



Queensland

Health Legislation Amendment Regulation (No. 2) 2019

Subordinate Legislation 2019 No. 117

made under the

Health Act 1937

Public Health Act 2005

Radiation Safety Act 1999

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Part 1 Preliminary

1 Short title

This regulation may be cited as the *Health Legislation Amendment Regulation (No. 2) 2019*.

2 Commencement

This regulation commences on 1 July 2019.

Part 2 Amendment of Health (Drugs and Poisons) Regulation 1996

3 Regulation amended

This part amends the *Health (Drugs and Poisons) Regulation 1996*.

4 Amendment of s 78A (Approved drug—nabiximols)

(1) Section 78A, heading—

omit, insert—

78A Medicinal cannabis

(2) Section 78A(1)—

omit, insert—

(1) Subject to section 74(3), a person must not dispense, prescribe, supply or use a controlled drug that is medicinal cannabis unless the person—

(a) is a specialist medical practitioner; or

(b) is another doctor who dispenses, prescribes, supplies or uses the medicinal cannabis under an approval.

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Maximum penalty—80 penalty units.

(3) Section 78A(3)—

omit.

5 Amendment of s 79 (Prescribing controlled drugs)

Section 79(4)(j), ‘dronabinol’—

omit, insert—

medicinal cannabis

6 Amendment of s 82 (Conditions of dispensing)

Section 82(2)(g), ‘dronabinol or nabiximols’—

omit, insert—

medicinal cannabis

7 Amendment of s 118 (Storage of controlled drugs at institutions)

(1) Section 118(1)(a) and (b)—

omit, insert—

(a) if the controlled drug is medicinal cannabis—in a way that complies with the medicinal cannabis security standard; or

(b) otherwise—

(i) in a receptacle that complies with appendix 6 of this regulation; or

(ii) in another place (a *secure place*) an inspector who inspects the place is reasonably satisfied is at least as secure as a receptacle mentioned in subparagraph (i).

(2) Section 118(2)(a), (b) and (c)—

omit, insert—

- (a) if the controlled drug is medicinal cannabis—ensure the drug is stored in a way that complies with the medicinal cannabis security standard; or
- (b) otherwise—
 - (i) ensure the drug is stored in the receptacle or secure place mentioned in subsection (1)(b); and
 - (ii) always keep the receptacle or secure place locked (other than when a controlled drug is being put into or taken out of the receptacle or secure place); and
 - (iii) ensure the key or combination to, or other way used to personally access, the receptacle or secure place can not be used by a person who is not authorised to possess a controlled drug at the institution.

8 Amendment of s 119 (Storage of controlled drugs generally)

(1) Section 119(1)(a) and (b)—

omit, insert—

- (a) if the controlled drug is medicinal cannabis—in a way that complies with the medicinal cannabis security standard; or
- (b) otherwise—
 - (i) in a receptacle that complies with appendix 6 of this regulation; or
 - (ii) in another place (a *secure place*) an inspector who inspects the place is reasonably satisfied is at least as secure

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as a receptacle mentioned in
subparagraph (i).

(2) Section 119(2), ‘The’—

omit, insert—

For subsection (1)(b), the

9 Amendment of s 120 (Notice required if lengthy treatment with controlled drug)

(1) Section 120, heading—

omit, insert—

120 Request for information about treatment with controlled drug

(2) Section 120(1) and (2)—

omit, insert—

(1) This section applies if the chief executive reasonably suspects a relevant practitioner has administered, dispensed, prescribed or supplied a controlled drug in the treatment of a patient.

(3) Section 120(3), ‘additional’—

omit, insert—

stated

(4) Section 120(3) to (5)—

renumber as section 120(2) to (4).

10 Insertion of new ch 5, pt 2, div 5

Chapter 5, part 2—

insert—

Division 5 Transitional provision for Health Legislation

Amendment Regulation (No. 2) 2019

316 Definition for division

In this division—

repealed regulation means the repealed *Public Health (Medicinal Cannabis) Regulation 2017*.

317 Existing manufacturing approvals continue

- (1) This section applies if, immediately before the commencement, a person was the holder of a manufacturing approval under the repealed regulation, part 2.
- (2) If the manufacturing approval authorised the manufacture of a controlled drug, the person is taken to hold a controlled drug manufacturer licence under this regulation—
 - (a) to manufacture the controlled drug; and
 - (b) for the same term that applied to the approval immediately before the commencement; and
 - (c) subject to a condition under the repealed regulation, section 23 that applied to the approval immediately before the commencement.
- (3) If the manufacturing approval authorised the manufacture of a restricted drug, the person is taken to hold a restricted drug manufacturer licence under this regulation—
 - (a) to manufacture the restricted drug; and
 - (b) for the same term that applied to the approval immediately before the commencement; and

[s 10]

- (c) subject to a condition under the repealed regulation, section 23 that applied to the approval immediately before the commencement.
- (4) This section applies despite section 22.

318 Existing wholesaling approvals continue

- (1) This section applies if, immediately before the commencement, a person was the holder of a wholesaling approval under the repealed regulation, part 2.
- (2) If the wholesaling approval authorised the wholesale of a controlled drug, the person is taken to hold a controlled drug wholesaler licence under this regulation—
 - (a) to sell the controlled drug by wholesale; and
 - (b) for the same term that applied to the approval immediately before the commencement; and
 - (c) subject to a condition under the repealed regulation, section 23 that applied to the approval immediately before the commencement.
- (3) If the wholesaling approval authorised the wholesale of a restricted drug, the person is taken to hold a restricted drug wholesaler licence under this regulation—
 - (a) to sell the restricted drug by wholesale; and
 - (b) for the same term that applied to the approval immediately before the commencement; and
 - (c) subject to a condition under the repealed regulation, section 23 that applied to the

approval immediately before the commencement.

- (4) This section applies despite section 22.

319 Exemption from fees

- (1) This section applies if a person is taken to hold a drug licence under section 317 or 318.
- (2) Despite section 17(1)(b), the person is not required to pay a fee for an application for the renewal of the drug licence.

11 Amendment of appendix 1 (Provisions not applying to morphine or opium in compounded preparations)

Appendix 1, entry for section 120, 'Notice required if lengthy treatment with controlled drug'—

omit, insert—

Request for information about treatment with controlled drug

12 Amendment of appendix 9 (Dictionary)

Appendix 9—

insert—

cannabis product means a product—

- (a) that is or was any part of a plant of the genus *Cannabis*, whether living or dead; or
- (b) otherwise derived, wholly or in part, from any part of a plant of the genus *Cannabis*, whether living or dead; or
- (c) that has, or is intended by the manufacturer of the product to have, a pharmacological effect that is substantially similar to the

-
- (a) each of the following respiratory diseases is prescribed—
 - (i) cancer;
 - (ii) chronic obstructive pulmonary disease, including chronic bronchitis and emphysema;
 - (iii) pneumoconiosis, including asbestosis, coal worker's pneumoconiosis, mixed-dust pneumoconiosis and silicosis; and
 - (b) the type of dust prescribed is inorganic dust.

49B Prescribed medical practitioners—Act, s 279AA

For section 279AA of the Act, definition *prescribed medical practitioner*, the prescribed class of persons is medical practitioners registered under the Health Practitioner Regulation National Law as specialist health practitioners in either of the following specialties or specialty fields—

- (a) occupational and environmental medicine;
- (b) respiratory and sleep medicine.

49C Prescribed period—Act, s 279AF

For section 279AF(2) of the Act, the prescribed period is 30 days from the day the prescribed medical practitioner diagnoses the person as having a notifiable dust lung disease.

15 Replacement of s 60 (Paint—Act, s 60)

Section 60—
omit, insert—

14A Use licensee—Act, s 103K

- (1) For section 103K(1)(a) of the Act, a person is taken to hold a use licence if the person is registered under the Health Practitioner Regulation National Law to practice in the dental profession as a dentist, other than as a student.
- (2) The radiation source the person is allowed to use is an intra-oral dental radiation apparatus.
- (3) The radiation practice the person is allowed to carry out is intra-oral dental plain radiography.
- (4) It is a condition of the licence that the person uses the radiation source in compliance with the ‘Code of Practice for Radiation Protection in Dentistry (2005)’ published by ARPANSA.

Editor’s note—

A copy of the code mentioned in subsection (4) is available on ARPANSA’s website.

14B Transport licensee—Act, s 103K

- (1) For section 103K(1)(b) of the Act, a person is taken to hold a transport licence if the person holds an authority under a corresponding transport law to transport a radioactive substance.
- (2) The person may transport the radioactive substance into Queensland.
- (3) It is a condition of the licence that the person transports the radioactive substance in compliance with the ‘Code of Practice for the Security of Radioactive Sources (2019)’ published by ARPANSA.

Editor’s note—

A copy of the code mentioned in subsection (3) is available on ARPANSA’s website.

- (4) In this section—

[s 20]

authority includes an accreditation, approval, certification or licence.

corresponding transport law means a law of another State or the Commonwealth relating to the transportation of radioactive substances.

20 Amendment of s 81 (Register of licensees—Act, s 207)

(1) Section 81, after ‘about licensees’—

insert—

, other than prescribed licensees,

(2) Section 81—

insert—

(2) Also, for section 207(1)(f) of the Act, the register must contain the following information about a prescribed licensee whose licence has been suspended or cancelled—

(a) if the licensee’s licence has been cancelled—

(i) the licensee’s name; and

(ii) the day the decision to cancel the licence takes effect;

(b) if the licensee’s licence has been suspended and the period of suspension has not ended—

(i) the licensee’s name; and

(ii) the day the decision to suspend the licence takes effect; and

(iii) the period of the suspension.

ENDNOTES

- 1 Made by the Governor in Council on 27 June 2019.
- 2 Notified on the Queensland legislation website on 28 June 2019.
- 3 The administering agency is Queensland Health.

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