

Queensland



Drugs Misuse Act 1986

DRUGS MISUSE REGULATION 1987

**Reprinted as in force on 5 October 2001
(includes amendments up to SL No. 174 of 2001)**

Reprint No. 4

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Information about this reprint

This regulation is reprinted as at 5 October 2001. The reprint shows the law as amended by all amendments that commenced on or before that day (Reprints Act 1992 s 5(c)).

The reprint includes a reference to the law by which each amendment was made—see list of legislation and list of annotations in endnotes.

Minor editorial changes allowed under the provisions of the Reprints Act 1992 have also been made to use aspects of format and printing style consistent with current drafting practice (s 35).

This page is specific to this reprint. See previous reprints for information about earlier changes made under the Reprints Act 1992. A table of earlier reprints is included in the endnotes.

Also see endnotes for information about—

- **when provisions commenced**
- **editorial changes made in earlier reprints.**

Queensland



DRUGS MISUSE REGULATION 1987

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DRUGS MISUSE REGULATION 1987

[as amended by all amendments that commenced on or before 5 October 2001]

PART 1—PRELIMINARY

1 Short title

This regulation may be cited as the *Drugs Misuse Regulation 1987*.

4 Dictionary

The dictionary in schedule 9 defines particular words used in this regulation.

PART 2—SEARCH WARRANT NOTICES AND RECORD

5 Notice to justice before whom a complaint to ground a search warrant is to be sworn

A justice of the peace (other than a stipendiary magistrate) who has before him or her a complaint to ground a search warrant shall be given a notice in the approved form by the complainant and shall read such notice prior to administering an oath for the purposes of such complaint.

6 Record of proceedings

(1) A justice of the peace who has a complaint sworn before him or her to ground a search warrant shall cause a record to be made of such proceedings in the approved form.

(2) The justice of the peace shall retain or cause to be retained by the nearest clerk of the court the complaint to ground a search warrant and record of proceedings for 2 years or such longer period as may be required in the particular case.

7 Notice to occupier of place entered pursuant to warrant

(1) A police officer to whom a search warrant has been issued shall prepare an occupier's notice in the approved form.

(2) A police officer executing a search warrant shall—

- (a) upon entry into or on the place to which the warrant relates, or at the first reasonable opportunity thereafter, serve the occupier's notice on a person who appears to be an occupier of that place; and
- (b) if no such person is then present, or if service is not practicable for any other reason, serve the occupier's notice by leaving it in a conspicuous location in or on that place.

(3) Subsection (2) does not apply where the police officer executing a search warrant has reasonable grounds to believe that service of an occupier's notice would frustrate or otherwise hinder the investigation of the offence in respect of which the search warrant was issued.

PART 3—SYRINGES AND DANGEROUS DRUGS DISPOSAL PROCEDURES

9 Prescribed procedures for the disposal of hypodermic syringes and needles

For the purposes of section 10(4A) of the Act, the prescribed procedures for the disposal of a hypodermic syringe or needle shall be as follows—

- (a) by placing the hypodermic syringe or needle in a rigid wall, puncture resistant container and that container is sealed or securely closed in such a manner that its contents are incapable of causing injury to any person; or
- (b) by giving the hypodermic syringe or needle to a person who is a medical practitioner, pharmacist or person or a member of a class of persons referred to as authorised in section 10(3) of the Act.

10 Prescribed procedure for disposal of dangerous drugs

For the purposes of section 52A¹ of the Act, the prescribed procedure for the disposal of a thing shall be as follows—

- (a) in the case where the thing is a trace amount of a dangerous drug contained in a hypodermic syringe or needle, by disposing of the hypodermic syringe or needle in accordance with the procedures prescribed in section 9; or
- (b) in any other case, at the first reasonable opportunity, by giving—
 - (i) such thing; and
 - (ii) where such thing is contained in a hypodermic syringe or needle, such syringe or needle;

to an officer authorised to exercise the powers contained in the *Health Act 1937*, section 132.²

PART 4—CONTROLLED SUBSTANCES

11 Other act that is a relevant transaction—Act, s 43C(b)

Any act by which a controlled substance is supplied³ by a person, in or in connection with the person's business, to anyone else is a relevant transaction for the supply of a controlled substance.

Example—

A and B are partners in a chain of pharmacies. They make cold tablets to sell in the pharmacies by compounding ephedrine (a controlled substance) with other substances.

The partners sell some of the left over ephedrine to a pharmaceutical research company and give the rest away.

Both the sale and gift of ephedrine are relevant transactions.

1 Section 52A (Prescribed persons permitted to receive and dispose of dangerous drugs) of the Act

2 *Health Act 1937*, section 132 (Powers of officers)

3 Under section 43A of the Act, “supply” means give, distribute, sell or supply.

12 Documents and proof of identity required for supply of a controlled substance—Act, s 43D(1)(a)

(1) This section applies to a person who supplies a controlled substance under a relevant transaction to anyone else (a “**recipient**”).

(2) The person must, before supplying the substance, obtain from the recipient a written order for the supply of the substance showing the following information—

- (a) the recipient’s name and address, and if the recipient purports to obtain the substance for another person, the other person’s name and address;
- (b) the date and number of the order;
- (c) the name and quantity of the substance to be supplied;
- (d) the purpose for which the substance is to be supplied.

(3) If the recipient is an individual, the person must, before supplying the substance, require the recipient to produce an official document containing the recipient’s photograph (for example, a passport or drivers licence) as evidence of the recipient’s identity.

(4) The person must, immediately the person supplies the substance under the transaction, make an invoice for the supply of the substance showing the following details—

- (a) the recipient’s name and address;
- (b) the recipient’s order number for the supply of the substance;
- (c) the date the substance was supplied;
- (d) the name and quantity of the substance supplied.

Maximum penalty for subsection (4)—20 penalty units.

13 Details about supply of controlled substance to be recorded in register

(1) The following details about a relevant transaction for the supply of a controlled substance must be recorded in the register⁴—

4 See section 43D(1)(c) of the Act for the requirement to keep the register.

- (a) the name and address of the recipient and, if the recipient purports to obtain the substance for another person, the other person's name and address;
- (b) the recipient's order number for the supply of the substance;
- (c) the invoice number for the supply of the substance;
- (d) if the recipient is—
 - (i) a company—its Australian Company Number; or
 - (ii) an individual—the type of official document produced under section 12(3) and the following details about the document—
 - (A) who issued it;
 - (B) its serial number or other identifying number or mark;
- (e) the name and quantity of the substance supplied;
- (f) the date the substance was supplied;
- (g) the purpose for which the substance was supplied.

(2) The details must be recorded in the register as soon as practicable, but in no case later than 7 days, after the day the person supplied the substance under the transaction.

(3) Nothing in this section prevents the keeping of a single register for the Act and another Act if—

- (a) the keeping of the single register is not contrary to the other Act; and
- (b) the details recorded under subsection (1) are easily identifiable in the single register.

14 Details about loss or theft of controlled substance to be recorded in register

The following details of the reporting to a police officer of the loss or theft of a controlled substance must be recorded in the register—

- (a) the day and place the report was made;
- (b) the name and registered number of the officer to whom the report was made;
- (c) the name and quantity of the substance lost or stolen.

15 Keeping of register, invoice and other documents

(1) This section applies to the register and other documents mentioned in the Act, section 43D(1)⁵ and the invoice required under section 12(4).

(2) A person who supplies a controlled substance under a relevant transaction must keep the register, documents or invoice—

- (a) at the principal or only place in Queensland where the person engages in relevant transactions; and
- (b) for 2 years from the day the person supplied the substance under the transaction.

PART 5—TRIAL PLANTING OF CANNABIS SATIVA FOR COMMERCIAL FIBRE PRODUCTION

Division 1—Preliminary

16 Purpose of pt 5

This part states the extent of the exemptions necessary for achieving the purpose of part 5B⁶ of the Act and makes other necessary provisions for achieving the purpose.

17 Limit of effect of exemption

The exemption of a person under this part from section 6, 8 or 9 of the Act has effect, in relation to the supply, production or possession of cannabis sativa only to the extent the supply, production or possession is for the purpose, directly or indirectly, of research carried out under part 5B of the Act.

5 Section 43D (Requirements for supply of controlled substance under relevant transactions) of the Act

6 Part 5B (Trial planting of cannabis sativa for commercial fibre production) of the Act

18 Conditions

The conditions for each exemption are in schedule 7.

Division 2—Exemptions

19 Exemptions for seed suppliers

A seed supplier is exempt from—

- (a) section 6 of the Act, for supplying cannabis seeds to a person who may lawfully possess them; and
- (b) section 9 of the Act, for possessing cannabis seeds to supply them to a person who may lawfully possess them.

20 Exemptions for carriers

(1) A carrier who is engaged or employed by a seed supplier to transport cannabis seeds is exempt from sections 6 and 9 of the Act, for transporting the seeds to a person who may lawfully possess them.

(2) A carrier who is engaged or employed by someone who may lawfully possess cannabis plant material to transport the plant material is exempt from sections 6 and 9 of the Act, for transporting the plant material to a person who may lawfully possess it.

(3) An exemption under subsection (1) or (2) is for the time necessary to transport the seeds or plant material to the person to whom they are consigned.

21 Exemptions for growers

(1) A grower is exempt from section 6 of the Act—

- (a) for supplying cannabis seeds to—
 - (i) a carrier, for transporting the seeds to another grower or an authorised officer; or
 - (ii) a person who may lawfully possess the seeds; and
- (b) for supplying cannabis plant material to—
 - (i) a carrier, for transporting the plant material to a person who may lawfully possess it; or

(ii) a person who may lawfully possess the plant material.

(2) A grower is exempt from section 8 of the Act for producing cannabis plants on land for which the grower is named in schedule 8.

(3) A grower is exempt from section 9 of the Act for possessing cannabis seeds and cannabis plants on land for which the grower is named in schedule 8.

22 Exemptions for authorised officer

An authorised officer is exempt from—

- (a) section 6 of the Act, for supplying cannabis seeds to—
 - (i) a grower; or
 - (ii) a DPI researcher responsible for the management and control of the Australian Tropical Crops Genetic Resource Centre, for safe keeping;⁷ and
- (b) section 9 of the Act, for possessing cannabis seeds and cannabis plant material for ensuring compliance with this part, including the conditions in schedule 7.

23 Exemptions for DPI researchers

A DPI researcher is exempt from—

- (a) section 6 of the Act, for supplying—
 - (i) cannabis seeds to a grower; and
 - (ii) cannabis plants to another DPI researcher; and
 - (iii) cannabis plant material to a non-DPI researcher; and
- (b) section 8 of the Act, for producing cannabis plants and cannabis seeds; and
- (c) section 9 of the Act, for possessing cannabis seeds, cannabis plants, or cannabis plant material.

⁷ The Centre is located at the Department of Primary Industries' Biloela Research Station.

24 Exemptions for non-DPI researchers

A non-DPI researcher is exempt from—

- (a) section 6 of the Act, for supplying cannabis seeds, cannabis plants, or cannabis plant material in the form of a processed fibre product, to a DPI researcher or another non-DPI researcher; and
- (b) section 8 of the Act, for producing cannabis plants and cannabis seeds; and
- (c) section 9 of the Act, for possessing cannabis seeds, cannabis plants, or cannabis plant material.

25 Exemptions for employees etc.

25.(1) A person (“**employee**”) who is engaged or employed by a person (“**employer**”) who may lawfully possess cannabis seeds or cannabis plant material is exempt from sections 8 and 9 of the Act, but only for the employee to perform functions necessary for the purpose of the employer’s lawful possession of the seeds or plant material.

(2) A person who is a member of the immediate family of a grower is exempt from sections 8 and 9 of the Act, but only for helping the grower perform functions normally associated with growing and harvesting cannabis plants.

(3) A person, other than a person exempted under subsection (1) or (2), who performs functions for another person who may lawfully possess cannabis plants, cannabis plant material or cannabis seeds is exempt from sections 8 and 9 of the Act for performing functions necessary for the purpose of the other person’s lawful possession of the plants, plant material or seeds.

Example for subsection (3)—

An authorised officer may ask a person in control of a drying oven used to extract moisture from plant material to dry plant material to enable the tetrahydrocannabinol content of the plant material to be determined.

Division 3—Other provisions

26 Appointment of authorised officers

The commissioner may appoint a person other than a police officer as an authorised officer under this part if—

- (a) the commissioner considers the person has the necessary expertise or experience to be an authorised officer; or
- (b) the person has satisfactorily finished training approved by the commissioner.

27 Authorised officer's identity card

(1) The commissioner must give each authorised officer an identity card.

(2) The identity card must—

- (a) contain a recent photo of the person; and
- (b) be signed by the person; and
- (c) identify the person as an authorised officer under this part; and
- (d) state an expiry date.

(3) A person who stops being an authorised officer must return the person's identity card to the commissioner as soon as possible (but within 21 days) after the person stops being an authorised officer, unless the person has a reasonable excuse.

Maximum penalty—20 penalty units.

(4) This section does not prevent the giving of a single identity card to a person for this part and for other purposes, whether or not for this Act.

28 Production or display of authorised officer's identity card

(1) An authorised officer may exercise a power in relation to someone only if—

- (a) the authorised officer first produces the authorised officer's identity card for the other person's inspection; or
- (b) the authorised officer has the authorised officer's identity card displayed so it is clearly visible to the other person.

(2) However, if for any reason it is not practicable to comply with subsection (1) before exercising the power, the authorised officer must produce the identity card for inspection by the person at the first reasonable opportunity.

29 Deferral of expiry of Act, pt 5B (Act, s 43Y)

The expiry of part 5B⁸ of the Act is deferred by 1 year.⁹

PART 6—TRANSITIONAL PROVISION**30 Transitional provision for Drugs Misuse Amendment Regulation (No. 2) 2001**

(1) To remove doubt, it is declared that this regulation, as in force immediately before the commencement of the *Drugs Misuse Amendment Regulation (No. 2) 2001* (the “**amending regulation**”), continues to apply in relation to an offence against the Act committed before the commencement of the amending regulation.

(2) Proceedings for an offence against the Act committed before the commencement of the amending regulation may be continued or started as if the amending regulation had not been made.

8 Part 5B (Trial planting of cannabis sativa for commercial fibre production) of the Act

9 The part will now expire at the end of the day on 18 December 2002.

SCHEDULE 1

DANGEROUS DRUGS

sections 4, 5, 6, 8, 8A, 9 and 59 of the Act

Amphetamine

Cocaine

Heroin

Lysergide

Methylamphetamine

Phencyclidine

SCHEDULE 2**DANGEROUS DRUGS**

sections 4, 5, 6, 8, 8A, 9 and 59 of the Act

Acetorphine

Acetyldihydrocodeine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of acetyldihydrocodeine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of acetyldihydrocodeine

Acetylmethadol

Acetylmorphines

Alfentanil

Alkoxyamphetamines and bromo-substituted alkoxyamphetamines except where separately specified

Alkoxyphenethylamines and alkyl-substituted alkoxyphenethylamines except where separately specified

Allylprodine

Alphacetylmethadol

Alphameprodine

Alphamethadol

Alphaprodine

Anileridine

Barbituric acid and any 5,5 disubstituted derivatives of barbituric acid, whether or not further substituted at position 1 of the ring

Benzethidine

Benzylmorphine

SCHEDULE 2 (continued)

Betacetylmethadol

Betameprodine

Betamethadol

Betaprodine

Bezitramide

4-Bromo-2,5-dimethoxyamphetamine

4-Bromo-2,5-dimethoxyphenethylamine

Bufotenine

Buprenorphine

Cannabinoids except tetrahydrocannabinols

Cannabis sativa

Clonitazene

Coca leaf

Codeine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 30 mg or less of codeine per dosage unit; or
- (b) in undivided preparations containing 1% or less of codeine

Codeine-N-oxide

Codoxime

4-Cyano-1-Methyl-4-Phenylpiperidine

4-Cyano-2-Dimethylamino-4,4-Diphenylbutane

Desomorphine

Diampromide

Diethylthiambutene

N,N-Diethyltryptamine

SCHEDULE 2 (continued)

Difenoxin except in preparations containing 0.5 mg or less of difenoxin and a quantity of atropine sulphate equivalent to not less than 5% of the dose of difenoxin per dosage unit

Dihydrocodeine except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of dihydrocodeine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of dihydrocodeine

Dihydromorphine

Dimenoxadol

Dimepheptanol

2,5-Dimethoxyamphetamine

2,5-Dimethoxy-4-Ethylamphetamine (DOET)

2,5-Dimethoxy-4-Methylamphetamine

Dimethylamino-1,2-Diphenylethane

3-(1,2-Dimethylheptyl)-1-Hydroxy-7,8,9,10-Tetrahydro-6,6,9-Trimethyl-6-H-Dibenzo(b,d)Pyran

Dimethylthiambutene

N,N-Dimethyltryptamine

Dioxaphetyl butyrate

Diphenoxylate except in preparations containing 2.5 mg or less of diphenoxylate and a quantity of atropine sulphate equivalent to not less than 1% of the dose of diphenoxylate per dosage unit

Dipipanone

Drotebanol

Ecgonine, its esters and derivatives which are convertible to ecgonine and cocaine

Ethylmethylthiambutene

SCHEDULE 2 (continued)

Ethylmorphine except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of ethylmorphine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of ethylmorphine

N-Ethyl-1-Phencyclohexylamine

Etonitazine

Etorphine

Etoxeridine

Fenethylamine

Fentanyl

Furethidine

Gamma hydroxybutyric acid

Hydrocodone

Hydromorphanol

Hydromorphone

Hydroxypethidine

Isomethadone

Ketamine

Ketobemidone

Levophenacetyl-morphine

Lysergamide and N-alkyl derivatives of lysergamide other than lysergide

Lysergic acid

Mecloqualone

Mescaline (3,4,5-Trimethoxyphenethylamine)

Metazocine

Methadone

SCHEDULE 2 (continued)

Methaqualone

Methcathinone

5-Methoxy-3,4-Methylenedioxyamphetamine (MMDA)

4-Methylaminorex

Methyldesorphine

Methyldihydromorphine

3,4-Methylenedioxyamphetamine

3,4-methylenedioxyethylamphetamine (MDEA)

3,4-Methylenedioxymethamphetamine (MDMA)

2-Methyl-3-Morpholino-1, 1-Diphenylpropane Carboxylic acid

Methylphenidate

4-Methylthioamphetamine (4-MTA)

1-Methyl-4-Phenylpiperidine-4-Carboxylic acid

Metopon

Moramide

Morpheridine

Morphine

Morphine methobromide

Morphine-N-oxide

Myrophine

Nabilone

Nicocodine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of nicocodine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of nicocodine

SCHEDULE 2 (continued)

Nicodicodeine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of nicodicodeine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of nicodicodeine

Nicomorphine

Noracymethadol

Norcodeine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of norcodeine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of norcodeine

Norlevorphanol

Normethadone

Normorphine

Norpipanone

Opium

Oxycodone

Oxymorphone

Papaver orientale

Papaver setigerum

Papaver somniferum L. except the seed thereof which seed has been rendered sterile

Parahexyl

Paramethoxyamphetamine (PMA)

Pentazocine

Pethidine

SCHEDULE 2 (continued)

Phenadoxone

Phenampromide

Phenazocine

Phendimetrazine

Phenmetrazine

Phenomorphane

Phenoperidine

1-(1-Phenylcyclohexyl)pyrrolidine

4-Phenylpiperidine-4-Carboxylic acid ethyl ester

Pholcodine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

(a) in divided preparations containing 100 mg or less of pholcodine per dosage unit; or

(b) in undivided preparations containing 2.5% or less of pholcodine

Piminodine

Piritramide

Proheptazine

Properidine

Propiram

Psilocin

Psilocybin

Racemethorphan

Racemoramide

Racemorphan

Sufentanil

Tetrahydrocannabinols including their alkyl homologues except where separately specified; and their corresponding carboxylic acids

Thebacon

SCHEDULE 2 (continued)

Thebaine

1-(1-(2-thienyl)cyclohexyl)piperidine

Tilidine

Trimeperidine

3, 4, 5-Trimethoxyamphetamine (TMA)

SCHEDULE 2A**DANGEROUS DRUGS**

sections 4, 5, 6, 8, 8A, 9 and 59 of the Act

Androisoxazole

Atamestane

Bambuterol

Bolandiol

Bolesterone

Bolazine

Boldenone (dehydrotestosterone)

Bolenol

Bolmantalate

Calusterone

Chlorandrostenolone

4-Chloromethandienone

Chloroxydienone

Chloroxymesterone (dehydrochloromethyltestosterone)

Clembuterol

Clostebol (4-chlorotestosterone)

Danazol

Dihydrolone

Dimethandrostanolone

Drostanolone

Enestebol

Ephedrine

Epitiostanol

SCHEDULE 2A (continued)

Ethyldienolone
Ethyloestranol
Fenoterol
Flunitrazepam
Fluoxymesterone
Formebolone (formyldienolone)
Formoterol
Furazabol
Hydroxystenozol
Mebolazine
Mepitiostane
Mesabolone
Mestanolone (androstalone)
Mesterolone
Methandienone
Methandriol
Methenolone
Methylclostebol
Methyltestosterone
Methyltrienolone
Metribolone
Mibolerone
Nandrolone
Norandrostenolone
Norbolethone
Norclostebol
Norethandrolone
Normethandrone

SCHEDULE 2A (continued)

Ovandrotone

Oxabolone

Oxandrolone

Oxymesterone

Oxymethalone

Prasterone

Propetandriol

Quinbolone

Reproterol

Salbutamol, except in metered aerosols or dry powder or capsules of dry powder for inhalation

Silandrone

Somatropin

Stanolone

Stanazolol

Stenbolone

Terbutaline, except in metered aerosols for inhalation

Testolactone

Testosterone, except in implant preparations for growth promotion in animals

Thiomesterone (tiomesterone)

Trenbolone (trienbolone, trienolone), except in implant preparations for use in animals

Trestolone

Any other anabolic and androgenic steroidal agent

SCHEDULE 3**SPECIFIED QUANTITIES FOR PARTICULAR
DANGEROUS DRUGS**

sections 4, 8, 9, 52A and 59 of the Act

Dangerous drug	Quantity of dangerous drug
Amphetamine	2.0 g
Barbituric Acid and any 5,5 disubstituted derivatives of barbituric acid whether or not further substituted at position 1 of the ring	50.0g
4-Bromo-2,5-dimethoxyamphetamine	0.5g
4-Bromo-2,5-dimethoxyphenethylamine	2.0 g
Cannibis sativa	500.0 g or, if the dangerous drug consists of plants the aggregate weight of which is less than 500.0 g, 100 plants
Cocaine	2.0 g
Codeine	10.0 g
N,N-Diethyltryptamine	2.0g
2,5-Dimethoxy-4-Ethylamphetamine (DOET)	2.0 g
2,5-Dimethoxy-4-Methylamphetamine	2.0 g
N,N-Dimethyltryptamine	2.0 g
Fenethylline	2.0 g
Fentanyl	0.01 g
Gamma hydroxybutyric acid	2.0 g
Heroin	2.0 g
Hydromorphone	2.0 g

SCHEDULE 3 (continued)

Dangerous drug	Quantity of dangerous drug
Lysergide	0.004 g
Methadone	2.0 g
Methcathinone	2.0 g
5-Methoxy-3,4-Methylenedioxyamphetamine (MMDA)	2.0 g
4-Methylaminorex	2.0 g
Methylamphetamine	2.0 g
3,4-methylenedioxyethylamphetamine (MDEA)	2.0 g
3,4-Methylenedioxymethamphetamine (MDMA)	2.0 g
4-Methylthioamphetamine (4-MTA)	2.0 g
Moramide	2.0 g
Morphine	2.0 g
Opium	20.0 g
Paramethoxyamphetamine (PMA)	2.0 g
Pethidine	10.0 g
Phencyclidine	0.5 g
Psilocin	0.10 g
Psilocybin	0.10 g
Tetrahydrocannabinols including their alkyl homologues except where separately specified; and their corresponding carboxylic acids	2.0 g
3,4,5-Trimethoxyamphetamine (TMA)	2.0 g

SCHEDULE 4**SPECIFIED QUANTITIES FOR PARTICULAR
DANGEROUS DRUGS**

sections 4, 8, 9 and 59 of the Act

Dangerous drug	Quantity of dangerous drug
Amphetamine	200.0 g
Cocaine	200.0 g
Heroin	200.0 g
Lysergide	0.4 g
Methylamphetamine	200.0 g
Phencyclidine	50.0 g

SCHEDULE 5**DANGEROUS DRUGS**

sections 4, 51 and 59 of the Act

Barbituric acid and any 5,5 disubstituted derivatives of barbituric acid, whether or not further substituted at position 1 of the ring

Buprenorphine

Codeine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 30 mg or less of codeine per dosage unit; or
- (b) in undivided preparations containing 1% or less of codeine

Difenoxin except in preparations containing 0.5 mg or less of difenoxin and a quantity of atropine sulphate equivalent to not less than 5% of the dose of difenoxin per dosage unit

Dihydrocodeine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of dihydrocodeine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of dihydrocodeine

Diphenoxylate except in preparations containing 2.5 mg or less of diphenoxylate and a quantity of atropine sulphate equivalent to not less than 1% of the dose of diphenoxylate per dosage unit

Ethylmorphine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of ethylmorphine per dosage unit; or

SCHEDULE 5 (continued)

- (b) in undivided preparations containing 2.5% or less of ethylmorphine

Hydrocodone

Hydromorphone

Ketamine

Methadone

Methylphenidate

Moramide

Morphine

Nicocodine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of nicocodine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of nicocodine

Nicodicodine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of nicodicodine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of nicodicodine

Norcodeine, except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of norcodeine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of norcodeine

Normethadone

Oxycodone

Pentazocine

SCHEDULE 5 (continued)

Pethidine

Phenazocine

Phendimetrazine

Phenmetrazine

Pholcodine except where it is compounded with 1 or more other medicaments in such a way that it cannot be readily extracted and where it is contained—

- (a) in divided preparations containing 100 mg or less of pholcodine per dosage unit; or
- (b) in undivided preparations containing 2.5% or less of pholcodine

Racemethorphan

Racemoramide

Racemorphan

SCHEDULE 6**CONTROLLED SUBSTANCES**

sections 43A and 59 of the Act

1-Chloro-Phenyl-2-Aminopropane

1-Phenyl-2-Chloropropane

1-Phenyl-2-Methylaminopropane

1-Phenyl-2-Nitro propene

Acetic Anhydride

Benzyl Cyanide

Boron Tribromide

Ephedrine

Hydriodic Acid

Hypophosphorous acid

Phenyl Acetic Acid

Phenylpropanolamine

Phenyl-2-Propanone

Phenyl-2-Propanone Oxime

Pseudoephedrine

Pyridine

Red phosphorous

SCHEDULE 7

EXEMPTION CONDITIONS

section 18

1 Seed suppliers

A seed supplier must—

- (a) keep cannabis seeds locked in a secure place when not otherwise required for the purpose of part 5; and
- (b) keep records of the source, quantity and delivery details of all cannabis seeds bought or otherwise obtained and the supply of all seeds to someone else, including to a carrier engaged or employed by the seed supplier to transport the seeds; and
- (c) pack all cannabis seeds to be delivered to someone else by a carrier in a way that ensures, as far as reasonably practicable, the seeds can not be lost if the package is damaged; and
- (d) ensure the only identifying information on the outside of a package containing cannabis seeds to be delivered to someone else by a carrier is information identifying the seed supplier and the person to whom the package is consigned; and
- (e) allow an authorised officer access at any reasonable time to premises the person owns, occupies or uses for the purpose for which the exemption is given and, if asked—
 - (i) give the authorised officer reasonable help in exercising the officer's powers; or
 - (ii) produce to the authorised officer for inspection the records mentioned in paragraph (b).

2 Growers

(1) A grower must—

- (a) keep records of the source, quantity and delivery details of all cannabis seeds obtained from a seed supplier, and, if the seed

SCHEDULE 7 (continued)

supplier engaged or employed a carrier to deliver the seeds to the grower, the name of the person who delivered them; and

- (b) if the grower receives a package containing cannabis seeds and the package appears to have been tampered with—inform an authorised officer that the package appears to have been tampered with as soon as reasonably practicable after receiving it; and
- (c) ensure each season's crop of cannabis plants is—
 - (i) if practicable, planted and harvested in the presence of an authorised officer; and
 - (ii) fenced to the satisfaction of an authorised officer; and
 - (iii) defoliated, or harvested in a way that defoliates the crop, before the crop develops flowering heads; and
- (d) ensure crop residue is ploughed in or otherwise destroyed as soon as practicable, but in any event within 14 days, after harvesting the plants; and
- (e) if the crop fails—ensure any plants left standing are ploughed in or otherwise destroyed; and
- (f) if the crop develops flowering heads before it can be harvested—as soon as practicable after the flowering heads develop, ensure the leaves and flowering heads of the plants are destroyed otherwise than by ploughing them in.

(2) A grower must allow an authorised officer entry to the site used for growing cannabis plants at reasonable times to monitor the following, and if asked, give the authorised officer reasonable help in exercising the officers powers—

- (a) the erection of any signs the authorised officer reasonably requires the grower to erect at the site;
- (b) the planting of the cannabis seeds;
- (c) the conduct of trials, experiments or procedures for the purposes of research;
- (d) crop sampling for tetrahydrocannabinol testing;
- (e) the harvesting of the crop;

SCHEDULE 7 (continued)

(f) the removal or destruction of crop residues or leaves and flowering heads.

(3) A grower must allow an authorised officer at reasonable times—

(a) to examine evidence of the source, physical purity, quality, quantity and location of cannabis seeds held by the grower; and

(b) to examine records kept and operating procedures used for the purpose of research; and

(c) to participate in the collation and publication of trial results; and

(d) to destroy, or supervise the destruction of, cannabis plants if the tetrahydrocannabinol content of plants taken in a random sample of the plants is more than 0.35%; and

(e) to take samples of cannabis seeds and cannabis plant material.

(4) For subsection (3), the grower must produce to the authorised officer, or otherwise allow the authorised officer access to, anything the authorised officer reasonably requires for examination.

(5) If an authorised officer requires, a grower must, after the last planting of cannabis seeds for a particular growing season—

(a) destroy any cannabis seeds not planted during the growing season, in the way and within the time required by the authorised officer; or

(b) give the cannabis seeds to the authorised officer for safe keeping at the Australian Tropical Crops Genetic Resource Centre.

3 Authorised officers

An authorised officer must—

(a) keep cannabis seeds in his or her possession locked in a secure place when not otherwise required for the purpose of part 5; and

(b) keep records of the source, quantity and delivery details of all cannabis seeds obtained from a person who may lawfully possess them, and, if the person engaged or employed another person to deliver the seeds to the authorised officer, the name of the person who delivered them; and

SCHEDULE 7 (continued)

- (c) keep records relating to anything else done by the authorised officer including, for example, samples of cannabis plant material taken and inspections of cannabis plants; and
- (d) deliver cannabis seeds given to him or her by a grower to the Australian Tropical Crops Genetic Resource Centre; and
- (e) keep cannabis plant material taken by the authorised officer for testing in a secure place until it can be delivered to someone else who may lawfully possess the plant material.

4 Researchers**(1) A researcher must—**

- (a) keep records of the source, quantity and delivery details of cannabis seeds obtained from a seed supplier, and, if the seeds are delivered by a carrier, the name of the person who delivered them;
- (b) if the researcher receives a package containing cannabis seeds and the package appears to have been tampered with—inform an authorised officer that the package appears to have been tampered with as soon as reasonably practicable after receiving it; and
- (c) keep cannabis seeds in his or her possession locked in a secure place when not otherwise required for the purpose of part 5; and
- (d) keep cannabis plant material in his or her possession in a secure place when not otherwise required for the purpose of part 5; and
- (e) keep records of all cannabis seeds or cannabis plant material bought or otherwise obtained and the supply or use of all seeds; and
- (f) allow an authorised officer access at any reasonable time to premises the person owns, occupies or uses for the purpose for which the exemption is given and, if asked—
 - (i) give the authorised officer reasonable help in exercising the officer's powers; or
 - (ii) produce to the authorised officer for inspection the records mentioned in paragraph (a).

SCHEDULE 7 (continued)

(2) A researcher must ensure—

- (a) each season's crop of cannabis plants is—
 - (i) if practicable, planted and harvested in the presence of an authorised officer; and
 - (ii) fenced to the satisfaction of the authorised officer; and
 - (iii) if the crop is grown as a field trial—defoliated, or harvested in a way defoliates the crop, before the crop can develop flowering heads; and
- (b) crop residue is ploughed in or otherwise destroyed within 14 days after harvesting the crop; and
- (c) if the crop fails—any cannabis plants left standing are ploughed in or otherwise destroyed; and
- (d) if the crop is grown as a field trial and develops flowering heads before it can be harvested—as soon as practicable after the flowering heads develop, the researcher destroys the leaves and flowering heads of the cannabis plants otherwise than by ploughing them in.

(3) A researcher must allow an authorised officer entry to the site used for the trial planting at reasonable times to monitor—

- (a) the erection of any signs the authorised officer reasonably requires the researcher to erect at the site; or
- (b) the planting of the cannabis seeds; or
- (c) the conduct of trials, experiments or procedures for the purpose of research; or
- (d) crop sampling for tetrahydrocannabinol testing; or
- (e) the harvesting of the crop; or
- (f) the removal or destruction of crop residues.

(4) A researcher must allow an authorised officer at reasonable times—

- (a) to examine evidence of the source, physical purity, quality, quantity and location of cannabis seeds held by the researcher; and

SCHEDULE 7 (continued)

- (b) to examine records kept and operating procedures used for the purpose of research; and
- (c) to participate in the collation and publication of trial results; and
- (d) to destroy, or supervise the destruction of, the cannabis plants if the tetrahydrocannabinol content of plants taken in a random sample of the crop is—
 - (i) for plant breeding—more than 1%; or
 - (ii) for field trials—more than 0.35%; and
- (e) to take samples of cannabis seeds and cannabis plant material.

(5) For subsection (4), the researcher must produce to the authorised officer, or otherwise allow the authorised officer access to, anything the authorised officer reasonably requires for examination.

(6) If an authorised officer requires, a researcher must, after the last planting for a particular growing season—

- (a) destroy any cannabis seeds not planted during the growing season in the way and within the time required by the authorised officer; or
- (b) give the seeds to the authorised officer for safe keeping at the Australian Tropical Crops Genetic Resource Centre.

(7) In this section—

“researcher” means—

- (a) a DPI researcher; or
- (b) a non-DPI researcher.

SCHEDULE 8**PERSONS EXEMPTED UNDER ACT, PART 5B**

section 21

1. Wherewell Pty Ltd of Bellmar, Bowenville Norwin Road, Bowenville, for Lot 4 on RP 54881, Parish of West Prairie, County of Aubigny.
2. Trevor Keith Burt of Wyena Park, Dalby, for Tavoy for Lot 50 on DY 590, Parish of Weale, County of Derby.
3. Selwyn John Rowlings of Department of Primary Industries, 13 Holberton Street, Toowoomba for Formartin Growers' Group, for Lots 419 and 420 on A 34967 and Lot 2 on RP 24747, Parish of Jondaryan, County of Augbiny.

SCHEDULE 9**DICTIONARY**

section 4

“authorised officer”, for part 5, see section 26.

“cannabis plant”, for part 5, means a low level drug content cannabis sativa plant.

“cannabis plant material”, for part 5, means plant material from a low level drug content cannabis sativa plant, other than leaves and flowering heads.

“cannabis seeds”, for part 5, means seeds of cannabis sativa that will produce low level drug content cannabis sativa plants.

“carrier” includes an employee of the carrier.

“commissioner” means the commissioner of the police service.

“DPI researcher” means a public service officer—

- (a) who is employed in the department within which the *Agricultural Standards Act 1994* is administered; and
- (b) whose duties include plant breeding.

“grower” means a person mentioned in schedule 8.

“non-DPI researcher” means a person, other than a DPI researcher, who has, or has access to, facilities that are used or intended to be used by the person for research into the development and use of fibre crops for commercial purposes and is identified in writing by the commissioner as a non-DPI researcher.

“occupier’s notice” means an occupier’s notice referred to in section 7.

“record of proceedings” means a record of proceedings referred to in section 6.

“search warrant” means a search warrant issued under section 18 of the Act.

“seed supplier” means—

SCHEDULE 9 (continued)

- (a) a person who, for trade or commerce, sells or otherwise provides seeds to someone else and is identified in writing by the commissioner as a seed supplier; or
- (b) a public service officer engaged in research or plant breeding who is authorised by the Minister by gazette notice to supply seeds to someone else.

“supply”—

- (a) for part 4, see section 43A¹⁰ of the Act; or
- (b) for part 5, does not include administer.

ENDNOTES

1 Index to endnotes

		Page
2	Date to which amendments incorporated	44
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2 Date to which amendments incorporated

This is the reprint date mentioned in the Reprints Act 1992, section 5(c). Accordingly, this reprint includes all amendments that commenced operation on or before 5 October 2001. Future amendments of the Drugs Misuse Regulation 1987 may be made in accordance with this reprint under the Reprints Act 1992, section 49.

3 Key

Key to abbreviations in list of legislation and annotations

Key	Explanation	Key	Explanation
AIA	= Acts Interpretation Act 1954	prev	= previous
amd	= amended	(prev)	= previously
amdt	= amendment	proc	= proclamation
ch	= chapter	prov	= provision
def	= definition	pt	= part
div	= division	pubd	= published
exp	= expires/expired	R[X]	= Reprint No.[X]
gaz	= gazette	RA	= Reprints Act 1992
hdg	= heading	reloc	= relocated
ins	= inserted	renum	= renumbered
lap	= lapsed	rep	= repealed
notfd	= notified	s	= section
o in c	= order in council	sch	= schedule
om	= omitted	sdiv	= subdivision
orig	= original	SIA	= Statutory Instruments Act 1992
p	= page	SIR	= Statutory Instruments Regulation 1992
para	= paragraph	SL	= subordinate legislation
prec	= preceding	sub	= substituted
pres	= present	unnum	= unnumbered

4 Table of earlier reprints

TABLE OF EARLIER REPRINTS

[If a reprint number includes a roman letter, the reprint was released in unauthorised, electronic form only.]

Reprint No.	Amendments included	Reprint date
1	to reg pubd gaz 6 May 1989	23 April 1993
2	to SL No. 309 of 1996	4 December 1996
2A	to SL No. 303 of 1997	21 November 1997
2B	to SL No. 459 of 1997	14 August 1998
3	to SL No. 348 of 1998	5 February 1999
3A	to SL No. 41 of 1999	8 April 1999
3B	to SL No. 235 of 1999	2 December 1999
3C	to Act No. 28 of 2000	11 August 2000
3D	to SL No. 234 of 2000	15 September 2000
3E	to SL No. 52 of 2001	8 June 2001

5 List of legislation

Drugs Misuse Regulation 1987

made by the Administrator of the Government on 29 October 1987
pubd gaz 31 October 1987 pp 836–47
commenced 31 October 1987 (see s 2)
exempted from application of SIA pt 7 (see SIA sch 2A)

Note—This regulation contains provisions relocated from the Drugs Misuse Act 1986. A list of legislation for the relocated provisions of the Drugs Misuse Act 1986 appears below.

as amended by—

regulations published gazette (pre SL series)—

17 December 1988 pp 2214–5
commenced on date of publication

6 May 1989 pp 230–1
commenced 6 May 1989 (see s 2)

Drugs Misuse Amendment Regulation (No. 1) 1996 SL No. 309

notfd gaz 8 November 1996 pp 959–61
ss 1–2 commenced on date of notification
remaining provisions commenced 8 November 1996 (see s 2)

**List of legislation to Drugs Misuse Act 1986 No. 36 schs 1–6—before relocation to
Drugs Misuse Regulation 1987 as schs 1–6 (see 1996 No. 49 s 21)—****Original relocated Act****Drugs Misuse Act 1986 No. 36 schs 1–6**

date of assent 5 September 1986

ss 1–2 commenced on date of assent

remaining provisions commenced 27 October 1986 (proc pubd gaz 25 October 1986
p 1242)

as amended by—

Drugs Misuse Act Amendment Act 1987 No. 53

date of assent 1 October 1987

ss 1–2 commenced on date of assent

s 10(a)(iii) commenced 6 May 1989 (proc pubd gaz 6 May 1989 p 213)

remaining provisions commenced 31 October 1987 (proc pubd gaz 31 October 1987
p 819)**Drugs Misuse Act Amendment Act 1989 No. 34**

date of assent 28 April 1989

ss 1–2 commenced on date of assent

remaining provisions commenced 6 May 1989 (proc pubd gaz 6 May 1989 p 213)

Statute Law (Miscellaneous Provisions) Act 1990 No. 88 s 3 sch

date of assent 6 December 1990

commenced on date of assent

Drugs Misuse Amendment Act 1995 No. 18

date of assent 11 April 1995

ss 1–2 commenced on date of assent

s 6 commenced 8 December 1995 (1995 SL No. 358)

remaining provisions commenced 12 April 1996 (automatic commencement under
AIA s 15DA(2))**Drugs Misuse Amendment Act 1996 No. 49**

date of assent 15 November 1996

commenced on date of assent

**List of legislation to Drugs Misuse Regulation 1987—after relocation of Drugs
Misuse Act 1986 No. 36 schs 1–6****Drugs Misuse Amendment Regulation (No. 1) 1997 SL No. 303**

notfd gaz 19 September 1997 pp 262–3

commenced on date of notification

Drugs Misuse Amendment Regulation (No. 2) 1997 SL No. 459

notfd gaz 19 December 1997 pp 1770–77

commenced on date of notification

Drugs Misuse Amendment Regulation (No. 1) 1998 SL No. 348

notfd gaz 18 December 1998 pp 1551–7

commenced on date of notification

Drugs Misuse Amendment Regulation (No. 1) 1999 SL No. 41

notfd gaz 26 March 1999 pp 1450–3
commenced on date of notification

Drugs Misuse Amendment Regulation (No. 2) 1999 SL No. 235

notfd gaz 22 October 1999 pp 710–1
commenced on date of notification

Drugs Misuse Amendment Act 2000 No. 28 pt 1 s 26 sch

date of assent 27 July 2000
commenced on date of assent

Drugs Misuse Amendment Regulation (No. 1) 2000 SL No. 234

notfd gaz 8 September 2000 pp 134–5
commenced on date of notification

Drugs Misuse Amendment Regulation (No. 1) 2001 SL No. 52

notfd gaz 25 May 2001 pp 334–6
commenced on date of notification

Drugs Misuse Amendment Regulation (No. 2) 2001 SL No. 174

notfd gaz 21 September 2001 pp 230–31
commenced on date of notification

6 List of annotations

PART 1—PRELIMINARY

pt hdg ins 1996 SL No. 309 s 4

Commencement

s 2 om R2 (see RA s 37)

Repeal

s 3 om R1 (see RA s 40)

Dictionary

prov hdg sub 1996 SL No. 309 s 5(1)

s 4 sub 1998 SL No. 348 s 3(2)

def “**occupier’s notice**” reloc to sch 9 1998 SL No. 348 s 3(1)

def “**record of proceedings**” reloc to sch 9 1998 SL No. 348 s 3(1)

def “**search warrant**” reloc to sch 9 1998 SL No. 348 s 3(1)

def “**supply**” ins 1996 SL No. 309 s 5(3)

om 1998 SL No. 348 s 3(2)

def “**the Act**” sub 1989 reg pubd gaz 6 May 1989 pp 230–1

om 1996 SL No. 309 s 5(2)

PART 2—SEARCH WARRANT NOTICES AND RECORD

pt hdg ins 1996 SL No. 309 s 6

Notice to justice before whom a complaint to ground a search warrant is to be sworn

s 5 amd 1996 SL No. 309 s 7

Record of proceedings

s 6 amd 1996 SL No. 309 s 8

Notice to occupier of place entered pursuant to warrant

s 7 sub reg pubd gaz 17 December 1988 pp 2214–15
amd 1996 SL No. 309 s 9

Forms

s 8 om 1996 SL No. 309 s 10

PART 3—SYRINGES AND DANGEROUS DRUGS DISPOSAL PROCEDURES

pt hdg ins 1996 SL No. 309 s 11

Prescribed procedures for the disposal of hypodermic syringes and needles

s 9 ins reg pubd gaz 6 May 1989 pp 230–1

Prescribed procedure for disposal of dangerous drugs

s 10 ins reg pubd gaz 6 May 1989 pp 230–1

PART 4—CONTROLLED SUBSTANCES

pt 4 (ss 11–15) ins 1996 SL No. 309 s 12

PART 5—TRIAL PLANTING OF CANNABIS SATIVA FOR COMMERCIAL FIBRE PRODUCTION

pt hdg ins 1998 SL No. 348 s 4

Division 1—Preliminary

div hdg ins 1998 SL No. 348 s 4

Purpose of pt 5

s 16 ins 1998 SL No. 348 s 4

Limit of effect of exemption

s 17 ins 1998 SL No. 348 s 4

Conditions

s 18 ins 1998 SL No. 348 s 4

Division 2—Exemptions

div hdg ins 1998 SL No. 348 s 4

Exemptions for seed suppliers

s 19 ins 1998 SL No. 348 s 4

Exemptions for carriers

s 20 ins 1998 SL No. 348 s 4

Exemptions for growers

s 21 ins 1998 SL No. 348 s 4

Exemptions for authorised officer

s 22 ins 1998 SL No. 348 s 4

Exemptions for DPI researchers

s 23 ins 1998 SL No. 348 s 4

Exemptions for non-DPI researchers

s 24 ins 1998 SL No. 348 s 4

Exemptions for employees etc.

s 25 ins 1998 SL No. 348 s 4

Division 3—Other provisions

div hdg ins 1998 SL No. 348 s 4

Appointment of authorised officers

s 26 ins 1998 SL No. 348 s 4

Authorised officer's identity card

s 27 ins 1998 SL No. 348 s 4

Production or display of authorised officer's identity card

s 28 ins 1998 SL No. 348 s 4

Deferral of expiry of Act, pt 5B (Act, s 43Y)

s 29 ins 2001 SL No. 52 s 3

PART 6—TRANSITIONAL PROVISION

prov hdg ins 2001 SL No. 174 s 3

Transitional provision for Drugs Misuse Amendment Regulation (No. 2) 2001

s 30 ins 2001 SL No. 174 s 3

SCHEDULEand reg pubd gaz 17 December 1988 pp 2214–5
om 1996 SL No. 309 s 13**SCHEDULE 1—DANGEROUS DRUGS**sch hdg ins 1997 SL No. 459 s 3(1)
(prev 1986 No. 36 sch 1)
amd 1996 No. 49 s 15
reloc 1996 No. 49 s 21
amd 1997 SL No. 459 s 3(2); 2001 SL No. 174 s 4**SCHEDULE 2—DANGEROUS DRUGS**sch hdg ins 1997 SL No. 459 s 4(1)
(prev 1986 No. 36 sch 2)
amd 1987 No. 53 s 11; 1989 No. 34 s 22; 1996 No. 49 s 16
reloc 1996 No. 49 s 21
amd 1997 SL No. 303 s 3; 1997 SL No. 459 s 4(2)–(3); 1999 SL No. 41 s 3;
2001 SL No. 174 s 5**SCHEDULE 2A—DANGEROUS DRUGS**

ins 2000 No. 28 s 26 sch

SCHEDULE 3—SPECIFIED QUANTITIES FOR PARTICULAR DANGEROUS DRUGSsch hdg ins 1997 SL No. 459 s 5(1)
(prev 1986 No. 36 sch 3)
amd 1987 No. 53 s 12; 1989 No. 34 s 23
sub 1990 No. 88 s 3 sch
amd 1996 No. 49 s 17
reloc 1996 No. 49 s 21
amd 1997 SL No. 303 s 4; 1997 SL No. 459 s 5(2)–(3); 1999 SL No. 41 s 4

SCHEDULE 4—SPECIFIED QUANTITIES FOR PARTICULAR DANGEROUS DRUGS

sch hdg ins 1997 SL No. 459 s 6(1)
 (prev 1986 No. 36 sch 4)
 amd 1996 No. 49 s 18
 reloc 1996 No. 49 s 21
 amd 1997 SL No. 459 s 6(2); 2001 SL No. 174 s 6

SCHEDULE 5—DANGEROUS DRUGS

sch hdg ins 1997 SL No. 459 s 7(1)
 (prev 1986 No. 36 sch 5)
 amd 1987 No. 53 s 13; 1996 No. 49 s 19
 reloc 1996 No. 49 s 21
 amd 1997 SL No. 459 s 7(2)

SCHEDULE 6—CONTROLLED SUBSTANCES

(prev 1986 No. 36 sch 6)
 prev sch 6 om R1 (see RA s 40)
 pres sch 6 ins 1995 No. 18 s 8
 amd 1996 No. 49 s 20
 reloc 1996 No. 49 s 21
 amd 1997 SL No. 459 s 8

SCHEDULE 7—EXEMPTION CONDITIONS

sch hdg ins 1998 SL No. 348 s 5

Seed suppliers

s 1 ins 1998 SL No. 348 s 5

Growers

s 2 ins 1998 SL No. 348 s 5

Authorised officers

s 3 ins 1998 SL No. 348 s 5

Researchers

s 4 ins 1998 SL No. 348 s 5
 amd 1999 SL No. 235 s 3

SCHEDULE 8—PERSONS EXEMPTED UNDER ACT, PART 5B

ins 1998 SL No. 348 s 5
 sub 1999 SL No. 235 s 4; 2000 SL No. 234 s 3

SCHEDULE 9—DICTIONARY

ins 1998 SL No. 348 s 5
 def “**occupier’s notice**” reloc 1998 SL No. 348 s 3(1)
 def “**record of proceedings**” reloc 1998 SL No. 348 s 3(1)
 def “**search warrant**” reloc 1998 SL No. 348 s 3(1)