the following detail:
   a) the name of the medical practitioner who issued the prescription
   b) the name, strength, form and quantity of the restricted substance supplied
   c) the directions for use on the prescription
   d) the date the restricted substance was supplied.

6. The registered nurse must immediately send a copy of the order along with a record of the supply of the restricted substance to a public hospital Pharmacy Department by email or facsimile.

7. The registered nurse must supply to the patient or carer a copy of the relevant Consumer Medicines Information for the restricted substance.

3) Publication

This authority will be published in the NSW Health website.

4) Duration

This instrument commences on signature and expires on 31 July 2025, unless earlier revoked.

POISONS AND THERAPEUTIC GOODS ACT 1966

AUTHORITY

I, JUDITH MACKSON, Chief Pharmacist, a duly appointed delegate of the Secretary, NSW Health, make this instrument pursuant to clauses 170 and 171 of the *Poisons and Therapeutic Goods Regulation 2008*, for the purpose of section 10(4)(d) of the *Poisons and Therapeutic Goods Act 1966* and clause 53 of the *Poisons and Therapeutic Goods Regulation 2008.*

Judith Mackson
Chief Pharmacist
(Delegation Numbers: PH427, PH380, PH381)

24 February 2022

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1) Authority

This instrument authorises a registered nurse at a public hospital to supply the following restricted substances:

a) molnupiravir
b) nirmatrelvir plus ritonavir.

2) Conditions

1. The registered nurse must only supply the restricted substance:
   a) at a public hospital that does not have an onsite Pharmacy Department, and
   b) at a time when a medical practitioner is not onsite at the public hospital.

2. The restricted substance must only be supplied when ordered by a medical practitioner on the NSW Health *Prescription and Declaration – Oral antiviral medicines for COVID-19 (for use by GPs)* form sent to the registered nurse by email or facsimile.

3. The registered nurse must supply the restricted supply in the unopened pack supplied by a public hospital Pharmacy Department.

4. The registered nurse must label the pack containing the restricted substance with the following details:
   a) the patient's name
   b) the name, strength, form and quantity of the restricted substance supplied
   c) the directions for use on the prescription
   d) the date of supply
   e) the name, address and telephone number of the public hospital from where the restricted substance is supplied
   f) the words 'KEEP OUT OF REACH OF CHILDREN' in red on a white background.

5. The registered nurse must record the supply in the medical record of the person for who the prescription is issued with the following detail:
   a) the name of the medical practitioner who issued the prescription
   b) the name, strength, form and quantity of the restricted substance supplied
   c) the directions for use on the prescription
   d) the date the restricted substance was supplied.

6. The registered nurse must immediately send a copy of the order along with a record of the supply of the restricted substance to a public hospital Pharmacy Department by email or facsimile.

7. The registered nurse must supply to the patient or carer a copy of the relevant Consumer Medicines Information for the restricted substance.

3) Publication

This authority will be published in the NSW Health website.

4) Duration

This instrument commences on signature and expires on 30 September 2022, unless earlier revoked.