



Medicines and Poisons Act 2019

Medicines and Poisons (Monitored Medicines Database Testing) Regulation 2020

Current as at 1 May 2020

Reprint note

This is the last reprint before repeal. Repealed on 27 September 2021 by 2021 SL No. 140 s 245.

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Queensland

Medicines and Poisons (Monitored Medicines Database Testing) Regulation 2020

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Medicines and Poisons (Monitored Medicines Database Testing) Regulation 2020

Part 1 Preliminary

1 Short title

This regulation may be cited as the *Medicines and Poisons (Monitored Medicines Database Testing) Regulation 2020*.

2 Commencement

This regulation commences on 1 May 2020.

3 Purpose

The purpose of this regulation is to facilitate the establishment and testing of the monitored medicines database using information given and recorded under the *Health Act 1937*.

4 Definitions

In this regulation—

authority see the *Health (Drugs and Poisons) Regulation 1996*, appendix 9.

controlled drug see the *Health (Drugs and Poisons) Regulation 1996*, appendix 9.

dispenser see the *Health (Drugs and Poisons) Regulation 1996*, appendix 9.

monitored medicine treatment activity means an activity—

- (a) performed by the holder of an authority or a relevant approval; and

- (b) relevant registration information for a monitored medicine treatment activity performed under an opioid treatment program that—
 - (i) was given to, or recorded by or on behalf of, the chief executive under the *Health Act 1937* at any time before the commencement; and
 - (ii) is kept electronically by the chief executive.

(2) In this section—

relevant health information means the following information—

- (a) information in relation to a monitored medicine treatment activity;
- (b) information in relation to a relevant approval;
- (c) information kept, from time to time, on a register under the Health Practitioner Regulation National Law for a prescriber or dispenser who holds an authority or a relevant approval;
- (d) personal information to identify a prescriber or dispenser for accessing or using the monitored medicines database;
- (e) information recorded in relation to a prescriber's or dispenser's access to, or use of, the monitored medicines database.

relevant registration information, for a monitored medicine treatment activity performed under an opioid treatment program, means information about the registration of particular persons in the program.

7 Users of database—Act, s 227

For section 227(4) of the Act, definition *user*, each of the following entities is prescribed to be a user—

- (a) a prescriber;

[s 8]

- (b) a dispenser;
- (c) a person employed by the chief executive.

8 Purposes for disclosure—Act, s 227

For section 227(2) of the Act, each of the following purposes is prescribed—

- (a) for a user who is a prescriber or dispenser—testing the monitored medicines database;
- (b) for a user who is a person employed by the chief executive—establishing or testing the monitored medicines database, including training about using the database.