THE SOUTH AUSTRALIAN

GOVERNMENT GAZETTE

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All instruments appearing in this gazette are to be considered official, and obeyed as such

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GOVERNOR'S INSTRUMENTS

APPOINTMENTS, RESIGNATIONS AND GENERAL MATTERS

Department of the Premier and Cabinet
Adelaide, 7 May 2026

Her Excellency the Governor in Executive Council has been pleased to appoint the undermentioned to the HomeStart Finance Board of Management, pursuant to the provisions of the Urban Renewal Act 1995:

Member: from 7 May 2026 until 6 May 2029
Harry Iraklis Paul Patsouris

By command,

RHIANNON KATE PEARCE, MP
For Premier

T&F26/0008CS

Department of the Premier and Cabinet
Adelaide, 7 May 2026

Her Excellency the Governor in Executive Council has been pleased to appoint the Honourable Leon William Kennedy Bignell as Agent-General for South Australia in London, for a term commencing on 27 May 2026 and expiring on 26 May 2030 - pursuant to the Agent-General Act 1901.

By command,

RHIANNON KATE PEARCE, MP
For Premier

DPC26/016CS

REGULATIONS

South Australia

Controlled Substances (Poisons) Regulations 2026

under the *Controlled Substances Act 1984*

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Part 1—Preliminary

1—Short title

These regulations may be cited as the *Controlled Substances (Poisons) Regulations 2026*.

2—Commencement

These regulations come into operation on 1 September 2026.

3—Interpretation

(1) In these regulations—

Act means the *Controlled Substances Act 1984*;

address means street address;

approved electronic communication means an electronic communication of a kind approved from time to time by the Minister;

approved electronic form, in relation to a prescription for a drug, means—

(a) a form approved from time to time by the Secretary under the Commonwealth Regulations; or

(b) a form approved from time to time by the Minister,

for the giving of prescriptions for drugs in electronic form;

approved information technology requirements means information technology requirements approved from time to time by the Minister;

APVMA means the Australian Pesticides and Veterinary Medicines Authority of the Commonwealth;

Australian jurisdiction means the Commonwealth or a State or Territory of the Commonwealth;

Chief Executive has the same meaning as in the *Health Care Act 2008*;

Commonwealth Regulations means the *National Health (Pharmaceutical Benefits) Regulations 2017* of the Commonwealth;

correctional institution has the same meaning as in the *Correctional Services Act 1982*;

council has the same meaning as in the *Local Government Act 1999*;

council subsidiary means a subsidiary of a council established under the *Local Government Act 1999*;

data source entity means any of the following:

(a) eRx Script Exchange Pty Ltd;

(b) Medication Knowledge Pty Ltd;

(c) the National Prescription Delivery Service;

(d) any other prescription exchange service operating in an Australian jurisdiction;

dental hygienist means a person registered under the *Health Practitioner Regulation National Law*—

(a) to practise in the dental profession (other than as a student); and

(b) in the dental hygienists division of that profession;

dental therapist means a person registered under the *Health Practitioner Regulation National Law*—

(a) to practise in the dental profession (other than as a student); and

(b) in the dental therapists division of that profession;

diesel fuel means a petroleum or shale product used or capable of being used in propelling a diesel engined motor vehicle;

dispense means to supply a drug in accordance with a prescription for that drug;

domestic partner means a person who is a domestic partner within the meaning of the *Family Relationships Act 1975*, whether declared as such under that Act or not;

drug means a poison designed for human or animal therapeutic use;

electronic communication has the same meaning as in the *Electronic Communications Act 2000*;

electronic prescription means a prescription given in an approved electronic form;

enrolled nurse means a person registered under the *Health Practitioner Regulation National Law*—

(a) to practise in the nursing and midwifery profession as a nurse (other than as a student); and

(b) in the enrolled nurses division of that profession;

fax includes e-fax;

health service facility means a hospital, nursing home or other facility at which a health service is provided for the public or any section of the public for the purpose of curing, alleviating, diagnosing or preventing the spread of any mental or physical illness, disease, injury, abnormality or disability;

information technology requirements has the same meaning as in the *Electronic Communications Act 2000*;

liquefied petroleum gas means a hydrocarbon fluid composed predominantly of any of the following hydrocarbons or mixtures of all or any of them:

(a) propane (C_3H_8);

(b) propylene (C_3H_6);

(c) butane (C_4H_{10});

(d) butylene (C_4H_8);

medication chart prescription has the same meaning as in the Commonwealth Regulations;

Metropolitan Adelaide means Metropolitan Adelaide as defined by General Registry Office Plan 639/93;

monitored drug means any of the following:

(a) any S8 poison;

(b) any S4 poison that is a benzodiazepine;

(c) any S4 poison that contains Codeine;

(d) any of the following S4 poisons:

(i) Gabapentin;

(ii) Pregabalin;

(iii) Quetiapine;

(iv) Tramadol;

(v) Zolpidem;

(vi) Zopiclone;

monitored drugs database means an electronic database kept by the Department that contains information relating to the sale, supply, prescription, administration and use of monitored drugs;

motor spirit means petrol or another petroleum or shale product used or capable of being used in propelling a motor vehicle (other than diesel fuel or liquefied petroleum gas);

National Health Act means the *National Health Act 1953* of the Commonwealth;

National Health (Continued Dispensing) Determination means the determination of that name, as in force from time to time, made under section 89A(3) of the National Health Act;

optometrist means a person registered under the *Health Practitioner Regulation National Law* to practise in the optometry profession (other than as a student);

oral health therapist means a person registered under the *Health Practitioner Regulation National Law*—

(a) to practise in the dental profession (other than as a student); and

(b) in the oral health therapists division of that profession;

petroleum product means a volatile solvent comprised of—

(a) motor spirit; or

(b) diesel fuel; or

(c) liquefied petroleum gas;

pharmaceutical benefit has the same meaning as in Part VII of the National Health Act;

podiatrist means a person registered under the *Health Practitioner Regulation National Law* to practise in the podiatry profession (other than as a student);

poison—see regulation 5;

Poisons Standard means the current Poisons Standard as defined in the Commonwealth Act and as modified by deleting Sections 54(1) and (2), 57, 58, 59, 60 and 64 and Appendices B, D and J;

prescribed (continued dispensing) pharmaceutical benefit means a pharmaceutical benefit listed in the National Health (Continued Dispensing) Determination as a pharmaceutical benefit that may be supplied under section 89A of the National Health Act by approved pharmacists without a prescription;

prescriber means a person who lawfully gives a prescription for a drug;

record means—

(a) a documentary record; or

(b) a record made by an electronic, electromagnetic, photographic or optical process; or

(c) any other kind of record;

registered nurse means a person registered under the *Health Practitioner Regulation National Law*—

(a) to practise in the nursing and midwifery profession as a nurse (other than as a student); and

(b) in the registered nurses division of that profession;

scheduled medicine means a medicine that contains a substance included in a schedule of the Poisons Standard;

S4 drug means—

(a) an S4 poison; or

(b) a substance designed for human or animal therapeutic use that has been approved by—

(i) the TGA for inclusion in the Australian Register of Therapeutic Goods; or

(ii) APVMA for inclusion in the Public Chemical Registration Information System (PUBCRIS),

but has not yet been—

(c) listed in some other schedule of the Poisons Standard; or

(d) exempted from listing in the Poisons Standard;

section 22 poison means a poison to which section 22 of the Act applies by virtue of regulation 27;

spouse—a person is the spouse of another if they are legally married;

TGA means the Therapeutic Goods Administration of the Commonwealth;

Vaccine Administration Code means the document of that name published by the Department as in force from time to time.

(2) In these regulations, a reference to an ***S1 poison*** is a reference to a poison listed in Schedule 1 of the Poisons Standard, a reference to an ***S2 poison*** is a reference to a poison listed in Schedule 2 of the Poisons Standard, and so on.

(3) In these regulations, ***incorporated hospital*** and ***SAAS*** have the same respective meanings as in the *Health Care Act 2008*.

(4) For the purposes of these regulations—

(a) an electronic prescription for a drug is ***presented*** when it is accessed electronically for the purpose of dispensing the drug; and

(b) a prescription for a drug given to a pharmacist by fax is ***presented*** when a faxed copy of the prescription is transmitted to the pharmacy at which the drug is to be dispensed.

4—Application of regulations

These regulations do not apply in relation to—

(a) a poison when contained in a product that is listed in Appendix A of the Poisons Standard; or

(b) a poison listed in Appendix G of the Poisons Standard when contained in a preparation in a concentration not exceeding the concentration specified in Appendix G for that poison; or

(c) a poison that is listed in any of the Schedules 1 to 6 (but is not listed in Schedule 7 or 8) of the Poisons Standard when contained in a preparation in a concentration not exceeding 10 milligrams per litre or 10 milligrams per kilogram.

Part 2—Controlled substances

5—Declaration of poisons (section 12(1) of Act)

(1) Pursuant to section 12(1) of the Act, the following substances (whether in a pure form, or contained in a preparation or admixture) are declared to be poisons:

(a) the primary substances listed in Schedules 1 to 8 and Schedule 10 of the Poisons Standard;

(b) section 17A, 17B and 17C precursors;

(c) the following related substances, but subject to any express exclusion contained in the Poisons Standard:

(i) the artificial form of a primary substance;

(ii) if a primary substance is a plant (other than a plant included in Schedule 8 of the Poisons Standard)—that plant, or any part of that plant, when packed or prepared for therapeutic use;

(iii) every salt, active principle or derivative (including an ester or ether) of a primary substance and every salt of such an active principle or derivative;

(iv) every alkaloid of a primary substance and every salt of such an alkaloid;

(v) every stereoisomer of a primary substance and every salt of such a stereoisomer.

(2) A related substance will be taken to be included in the Schedule, or Schedules, of the Poisons Standard in which the primary substance to which it is related is included.

(3) A reference in these regulations to a poison will be taken to include a reference to the primary substance and its related substances (in each case whether in a pure form, or contained in a preparation or admixture).

6—Declaration of prescription drugs (section 12(2) of Act)

Pursuant to section 12(2) of the Act, S4 poisons and S8 poisons are declared to be prescription drugs.

7—Declaration of drugs of dependence (section 12(3) of Act)

Pursuant to section 12(3) of the Act, S8 poisons are declared to be drugs of dependence.

8—Declaration of volatile solvents (section 12(7) of Act)

Pursuant to section 12(7) of the Act, the following are declared to be volatile solvents:

(a) the following substances (whether in their natural or artificial form):

Acetone (dimethyl ketone, propanone)

Amyl nitrite (isopentyl nitrite)

Bromochlorodifluoromethane (BCF)

Butane

Butanone (methyl ethyl ketone)

Butyl nitrite

Carbon tetrachloride

Chlorofluorocarbons and fluorocarbons except where separately specified

Chloroform

Dichloromethane (methylene chloride)

Diethyl ether (ethoxyethane)

Dimethyl ether (methoxymethane)

Enflurane

Ethyl acetate

Ethyl chloride (chloroethane)

Halothane

Heptane

Hexane

Isoamyl nitrite

Isobutane (2-methylpropane)

Isobutyl nitrite (2-methylpropyl nitrite)

Isoflurane

Methoxyflurane

Methyl acetate

Methyl isobutyl ketone (4-methylpentan-2-one)

Methyl tert-butyl ether

Nitrous oxide

Octane

Octyl nitrite

Pentane

Petrol

Propane

Sevoflurane

Tetrachloroethylene (perchloroethylene, tetrachloroethene)

Toluene (methylbenzene)

1,1,1-trichloroethane (methylchloroform)

Trichloroethylene (trichloroethene)

Xylene (xylol);

(b) structural isomers of a substance specified in paragraph (a);

(c) preparations or admixtures containing any proportion of a substance specified in paragraph (a);

(d) preparations or admixtures containing any proportion of structural isomers of a substance specified in paragraph (a).

Part 3—Application of Part 4 of Act (general offences)

9—Manufacture, production and packing (section 13 of Act)

Section 13 of the Act applies to all S1 poisons, S2 poisons, S3 poisons and S7 poisons.

10—Exemption from section 13 of Act

The holder of a licence under the Commonwealth Act to manufacture goods is exempt from the requirement to hold a licence under section 13 of the Act in respect of the manufacture of those goods.

11—Sale by wholesale (section 14 of Act)

Section 14 of the Act applies to all S1 poisons, S2 poisons, S3 poisons and S7 poisons.

12—Sale or supply to end user (section 15 of Act)

(1) Subject to subregulation (2), section 15 of the Act applies to all S1 poisons, S2 poisons, S3 poisons and S7 poisons.

(2) Section 15 of the Act does not apply in relation to—

(a) adrenaline (epinephrine), when supplied by a council subsidiary or health service facility for administration to a person as part of an immunisation program delivered by the council, council subsidiary or health service facility; or

(b) an S3 poison in circumstances where the poison is sold by retail or supplied, for purposes relating to the provision of first aid, in accordance with a scheme approved by the Minister for the purposes of this subregulation.

13—Directions to be given for safe and proper use of S3 poisons sold by retail etc

(1) Subject to subregulation (2), a person who sells by retail or supplies an S3 poison must personally (not through an assistant) give oral directions, supplemented wherever practicable with written directions, for the safe and proper use of the poison to the person purchasing or being supplied with the poison.

Maximum penalty: $3 000.

(2) An interpreter may be used to assist in the giving of oral directions if the person purchasing or being supplied with the poison is not sufficiently familiar with the English language.

14—Special provisions relating to sale or supply of pseudoephedrine

(1) A person must not sell or supply pseudoephedrine unless a prescribed identification document or a birth certificate is produced by the person to whom the pseudoephedrine is to be sold or supplied.

Maximum penalty: $3 000.

(2) A person who sells or supplies pseudoephedrine must make and keep a record of the following information:

(a) the name and address of the person to whom the pseudoephedrine is being sold or supplied;

(b) the form of prescribed identification document produced by the person to whom the pseudoephedrine is being sold or supplied;

(c) the unique identification number (if any) on the prescribed identification document produced;

(d) the date of the sale or supply;

(e) the directions given for the safe and proper use of the pseudoephedrine;

(f) the trade name or the approved name of the pseudoephedrine being sold or supplied, or, if it does not have either a trade name or approved name, its ingredients and the form, strength and quantity sold or supplied;

(g) a unique identifier enabling those records to be linked with the pseudoephedrine sold or supplied.

Maximum penalty: $3 000.

(3) Subregulations (1) and (2) do not apply in relation to—

(a) the sale of pseudoephedrine by wholesale; or

(b) the sale or supply of pseudoephedrine in the course of professional practice by—

(i) a pharmacist in a hospital; or

(ii) a registered health practitioner other than a pharmacist; or

(iii) a veterinary surgeon.

(4) A person who makes a record under subregulation (2) must keep it in an electronic form that is accessible via the internet by the Chief Executive and the Commissioner of Police.

Maximum penalty: $3 000.

(5) In this regulation—

Australian student identification card means a card issued by an Australian educational institution to identify a person studying at the institution;

birth certificate of a person means a certified copy of, or extract from, a register of births kept under an Australian law, or under the law of the country in which the person was born;

driver's licence means—

(a) a driver's licence issued under the *Motor Vehicles Act 1959*; or

(b) an interstate licence, interstate learner's permit or foreign licence within the meaning of that Act;

prescribed identification document means a current—

(a) driver's licence; or

(b) firearms licence; or

(c) passport (other than an Australian passport); or

(d) proof of age card; or

(e) Australian student identification card,

that bears a photograph of the holder;

proof of age card means a proof of age card issued by the Registrar of Motor Vehicles or by a corresponding public authority of another State or a Territory of the Commonwealth.

15—Sale of certain poisons (section 16 of Act)

(1) Section 16 of the Act applies to all S7 poisons.

(2) For the purposes of section 16(4) of the Act, a person who sells S7 poisons must keep records of such matters as are specified in Part 2 Section 56(1) of the Poisons Standard.

16—Declaration of precursors (sections 17A, 17B and 17C of Act)

(1) Section 17A of the Act applies to the following poisons:

1-Chlorophenyl-2-aminopropane

3,4-Methylenedioxyphenylpropan-2-one (PMK)

1-Phenyl-2-bromopropane

1-Phenyl-1-chloro-2-methylaminopropane

1-Phenyl-2-chloropropane

1-Phenyl-2-iodopropane

1-Phenyl-2-nitropropene.

(2) Section 17B of the Act applies to the following poisons:

Chemical name	Alternative name	CAS number
Acetic anhydride		108-24-7
4-Allylpyrocatechol	2-Hydroxychavicol	1126-61-0
alpha Phenylacetonitrile	alpha Acetyl Phenylacetonitrile	4468-48-8
4-Amino-butanoic acid	Piperidinic acid	56-12-2
Anethole	trans-Anethole	4180-23-8
		104-46-1
Bromobenzene	Phenylbromide	108-86-1
Bromosafrole		38589-39-8
Boron tribromide		10294-33-4
1,4-Butanediol	Tetramethylene Glycol	110-63-4
1-Chlorophenyl-2-aminopropane		
Ephedrine (including salts)	L-Ephedrine	50-98-6
Ethyl phenylacetate	Benzene acetic acid, ethyl ester	101-97-3
Gamma butyrolactone		96-48-0
Gamma hydroxybutanoic acid (including salts)	Gamma hydroxybutyric acid	
Hydriodic acid	Hydrogen iodide	10034-85-2
4-Hydroxybutanal	4-Hydroxybutyraldehyde	5371-52-8
2-Hydroxytetrahydrofuran	Tetrahydro-2-furanol	1346-46-9
4-Hydroxybutanoic acid lactone	Gamma-valerolactone	9648-0
4-Hydroxybutanoic acid nitrile	4-Hydroxybutyronitrile	628-22-8
4-Hydroxypentanoic acid	Gamma Valerolactone	108-29-2
Hypophosphite salts		
Hypophosphorous acid	Phosphinic acid	6303-21-5
Lithium aluminium hydride	LAH	16853-85-3
Methcathinone	Ephedrone	
3,4-Methylenedioxy-phenylacetic acid	1,3-Benzodioxolo-5-acetic acid	2861-28-1
3,4-Methylenedioxyphenylpropan-2-one		4676-39-5
N-Methylephedrine		552-79-4
Methyl phenylacetate	Benzeneacetic acid, methyl ester	101-41-7
N-Methylpseudoephedrine		51018-28-1
Norpseudoephedrine		53643-20-2
2-Pyrrolidone	Gamma-butyrolactam	616-45-5
Phenylacetamide		103-81-1

Phenylacetic acid (including salts)		103-82-2
Phenylacetonitrile	Benzyl cyanide/Benzeneacetonitrile/ Benzyl nitrile	140-29-4
Phenylacetyl chloride		103-80-0
1-Phenyl-2-bromopropane	(+-)-2-Bromo-1-phenylpropane	2114-39-8
1-Phenyl-2-chloropropane		
1-Phenyl-2-iodopropane	(2-Iodopropyl)benzene	29527-87-5
1-Phenyl-2-nitropropene		
Phenylpropanolamine	Norephedrine	37577-28-9
1-Phenyl-2-propanone	Benzyl methyl ketone, Phenylacetone	103-79-7
1-Phenyl-2-propanone oxime		
1-Phenyl-2-propanol		14898-87-4
2-Phenyl-propanal	Hydratropic aldehyde	93-53-8
Phosphorus		7723-14-0
Phosphorous acid	Phosphonic Acid	10294-56-1
1-Phenyl-1-propanone	Phenylethylketone, Propiophenone	99-55-0
Piperonal	3,4-Methylenedioxy-benzaldehyde, Heliotropine	120-57-0
Pseudoephedrine (including salts)		
Pyridine		110-86-1
Safrole	5-(2-Propenyl)-1,3-Benzodioxide	94-59-7
Sassafras oil		8006-80-2
Sodium bis(2-methoxyethoxy) aluminium hydride	Sodium dihydrido-bis(2-methoxyethoxy) aluminate	22722-98-1
Sodium cyanoborohydride	Sodium borocyanohydride	25895-60-7

(3) Section 17C of the Act applies to the following poisons:

Acetaldehyde

N-Acetylanthranilic acid

Allylbenzene

Ammonium formate

Anthranilic acid

Benzaldehyde

1,3-benzodioxole

Benzyl bromide

Benzyl chloride

5-bromo-1,3-benzodioxole

Chromic acid (including salts)

Ergometrine

Ergotamine

Ethanamine

N-Ethylephedrine

N-Ethylpseudoephedrine

Eugenol

Formaldehyde

Formamide

Hydrobromic acid

Iodine (including iodide salts)

Isosafrole

Lithium

Lysergic acid

Magnesium

Mandelic acid

Mercuric chloride

Mercury

Methylamine

Methylammonium salts

N-methylformamide

Nitroethane

Nitromethane

Palladium (including salts)

Phenylalanine

Piperidine

Platinum

Potassium

Propionic anhydride

Raney nickel

Sodium

Sodium borohydride

Thionyl chloride

Thorium (including salts)

Trans-beta-methylstyrene.

17—End user statement for precursors (sections 17B and 17C of Act)

For the purposes of sections 17B(1)(c) and 17C(1)(a) of the Act, the form of end user statement in Schedule 1 is prescribed.

18—Regulation of prescription drugs—prescription by pharmacists of certain prescription drugs (section 18(1)(a)(iii) of Act)

(1) For the purposes of section 18(1)(a)(iii) of the Act, a pharmacist is authorised to prescribe a prescription drug (not being a drug of dependence) if—

(a) the pharmacist is included on the register of authorised pharmacists; and

(b) the drug is prescribed in accordance with the Community Pharmacist Prescribing Code.

(2) The Minister may authorise a pharmacist for the purpose of this regulation (an ***authorised pharmacist***) if the Minister is satisfied that the pharmacist meets the criteria and requirements set out by the Community Pharmacist Prescribing Code.

(3) The Minister must cause a register of authorised pharmacists to be kept and made available to the public on a website operated by, or on behalf of, the Department.

(4) In this regulation—

Community Pharmacist Prescribing Code means the document of that name published by the Department, as in force from time to time.

19—Regulation of prescription drugs—administration of certain S4 drugs (section 18(1d)(a)(iii) of Act)

(1) For the purposes of section 18(1d)(a)(iii) of the Act, a dental hygienist, dental therapist, oral health therapist or podiatrist is authorised to administer any of the following S4 drugs:

Articaine

Benzocaine

Bupivacaine

Levobupivacaine

Lidocaine

Mepivacaine

Prilocaine

Ropivacaine.

(2) For the purposes of section 18(1d)(a)(iii) of the Act, an optometrist is authorised to administer any of the following S4 drugs:

Eye drops containing 0.5% or less of tetracaine

Eye drops containing 1.0% or less of atropine sulfate monohydrate

Eye drops containing 1.0% or less of cyclopentolate hydrochloride

Eye drops containing 2.0% or less of homatropine hydrobromide

Eye drops containing 0.5% or less of lidocaine

Eye drops containing 0.5% or less of oxybuprocaine

Eye drops containing 2.0% or less of pilocarpine nitrate

Eye drops containing 0.5% or less of proxymetacaine

Eye drops containing 1.0% or less of tropicamide.

Note—

An optometrist's registration may also be endorsed under section 94 of the *Health Practitioner Regulation National Law* as being qualified to administer a scheduled medicine or class of scheduled medicines—see section 18(1d)(a)(ii) of the Act.

(3) For the purposes of section 18(1d)(a)(iii) of the Act, a registered health practitioner of a class determined by the Minister may administer a prescription drug (not being a drug of dependence) to a person if—

(a) the registered health practitioner has successfully completed a training program approved by the Minister from time to time for the purposes of this subregulation; and

(b) the drug is listed in the Vaccine Administration Code or is a drug approved by the Minister from time to time for the purposes of this subregulation; and

(c) the drug is administered as part of—

(i) an immunisation program delivered by—

(A) an incorporated hospital; or

(B) SAAS; or

(C) a council or council subsidiary; or

(ii) an immunisation program delivered by an organisation approved by the Minister for the purposes of this subregulation; and

(d) the drug is administered in accordance with—

(i) the Vaccine Administration Code; and

(ii) —

(A) in the case of a drug administered as part of the *National Immunisation Program—the National Immunisation Program Schedule* and the *Australian Immunisation Handbook*; or

(B) in any other case—requirements specified by the Minister.

(4) For the purposes of section 18(1d)(a)(iii) of the Act, a pharmacist may administer an S4 drug that is a vaccine to a person if—

(a) the pharmacist has successfully completed an immunisation education or training program accredited for pharmacists by—

(i) Australian Pharmacy Council Ltd; or

(ii) Health Education Services Australia Pty Ltd; or

(iii) any other person or body approved by the Minister for the purposes of this subregulation; and

(b) the drug is administered in accordance with—

(i) if instructions for the administration of the drug are in the *Australian Immunisation Handbook*—the *Australian Immunisation Handbook*; or

(ii) in any other case—requirements specified by the Minister.

(5) In this regulation—

Australian Immunisation Handbook means *The Australian Immunisation Handbook* published by the Commonwealth Department of Health and Ageing, as in force from time to time;

National Immunisation Program Schedule means the *National Immunisation Program Schedule* published by the Commonwealth Department of Health and Ageing, as in force from time to time.

20—Regulation of prescription drugs—prescription of certain S4 drugs by medical practitioners (section 18(2) of Act)

(1) For the purposes of section 18(2) of the Act—

(a) each of the S4 drugs listed in column 1 of the table below, when used for the purpose set out in column 2, is a prescribed prescription drug; and

(b) the qualifications and requirements specified in that table alongside a drug or list of drugs in column 3 are prescribed qualifications and requirements.

	Prescription drug	Use	Qualifications and requirements
1	Clomifene Cyclofenil Follitropin alpha (recombinant human follicle stimulating hormone) Follitropin beta (recombinant human follicle stimulating hormone) Luteinising hormone Urofollitrophin (follicle stimulating hormone)	Human use	A medical practitioner who— (a) is registered in the specialty of endocrinology or obstetrics and gynaecology; or (b) provides services to a fertility unit, an endocrinology unit or obstetrics and gynaecology unit of a teaching hospital in South Australia.
2	Acitretin Bexarotene Etretinate	Human use	A medical practitioner who— (a) is registered in the specialty of dermatology, oncology or haematology; or (b) is a medical registrar working under the supervision of a medical practitioner referred to in paragraph (a); or (c) is registered in some other specialty and is authorised by the Minister to prescribe such drugs.

3	Isotretinoin	Human internal use	A medical practitioner who— (a) is registered in the specialty of dermatology, oncology or haematology; or (b) is a medical registrar working under the supervision of a medical practitioner referred to in paragraph (a); or (c) is registered in some other speciality and is authorised by the Minister to prescribe such drugs.
4	Tretinoin	Human internal use	A medical practitioner who— (a) is registered in the specialty of oncology or haematology; or (b) is a medical registrar working under the supervision of a medical practitioner referred to in paragraph (a); or (c) is registered in some other speciality and is authorised by the Minister to prescribe such drugs.
5	Lenalidomide Pomalidomide Thalidomide	Human use	A medical practitioner who— (a) is registered in the specialty of oncology or haematology; or (b) is a medical registrar working under the supervision of a medical practitioner referred to in paragraph (a); or (c) is authorised by the Minister to prescribe such drugs.
6	Ambrisentan Bosentan Macitentan Sitaxentan	Human use	A medical practitioner who— (a) is registered as a specialist; or (b) is a medical registrar who is working under the supervision of a medical practitioner referred to in paragraph (a); or (c) is authorised by the Minister to prescribe such drugs.

7	Enzalutamide	Human use	A medical practitioner who— (a) is registered as a specialist; or (b) is a medical registrar who is working under the supervision of a medical practitioner referred to in paragraph (a); or (c) is authorised by the Minister to prescribe such drugs.
8	Riociguat	Human use	A medical practitioner who— (a) is registered as a specialist; or (b) is a medical registrar who is working under the supervision of a medical practitioner referred to in paragraph (a); or (c) is authorised by the Minister to prescribe such drugs.

(2) A medical practitioner who prescribes an S4 drug listed in the table in subregulation (1) (other than in item 1) must—

(a) inform the patient of the name of the drug and that the drug may cause birth defects; and

(b) provide the patient with written information about the drug and its potential side effects; and

(c) inform the patient of the dangers should the patient unlawfully supply the drug to another person; and

(d) if the patient is a female of child-bearing age—

(i) ensure that the possibility of pregnancy has been excluded prior to commencement of treatment; and

(ii) inform her that she must not become pregnant during treatment or within the prescribed period after completion of treatment; and

(e) obtain written consent for the treatment from the patient.

Maximum penalty: $5 000.

(3) In this regulation—

prescribed period means—

(a) in the case of treatment with a drug listed in item 2 of the table in subregulation (1) (other than bexarotene)—24 months;

(b) in the case of treatment with bexarotene or a drug listed in item 3, 4, 5 or 8 of that table—1 month;

(c) in the case of treatment with a drug listed in item 6 or 7 of that table—3 months.

21—Regulation of prescription drugs—prescription of certain S8 poisons by medical practitioners (section 18(2) of Act)

(1) For the purposes of section 18(2) of the Act—

(a) each of the S8 poisons listed in column 1 of the table below, when used for the purpose set out in column 2, is a prescribed prescription drug; and

(b) the qualifications and requirements specified in that table alongside a drug in column 3 are prescribed qualifications and requirements.

	Prescription drug	Use	Qualifications and requirements
1	N,α-dimethyl-3,4-(methylenedioxy)phenylethylamine (MDMA)	Human use, for the treatment of post-traumatic stress disorder	A medical practitioner— (a) who is registered in the specialty of psychiatry; and (b) for whom an authority under section 19(5) of the Commonwealth Act that covers MDMA is in force.
2	Psilocybine (Psilocybin)	Human use, for the treatment of treatment-resistant depression	A medical practitioner— (a) who is registered in the specialty of psychiatry; and (b) for whom an authority under section 19(5) of the Commonwealth Act that covers psilocybine is in force.

(2) A medical practitioner who prescribes an S8 poison listed in the table in subregulation (1) must, within 1 business day of prescribing the poison and for the purposes of the Chief Psychiatrist performing the Chief Psychiatrist's functions under the *Mental Health Act 2009*, give notice to the Chief Psychiatrist—

(a) in a form determined by the Chief Psychiatrist; and

(b) containing such information as the Chief Psychiatrist may determine.

22—Exemptions from section 18 of Act

(1) A council, council subsidiary or health service facility is exempt from section 18(1c)(d) of the Act in respect of the supply of an S4 drug under an immunisation program run by the council, council subsidiary or health service facility.

(2) A pharmacist who sells or supplies an S4 drug without dispensing a prescription is exempt from section 18(1b)(a) and (1c)(a) of the Act in relation to that sale or supply if—

(a) the drug is sold or supplied to a council, council subsidiary or health service facility for use in an immunisation program delivered by the council, council subsidiary or health service facility and the pharmacist has received a written order for the drug from the council, council subsidiary or health service facility; or

(b) the drug is for use by a person who holds a licence to sell, supply or administer an S4 drug and the pharmacist has received a written order for the drug from the licensee; or

(c) the drug is sold or supplied for the mass treatment of certain animals to the owner of the animals and—

(i) the pharmacist has received a written order for the drug from a veterinary surgeon; or

(ii) —

(A) the drug is an antibiotic; and

(B) the pharmacist has received a written order for the drug from an inspector appointed under the *Livestock Act 1997*; and

(C) the written order is on a form approved by the Chief Inspector of Stock under that Act and has been countersigned by the Chief Inspector; or

(d) the drug is sold or supplied to a registered health practitioner or veterinary surgeon authorised to sell, supply or administer S4 drugs and the pharmacist has received a written order for the drug from that practitioner or veterinary surgeon; or

(e) the drug is authorised or required by the law of any place to be carried on board a ship and the pharmacist has received a written order for the drug from the master or medical officer of the ship; or

(f) the drug is not one listed in the table in regulation 20(1) for the purposes of section 18(2) of the Act and the pharmacist—

(i) is satisfied that—

(A) the person for whom it is to be sold or supplied is being medically treated with the drug; and

(B) the continued sale or supply of that drug is essential to the health of that person; and

(C) there is good reason for the person's inability to produce a prescription for the drug; and

(ii) sells or supplies—

(A) where the pharmacist is satisfied that the person for whom it is to be sold or supplied is affected by an emergency specified by the Minister by notice under subregulation (3) and the sale or supply occurs during the period specified in relation to that emergency in the same notice—

- for drugs that are on the Pharmaceutical Benefits Scheme—no more than the standard Pharmaceutical Benefits maximum quantity; or
- for drugs that are not on the Pharmaceutical Benefits Scheme—the quantity that is contained in the smallest standard pack in which the drug is generally available; or

(B) in any other case—

- if the drug is a cream, ointment or liquid or one that is packaged in such a manner as to promote the safe and proper use of the drug—the smallest standard package or container made by the manufacturer; or

- if the drug is not a cream, ointment or liquid or other drug described above—no more than 3 days dosage of the drug; and

(iii) on the day on which the drug is sold or supplied, records—

(A) their name as the seller or supplier of the drug; and

(B) the date; and

(C) the trade name or the approved name of the drug, or, if it does not have either a trade name or approved name, its ingredients; and

(D) the name and address of the person for whom the drug is sold or supplied; and

(E) the form, strength and quantity of the drug; and

(F) the directions given for the safe and proper use of the drug, including (where appropriate) the route of administration of the drug; or

(g) the drug is a prescribed (continued dispensing) pharmaceutical benefit and the sale or supply is made in accordance with the conditions specified in the National Health (Continued Dispensing) Determination; or

(h) the drug is sold or supplied to a person in accordance with a scheme determined by the Minister from time to time for the purposes of this paragraph by notice in the Gazette.

(3) If the Minister is satisfied that an emergency (within the meaning of the *Emergency Management Act 2004*) is occurring in an area of the State, the Minister may, by notice in the Gazette, specify the emergency and a period of time in relation to that emergency for the purposes of subregulation (2)(f)(ii)(A).

(4) A notice under subregulation (3)—

(a) may specify the emergency by reference to any factor the Minister thinks fit (including, without limitation, by a description of the circumstances of the emergency, the area within which the emergency is occurring or by any description by which the emergency is commonly known); and

(b) may, from time to time as the Minister thinks fit, be varied or substituted by a new notice or be revoked.

(5) The holder of a licence under the Commonwealth Act to manufacture goods is exempt from section 18(1e) of the Act in respect of the manufacture of those goods.

(6) In this regulation—

Pharmaceutical Benefits Scheme means the Pharmaceutical Benefits Scheme under the National Health Act.

23—Exemptions from section 18A of Act

(1) A registered health practitioner authorised to prescribe or supply a drug of dependence is exempt from section 18A(1) of the Act in respect of the prescription or supply of such a drug for use by a person in respect of whom a section 18A authority exists if—

(a) in the case of a person who is receiving treatment in a hospital or correctional institution—

(i) the registered health practitioner notifies the holder of the section 18A authority that the practitioner has prescribed or supplied the drug or is intending to prescribe or supply the drug in respect of that person; and

(ii) the drug is only administered to the person while in the hospital or correctional institution; and

(iii) if the drug is solely for the treatment of drug dependence—the dose administered does not exceed the dose authorised; or

(b) in the case of a person who is being discharged from a hospital following treatment in the hospital—

(i) the registered health practitioner notifies the holder of the section 18A authority that the practitioner has prescribed or supplied the drug or is intending to prescribe or supply the drug in respect of that person; and

(ii) if the drug is solely for the treatment of drug dependence—the dose prescribed does not exceed the dose authorised; or

(c) in the case of a person not referred to in paragraph (a) or (b)—

(i) the registered health practitioner prescribing or supplying the drug—

(A) notifies the holder of the section 18A authority that the practitioner has prescribed or supplied the drug or is intending to prescribe or supply the drug in respect of that person; or

(B) is a medical practitioner (including a locum for the time being substituting for such a practitioner) in the same practice as the holder of the section 18A authority; and

(ii) the registered health practitioner prescribing or supplying the drug does so with the approval of the holder of the section 18A authority; and

(iii) the registered health practitioner prescribing or supplying the drug complies with the section 18A authority relating to the person for whom the drug is prescribed or to whom the drug is supplied.

(2) A registered health practitioner authorised to prescribe or supply a drug of dependence is exempt from section 18A(1) of the Act in relation to the prescription or supply of such a drug for a person in respect of whom a section 18A authority does not exist if—

(a) the drug (not being dextromoramide or pethidine) is for use by a person aged 70 years or more; or

(b) the drug (not being dextromoramide or pethidine) is for use by a person whose life expectancy is reasonably believed by the registered health practitioner principally responsible for treatment of the person, to be less than 12 months and—

(i) the registered health practitioner has informed the Minister of the person's name and address, date of birth and the nature of the condition for which the drug is prescribed; and

(ii) the prescription for the drug is endorsed either "Notified Palliative Care Patient" or "NPCP"; or

(c) the drug is for use by a person who is receiving treatment in a hospital or correctional institution and the duration of treatment of the person with the drug while the person is in the hospital or correctional institution does not exceed 14 days; or

(d) the drug is for use by a person who is being discharged from a hospital following treatment in the hospital and the duration of treatment of the person with the drug after discharge does not exceed 14 days; or

(e) —

(i) the drug is prescribed or supplied by a registered health practitioner solely for the treatment of drug dependence and as part of a drug treatment service program administered by an incorporated hospital; and

(ii) an application for a section 18A authority is made in respect of the person by a registered health practitioner practising at the drug treatment service program within 5 business days of the drug being prescribed or supplied to the person.

(3) In this regulation—

section 18A authority means an authority granted by the Minister to a registered health practitioner under section 18A of the Act to prescribe or supply a drug of dependence.

24—Sale or supply of volatile solvents (section 19 of Act)

(1) Section 19(3) of the Act applies to—

(a) nitrous oxide; and

(b) volatile solvents that are petroleum products.

(2) For the purposes of section 19(3) of the Act, the age prescribed is—

(a) in the case of nitrous oxide—18 years; or

(b) in the case of a volatile solvent that is a petroleum product—16 years.

(3) A person is exempt from section 19(3) of the Act in respect of the supply of nitrous oxide to a person who is under the age of 18 years if—

(a) the first person lawfully carries on a business of selling nitrous oxide; and

(b) the second person is an employee of the first person; and

(c) the supply is in the ordinary course of business.

(4) The Minister may, by notice in the Gazette, exempt any person from section 19(3) of the Act in respect of the sale or supply of nitrous oxide subject to such conditions (if any) as the Minister thinks fit.

(5) A person who sells or supplies a volatile solvent for use as an inhalant in medical or dental treatment is exempt from section 19 of the Act in respect of that sale or supply.

25—Special provisions relating to retail sale of nitrous oxide

(1) A person must not sell a substance that is, or purports to be, nitrous oxide by retail between the hours of 10 pm and 5 am on the following day.

Maximum penalty: $5 000.

(2) A person who sells a substance that is, or purports to be, nitrous oxide from premises by retail must ensure that the substance is—

(a) stored in a part of the premises to which members of the public are not permitted access; and

(b) stored in such a way that it is not visible to members of the public at the premises.

Maximum penalty: $5 000.

(3) A person who sells a substance that is, or purports to be, nitrous oxide from premises by retail must display a notice that complies with the requirements in subregulation (4) in a manner and position that is likely to attract the attention of customers.

Maximum penalty: $2 500.

Expiation fee: $315.

(4) A notice displayed under subregulation (3) must comply with the following requirements:

(a) the notice must display the following words:

IT IS UNLAWFUL TO SELL OR SUPPLY NITROUS OXIDE TO PERSONS UNDER THE AGE OF 18 YEARS. PERSONS MAY BE REQUIRED TO PRODUCE EVIDENCE OF AGE WHEN MAKING A PURCHASE;

(b) the words required to be displayed must appear on the notice in legible letters or numerals not less than 15 millimetres in height and be of a colour that contrasts with the background colour of the notice.

26—Automatic vending machines (section 20 of Act)

(1) Section 20(1) of the Act does not apply to—

(a) an S5 poison that is sold or supplied by means of an automatic vending machine located at a car washing facility provided that the first aid instructions, warning statements and safety directions for the poison specified in the Poisons Standard are displayed at the facility; or

(b) the following products sold or supplied by means of an automatic vending machine:

(i) condoms with or without spermicides or viricides;

(ii) lubricants with or without spermicides or viricides; or

(c) injecting equipment sold or supplied by way of an automatic vending machine at a location and site approved by the Minister; or

(d) an unscheduled medicine sold or supplied by way of an automatic vending machine provided that—

(i) the medicine is sold or supplied in the original unopened pack supplied by the manufacturer; and

(ii) the medicine is sold or supplied in a pack that contains not more than 2 adult doses of the medicine; and

(iii) the automatic vending machine is presented and located in such a way that makes unsupervised access by children unlikely.

(2) In this regulation—

injecting equipment means—

(a) alcohol swabs, needles, syringe filters, syringes, tourniquets, water for injection or winged infusion sets; or

(b) a kit containing 1 or more of the items specified in paragraph (a);

unscheduled medicine means a medicine that is included in the Australian Register of Therapeutic Goods and is not a scheduled medicine.

27—Possession of poisons (section 22 of Act)

(1) Section 22 of the Act applies to the following poisons:

4-aminopropiophenone

Acrolein

Arsenic as an S7 poison

Chloropicrin

Cyanides as S7 poisons

Cyanogen

DDT

Fluoroacetamide

Fluoroacetic acid

Hydrocyanic acid as an S7 poison

Methyl bromide

Mirex

Sodium fluoroacetate

Strychnine as an S7 poison

Thallium.

(2) A person is exempt from section 22 of the Act in respect of the possession of 4-aminopropiophenone if—

(a) —

(i) in the case of 4-aminopropiophenone that is contained in a capsule for use with a Canid Pest Ejector—the concentration of 4-aminopropiophenone in each cartridge does not exceed the constituent concentration specified in the Public Chemical Registration Information System (PUBCRIS) by APVMA; or

(ii) in the case of 4-aminopropiophenone that is a constituent of baits designed for destroying vertebrate animals—the concentration of 4-aminopropiophenone in each bait does not exceed 2%; and

(b) the total amount of 4-aminopropiophenone in the particular quantity of capsules or baits does not exceed 5 kilograms; and

(c) the person—

(i) has the written approval of the Minister to acquire and possess those capsules or baits; and

(ii) acquires the capsules or baits from a supplier approved by the Minister.

(3) A person is exempt from section 22 of the Act in respect of the possession of sodium fluoroacetate if—

(a) —

(i) in the case of sodium fluoroacetate that is contained in a capsule for use with a Canid Pest Ejector designed for destroying foxes or wild dogs—the concentration of sodium fluoroacetate in each capsule does not exceed 0.8%; or

(ii) in the case of sodium fluoroacetate that is a constituent of baits designed for destroying vertebrate animals—the concentration of sodium fluoroacetate in each bait does not exceed 0.04%; or

(iii) in the case of sodium fluoroacetate that is contained in a cartridge for use with a Felixer grooming trap—the concentration of sodium fluoroacetate in each cartridge does not exceed the constituent concentration specified in the Public Chemical Registration Information System (PUBCRIS) by APVMA; and

(b) the total amount of sodium fluoroacetate present in the particular quantity of capsules, cartridges or baits does not exceed 50 grams; and

(c) the person—

(i) has the written approval of the Minister to acquire and possess those capsules, cartridges or baits; and

(ii) acquires the capsules, cartridges or baits from a supplier approved by the Minister.

(4) A person is exempt from section 22 of the Act in respect of the possession of strychnine if—

(a) the person is the owner or occupier, or an agent or employee of an owner or occupier, of land that is situated outside a township and outside Metropolitan Adelaide; and

(b) the strychnine is a constituent of baits designed for destroying mice; and

(c) the quantity of baits in the person's possession does not exceed 5 kilograms; and

(d) the amount of strychnine present in any quantity of the baits does not exceed 0.5%.

(5) A person lawfully in possession of baits containing strychnine under this regulation must not use those baits except for the purpose of destroying mice in or around storage areas on land situated outside a township and outside Metropolitan Adelaide.

Maximum penalty: $3 000.

(6) The Minister may exempt a person who is licensed under the *Controlled Substances (Pesticides) Regulations 2017* from the requirement to hold a licence under section 22 of the Act in respect of the use of a pesticide that is a section 22 poison.

28—Packaging and labelling of poisons (section 24 of Act)

(1) For the purposes of section 24(b) of the Act, the package or container—

(a) must comply with the requirements set out in the Poisons Standard; and

(b) in the case of a package or container for an S2 poison, S3 poison, S4 poison or S8 poison, must—

(i) be impervious to, and incapable of chemical reaction with, the poison when the package or container is under conditions of temperature and pressure that are likely to be encountered in normal use; and

(ii) have sufficient strength and impermeability to prevent leakage of the poison during handling, transport and storage of the package or container under normal handling conditions; and

(iii) in the case of a package or container intended to be opened more than once—be able to be securely and readily closed and reclosed; and

(iv) in the case of a prescribed medicine—comply with the packaging requirements of Therapeutic Goods Order No 95.

(2) For the purposes of section 24(c) of the Act, a package or container in which a poison for human or animal therapeutic use is sold by retail on prescription, or is supplied on prescription—

(a) must have affixed to it a label that complies with Appendix L Clause 1 of the Poisons Standard; and

(b) must, in the case of a poison that is listed in column 1 of Appendix L Clause 2 of the Poisons Standard have affixed to it a label that contains the warning statements prescribed for that poison by Appendix F Clause 1 of that Standard; and

(c) must, in the case of a preparation for internal use by humans that contains a poison listed in Appendix K of the Poisons Standard, have affixed to it a label that contains the sedation warning statement 39, 40 or 90 as specified in Appendix F Clause 1 of that Standard.

(3) For the purposes of section 24(c) of the Act, a package or container in which a prescribed S3 poison is sold by retail, or is supplied—

(a) must have affixed to it a label that—

(i) complies with Appendix L Clause 1 of the Poisons Standard; and

(ii) in the case of pseudoephedrine—contains a unique identifier enabling that poison to be linked with the records required to be kept under regulation 14; and

(b) must, in the case of a preparation for internal use by humans that contains a poison listed in Appendix K of the Poisons Standard, have affixed to it a label that contains the sedation warning statement 39, 40 or 90 as specified in Appendix F Clause 1 of that Standard.

(4) For the purposes of section 24(c) of the Act, a package or container in which a poison designed for human or animal therapeutic use (other than a prescribed S3 poison) is sold by retail or is supplied—

(a) must have affixed to it the label appearing on the package or container for the poison as supplied by the manufacturer (being a label that complies with the Poisons Standard); or

(b) must have affixed to it—

(i) a label that complies with Appendix L Clause 1 of the Poisons Standard; and

(ii) in the case of a preparation for internal use by humans that contains a poison listed in Appendix K of that Standard—a label that contains the sedation warning statement 39, 40 or 90 as specified in Appendix F Clause 1 of that Standard.

(5) For the purposes of section 24(c) of the Act, a package or container in which a poison (other than a poison designed for human or animal therapeutic use or a prescribed S3 poison) is sold by retail or is supplied (other than on prescription) must have affixed to it a label that complies with the Poisons Standard.

(6) A registered health practitioner or veterinary surgeon who is authorised to prescribe, sell or supply a prescribed medicine is exempt from the requirement to comply with the packaging requirements of Therapeutic Goods Order No 95 in relation to the sale or supply of that prescribed medicine to a particular person if the registered health practitioner or veterinary surgeon believes that the person would suffer undue hardship through difficulty in opening a container that complies with the requirements of that Order.

(7) The Minister may grant an exemption from specified requirements of section 24(b) or (c) of the Act to a seller or supplier in respect of a particular product if the Minister is satisfied that the product is otherwise adequately packaged or labelled.

(8) The Minister may grant a seller or supplier, or a class of sellers or suppliers, an exemption from subregulation (1)(b)(iv) in relation to specified packaging requirements of Therapeutic Goods Order No 95 for a specified prescribed medicine.

(9) In this regulation—

prescribed medicine means—

(a) a medicine that contains a substance listed in Schedule 1 to Therapeutic Goods Order No 95 or a salt, ester or other derivative of such a substance; or

(b) a product that—

(i) contains a substance listed in Schedule 1 to Therapeutic Goods Order No 95 or a salt, ester or other derivative of such a substance; and

(ii) is intended solely for use in animals;

prescribed S3 poison means any of the following S3 poisons:

(a) dihydrocodeine in cough preparations;

(b) pseudoephedrine;

Therapeutic Goods Order No 95 means *Therapeutic Goods Order No. 95 - Child-resistant packaging requirements for medicines 2017* made under the Commonwealth Act on 29 November 2017, as in force from time to time.

29—Storage of poisons (section 25 of Act)

For the purposes of section 25 of the Act, the following requirements apply:

(a) a person must not store a poison in a container that—

(i) is normally used for containing food or beverages; or

(ii) is similar to a container that is normally used for containing food or beverages;

(b) a person must not store an S2 poison in premises where such a poison is sold by retail unless—

(i) it is stored in a part of the premises to which the public is not permitted access; or

(ii) if it is stored in a part of the premises to which the public is permitted access, it—

(A) is stored not less than 1.2 metres above floor level; or

(B) is enclosed in—

- a child-resistant package; or
- a blister pack; or
- a container approved by the Minister; or

(C) is stored in a container that has a capacity of not less than 5 litres; or

(D) is stored in a container that has a gross weight of not less than 5 kilograms;

(c) a person must not store an S3 poison or S4 poison in premises where such a poison is sold by retail unless it is stored in a part of the premises to which the public is not permitted access;

(d) a person must not store an S4 poison (other than a poison which has been lawfully dispensed to the person) in premises where such a poison is supplied or administered unless it is stored in a part of the premises to which the public is not permitted access;

(e) a person must not store an S6 poison or S7 poison in premises where such a poison is sold by retail except in accordance with the requirements of Part 2 Section 54(3) and (4) of the Poisons Standard;

(f) a person must not store a drug of dependence except in accordance with the requirements of the *Code of Practice for the Storage and Transport of Drugs of Dependence*, published by the Department, as in force from time to time;

(g) a person must not store pentobarbital in injectable preparations except in a locked container.

30—Consignment of poisons for transport

A person must not—

(a) consign a poison for transport unless it is packed in such a way as to avoid leakage arising from the ordinary risks of handling and transport; or

(b) consign for transport a drug of dependence except in accordance with the requirements of the *Code of Practice for the Storage and Transport of Drugs of Dependence*, published by the Department, as in force from time to time.

Maximum penalty: $5 000.

31—Transport of poisons (section 26 of Act)

For the purposes of section 26 of the Act, a person must not—

(a) transport an S2 poison, S3 poison, S4 poison or S8 poison in a vehicle in which any food, or component of food, for human or animal consumption is being transported unless that poison is carried in a part of the vehicle effectively separated from that part of the vehicle containing the food; or

(b) transport a drug of dependence except in accordance with the requirements of the *Code of Practice for the Storage and Transport of Drugs of Dependence*, published by the Department, as in force from time to time.

32—Prohibition on use of certain poisons for certain purposes (section 27 of Act)

(1) For the purposes of section 27 of the Act, a person must not sell, supply, purchase or use an S7 poison for a domestic purpose or domestic gardening purpose.

(2) For the purposes of section 27 of the Act, a person must not sell, supply, prescribe or use a poison listed in Schedule 10 of the Poisons Standard for the purpose or purposes indicated in relation to that poison in that Schedule (other than amygdalin for human therapeutic use).

33—Prohibition on use of certain poisons

(1) A person must not sell, supply, prescribe or use amygdalin for human therapeutic use unless—

(a) special access to amygdalin has been authorised in accordance with the requirements of sections 18 and 31A of the Commonwealth Act and regulation 12A of the *Therapeutic Goods Regulations 1990* made under that Act; and

(b) permission for the importation of amygdalin (subject to special access authorisation) has been granted under regulation 5H and Schedule 8 item 12AA of the *Customs (Prohibited Imports) Regulations 1956* of the Commonwealth.

Maximum penalty: $5 000.

(2) A person must not—

(a) prescribe, sell, supply or purchase a poison produced for the treatment of animals if the person knows, or if there are reasonable grounds for suspecting, that the poison is intended for human use; or

(b) administer to any person (including themselves) a poison produced for the treatment of animals; or

(c) use choramphenicol for the treatment of stock bred, raised or used for the purpose of providing a product for human consumption.

Maximum penalty: $5 000.

(3) In this regulation—

stock means—

(a) a bird or other animal; or

(b) a bee of the genus *Apis* or *Megachile*.

34—Restrictions on advertising (section 28 of Act)

(1) Section 28 of the Act applies to—

(a) all poisons listed in Schedule 10 of the Poisons Standard; and

(b) all S3 poisons other than those listed in Appendix H of the Poisons Standard; and

(c) all S4 poisons and S8 poisons; and

(d) all controlled drugs other than drugs of dependence.

(2) A person is exempt from section 28 of the Act if—

(a) the person only publishes an advertisement of a poison in a journal that is circulated predominantly among registered health practitioners, medical administrators, scientists working in medical laboratories or persons who are licensed to sell the poison by wholesale; or

(b) the person only publishes an advertisement of a poison that consists of a price list that complies with the *Price Information Code of Practice*, published by the TGA, as in force from time to time.

(3) In this regulation—

journal means a newsletter, magazine or other periodical, whether published for sale or for distribution without charge.

Part 4—Prescriptions and dispensing

35—How prescriptions are to be given

(1) Subject to this regulation, a prescriber must give a prescription for a drug—

(a) in writing; or

(b) in an approved electronic form.

Maximum penalty: $5 000.

(2) A prescriber may, if of the opinion that good reason exists for doing so, give a prescription for a drug to a pharmacist by—

(a) telephone; or

(b) fax; or

(c) an approved electronic communication.

(3) If a prescriber gives a prescription in writing, the prescriber must give the prescription to—

(a) in the case of a prescription for a drug for human use—

(i) the person for whom the drug is to be supplied; or

(ii) a person acting on behalf of the person for whom the drug is to be supplied; or

(b) in the case of a prescription for a drug for animal use—

(i) the owner of the animal; or

(ii) a person acting on behalf of the owner of the animal.

Maximum penalty: $5 000.

(4) If a prescription is given in an approved electronic form, the prescriber must—

(a) in the case of a prescription in a form approved by the Secretary under the Commonwealth Regulations—prepare and submit the prescription in accordance with any approved information technology requirements (as defined in the Commonwealth Regulations) by means of an eligible electronic communication (as defined in the Commonwealth Regulations); or

(b) in the case of a prescription in a form approved by the Minister—prepare and submit the prescription in accordance with approved information technology requirements (if any) by means of an approved electronic communication.

Maximum penalty: $3 000.

(5) If a prescription is prepared in an approved electronic form, the prescriber must include in the prescription—

(a) the date on which the prescription is given; and

(b) the prescriber's professional name, address and telephone number; and

(c) the full name and address of the person for whom the prescription is intended; and

(d) the name, dose form and (if relevant) the route of administration of the drug being prescribed; and

(e) if applicable—the strength of the drug being prescribed; and

(f) the dose of the drug to be administered to the person for whom the drug is being prescribed; and

(g) the frequency at which the drug is to be administered; and

(h) the total amount of the drug to be supplied each time the prescription is dispensed; and

(i) the total number of times the drug may be dispensed; and

(j) if the prescription is for a drug of dependence for human use—the date of birth of the person for whom the prescription is intended; and

(k) the words—

(i) "For dental treatment only" if the prescriber is a dentist; or

(ii) "For podiatric treatment only" if the prescriber is a podiatrist; or

(iii) "For animal treatment only" if the prescriber is a veterinary surgeon.

Maximum penalty: $3 000.

(6) If a prescription for a monitored drug for human use is prepared in an approved electronic form, the prescriber must—

(a) keep a record of—

(i) the details required by subregulation (5) to be included in the prescription; and

(ii) the date of birth of the person for whom the prescription has been prepared; and

(b) transmit that record electronically to a data source entity at, or immediately following, the time the record is created (unless subregulation (7) applies).

Maximum penalty: $5 000.

Expiation fee: $1 250.

(7) If a prescriber is unable to transmit a record relating to a prescription in accordance with subregulation (6)(b) because the electronic system used to keep the record is not compatible with the electronic system of a data source entity, the prescriber must transmit the record electronically to the Chief Executive so that it is received no later than—

(a) if the prescription is prepared on a day falling within the first 14 days of a month—the 21st day of that month; or

(b) if the prescription is prepared on any other day—the 7th day of the month following the month in which the prescription was prepared; or

(c) in any case, such later day as the Chief Executive may, on application, authorise.

Maximum penalty: $5 000.

Expiation fee: $1 250.

(8) The Minister may exempt a prescriber or class of prescribers from the operation of subregulation (6)(b) or (7) (or both) if satisfied that proper cause exists for the exemption.

(9) If a prescription is given to a pharmacist by telephone, the prescriber must give the pharmacist—

(a) the prescriber's professional name, address and telephone number; and

(b) the full name and address of the person for whom the prescription is intended (or, if the prescription is intended for an animal, the species of animal for which it is intended, the name and address of the owner of the animal and the name (if any) of the animal); and

(c) the name, dose form and (if relevant) the route of administration of the drug being prescribed; and

(d) if applicable—the strength of the drug being prescribed; and

(e) the dose of the drug to be administered to—

(i) the person for whom the drug is being prescribed; or

(ii) the animal in relation to which the drug is being prescribed,

(as the case may be);

(f) the total amount of the drug to be supplied; and

(g) the frequency at which the drug is to be administered; and

(h) if the prescription is for a drug of dependence for human use—the date of birth of the person for whom the prescription is intended.

Maximum penalty: $3 000.

(10) If a prescription is given to a pharmacist by telephone—

(a) the prescriber must, immediately after so giving the prescription, complete a prescription in writing that—

(i) clearly states that it is given in confirmation of the prescription given by telephone on the particular date on which it is so given; and

(ii) otherwise complies with these regulations; and

(b) the prescriber must forward the written prescription to the pharmacist—

(i) if the prescription is for a drug of dependence—within 24 hours of giving the prescription by telephone; or

(ii) in any other case—as soon as practicable after giving the prescription by telephone.

Maximum penalty: $3 000.

(11) If a prescription is given to a pharmacist by fax, the prescriber must forward the original prescription to the pharmacist—

(a) in the case of a prescription for a drug of dependence—within 24 hours of giving the prescription by fax; or

(b) in any other case—as soon as practicable after giving the prescription by fax,

unless the prescriber has endorsed the prescription given by fax with the name and address of a single pharmacy at which the prescription may be dispensed.

Maximum penalty: $3 000.

(12) If a prescription is given to a pharmacist by an approved electronic communication, the prescriber must comply with any requirements imposed by the Minister.

Maximum penalty: $3 000.

(13) This regulation does not apply to a prescriber who gives a prescription for a drug if—

(a) the prescription is a medication chart prescription; and

(b) the provisions of the Commonwealth Regulations applying to the giving of a medication chart prescription for a pharmaceutical benefit are complied with in relation to the giving of the prescription of the drug (whether or not the drug is a pharmaceutical benefit).

36—Written prescriptions

(1) A prescriber who writes a prescription for the supply of a drug must—

(a) date the prescription with the date on which the prescription is written and sign the prescription; and

(b) include on the prescription—

(i) their professional name, address and telephone number; and

(ii) the full name and address of the person for whom the prescription is intended or, if the prescription is intended for an animal, the species of animal for which it is intended, the name and address of the owner of the animal and the name (if any) of the animal; and

(iii) the words—

(A) "For dental treatment only" if the prescriber is a dentist; or

(B) "For podiatric treatment only" if the prescriber is a podiatrist; or

(C) "For animal treatment only" if the prescriber is a veterinary surgeon; and

(c) specify on the prescription—

(i) the name, dose form and (if relevant) the route of administration of the drug being prescribed; and

(ii) if applicable—the strength of the drug; and

(iii) the dose of the drug to be administered to the person for whom, or the animal for which, it is prescribed; and

(iv) the frequency at which the drug is to be administered; and

(v) the total amount of the drug to be supplied each time the prescription is dispensed; and

(vi) the total number of times the drug may be dispensed; and

(d) if the prescription is for a drug of dependence for human use, comply with the following additional requirements:

(i) include on the prescription the date of birth of the person for whom the prescription is intended;

(ii) express the total amount of the drug to be specified under subparagraph (c)(v) in both words and numerals.

Maximum penalty: $3 000.

(2) A prescriber who writes a prescription for the supply of a monitored drug for human use—

(a) must keep a record of—

(i) the details required to be included and specified under subregulation (1); and

(ii) the date of birth of the person for whom the prescription has been written; and

(b) if the record is kept in electronic form—must transmit that record electronically to a data source entity at, or immediately following, the time the record is created (unless subregulation (3) applies).

Maximum penalty: $5 000.

Expiation fee: $1 250.

(3) If a prescriber is unable to transmit a record relating to a prescription in accordance with subregulation (2)(b) because the electronic system used to keep the record is not compatible with the electronic system of a data source entity, the prescriber must transmit the record electronically to the Chief Executive so that it is received no later than—

(a) if the prescription is written on a day falling within the first 14 days of a month—the 21st day of that month; or

(b) if the prescription is written on any other day—the 7th day of the month following the month in which the prescription was prepared; or

(c) in any case, such later day as the Chief Executive may, on application, authorise.

Maximum penalty: $5 000.

Expiation fee: $1 250.

(4) The Minister may exempt a prescriber or class of prescribers from the operation of subregulation (2)(b) or (3) (or both) if satisfied that proper cause exists for the exemption.

(5) If a prescriber writes a prescription for an above average strength or potentially dangerous dose of a drug, the prescriber must—

(a) underline the statement of the dose of the drug on the prescription; and

(b) sign their initials alongside the underlined portion of the prescription referred to in paragraph (a).

Maximum penalty: $3 000.

(6) This regulation does not apply to a person who writes a prescription for a drug if—

(a) the prescription is a medication chart prescription; and

(b) the provisions of the Commonwealth Regulations applying to the giving of a medication chart prescription for a pharmaceutical benefit are complied with in relation to the giving of the prescription for the drug (whether or not the drug is a pharmaceutical benefit).

37—Giving prescriptions for monitored drugs—special provisions

(1) Before a prescriber gives a prescription for the supply of a monitored drug for human use (whether the prescription is given in writing, in an approved electronic form, by telephone, by fax or by an approved electronic communication), the prescriber must take all reasonable steps to check relevant information held in the monitored drugs database relating to the person for whom the drug is to be prescribed.

Maximum penalty: $5 000.

(2) Subregulation (1) does not apply if—

(a) the drug (not being dextromoramide or pethidine) is for use by a person aged 70 years or more; or

(b) the drug (not being dextromoramide or pethidine) is for use by a person whose life expectancy is reasonably believed by the registered health practitioner principally responsible for treatment of the person, to be less than 12 months and—

(i) the prescriber has informed the Minister of the person's name and address, date of birth and the nature of the condition for which the drug is prescribed; and

(ii) the prescription for the drug is endorsed either "Notified Palliative Care Patient" or "NPCP"; or

(c) the drug is for use by a person who is receiving treatment in a hospital or a correctional institution and the duration of treatment of the person with the drug while the person is in the hospital or correctional institution does not exceed 14 days; or

(d) the drug is for use by a person who is being discharged from a hospital following treatment in the hospital and the duration of treatment of the person with the drug after discharge does not exceed 14 days.

(3) Subregulation (1) does not apply to a person who gives a prescription for a drug if—

(a) the prescription is a medication chart prescription; and

(b) the provisions of the Commonwealth Regulations applying to the giving of a medication chart prescription for a pharmaceutical benefit are complied with in relation to the giving of the prescription for the drug (whether or not the drug is a pharmaceutical benefit).

38—Dispensing prescriptions

(1) If a pharmacist or medical practitioner dispenses a drug pursuant to a prescription, the pharmacist or medical practitioner must—

(a) in the case of a written prescription or electronic prescription—record in or on the prescription—

(i) the pharmacist's or medical practitioner's name, business name (if any) and business address; and

(ii) the date on which the drug is dispensed; and

(iii) the unique identifier applicable to the drug; or

(b) in the case of a prescription given by fax that is endorsed with the name and address of a single pharmacy at which the prescription is to be dispensed—endorse on the faxed copy—

(i) the pharmacist's or medical practitioner's name, business name (if any) and business address; and

(ii) the date on which the drug is dispensed; and

(iii) the unique identifier applicable to the drug.

Maximum penalty: $5 000.

(2) A pharmacist or medical practitioner who dispenses a drug pursuant to a prescription must, on the day on which the drug is dispensed, record the following information:

(a) the unique identifier applicable to the drug dispensed on the prescription;

(b) the name of the pharmacist or medical practitioner as the dispenser;

(c) the date on which the drug is dispensed;

(d) the trade name or the approved name of the drug, or, if it does not have either a trade or approved name, the ingredients of the drug;

(e) if the drug is dispensed for a person—

(i) the full name and address of the person; and

(ii) in the case of a monitored drug—the person's date of birth;

(f) if the drug is intended for an animal—the species of animal for which it is intended, the name and address of the owner of the animal and the name (if any) of the animal;

(g) the form, strength and quantity of the dispensed drug;

(h) the directions given for the safe and proper use of the dispensed drug;

(i) the name, address and business telephone number of the person who prescribed the drug;

(j) the number of times the prescription may be dispensed and (if the prescription so specifies) the intervals at which the drug may be dispensed;

(k) any instructions the prescriber has included in or on the prescription in relation to a specialised supply of the drug;

(l) if the prescription is endorsed for dispensing at a single pharmacy—the name and address of that pharmacy.

Maximum penalty: $5 000.

(3) A pharmacist or medical practitioner must not do the following:

(a) in the case of a prescription for an S4 poison that does not specify the number of times the drug is to be dispensed—dispense the drug more than once pursuant to the prescription;

(b) in the case of a prescription that specifies the number of times and the intervals at which the drug may be dispensed—dispense the drug more times than the number specified or at intervals less than those specified;

(c) in the case of a prescription that specifies the number of times, but not the intervals at which, the drug may be dispensed—dispense the drug more frequently than the pharmacist or medical practitioner considers appropriate.

Maximum penalty: $5 000.

(4) Despite subregulation (3)(b), if a pharmacist or medical practitioner is satisfied that a person—

(a) has lost a previously dispensed supply of a drug; or

(b) will, through absence from the State or otherwise, find it unduly difficult to have future supplies of a drug dispensed as needed,

the pharmacist or medical practitioner may (but is not obliged to) dispense a prescription for the person at an interval earlier than that specified in or on the prescription.

(5) If, under subregulation (4), a pharmacist or medical practitioner dispenses a drug of dependence at an earlier interval than that specified in or on the prescription, the pharmacist or practitioner must notify the prescriber of that fact in writing.

Maximum penalty: $5 000.

(6) If a prescription given by fax is endorsed with the name and address of a single pharmacy at which the drug may be dispensed, a pharmacist must not dispense the drug unless the pharmacist is on duty at that pharmacy.

Maximum penalty: $5 000.

(7) A pharmacist or medical practitioner must not dispense a drug if—

(a) the prescription for the drug—

(i) is presented or otherwise sought to be dispensed—

(A) in the case of a drug of dependence—more than 6 months after the date on which it was written; or

(B) in any other case—more than 12 months after the date on which it was written; or

(ii) has been cancelled; or

(iii) is partly or wholly illegible; or

(iv) does not comply with the Act or these regulations; or

(b) there are reasonable grounds for suspecting that the prescription has been altered, forged or obtained by false pretences.

Maximum penalty: $5 000.

(8) If a prescription for a drug that is to be dispensed for the first or only time is given in writing, a pharmacist or medical practitioner must not dispense the drug unless the original written prescription for the drug is presented to the pharmacist or medical practitioner.

Maximum penalty: $5 000.

(9) If a prescription for a drug that is to be dispensed for the first or only time is given by fax, a pharmacist or medical practitioner must not dispense the drug unless the faxed prescription is endorsed with the name and address of a single pharmacy at which the drug may be dispensed.

Maximum penalty: $5 000.

(10) If a prescription for a drug that is to be dispensed for the second or subsequent time is given in writing, a pharmacist or medical practitioner must not dispense the drug unless—

(a) the original written prescription for the drug and a written record (whether made on the prescription or on a separately attached repeat authorisation) of the number of times the drug has been dispensed are presented to the pharmacist or medical practitioner; or

(b) a duplicate or copy of the written prescription for the drug and a written record (made both on the duplicate or copy (as the case may be) and on a separately attached repeat authorisation) of the number of times the drug has been dispensed are presented to the pharmacist or medical practitioner.

Maximum penalty: $5 000.

(11) If a pharmacist or medical practitioner—

(a) dispenses a drug pursuant to a written prescription; and

(b) the drug is fully dispensed,

the pharmacist or medical practitioner must endorse the prescription with the word "CANCELLED" on the day on which the drug is dispensed.

Maximum penalty: $5 000.

(12) If a pharmacist—

(a) dispenses a drug pursuant to a prescription given by fax that is endorsed with the name of a single pharmacy at which the prescription may be dispensed; and

(b) the drug is fully dispensed,

the pharmacist must endorse the faxed copy of the prescription with the word "CANCELLED" on the day on which the drug is dispensed.

Maximum penalty: $5 000.

(13) If a pharmacist or medical practitioner—

(a) dispenses a drug pursuant to an electronic prescription; and

(b) the drug is fully dispensed,

the pharmacist or medical practitioner must record in or on the prescription, on the day that the prescription is dispensed, that the prescription is cancelled.

(14) A pharmacist or medical practitioner who dispenses a prescription for an S4 poison must, unless the prescription is for any reason forwarded to the Department or the Minister—

(a) in the case of a written prescription—

(i) retain the original or duplicate prescription for at least 1 year; and

(ii) keep the original or duplicate prescription readily available for inspection by an authorised officer during that period; or

(b) in the case of a prescription given by fax—

(i) retain the faxed copy of the prescription for at least 1 year; and

(ii) keep the faxed copy of the prescription readily available for inspection by an authorised officer during that period; or

(c) in the case of an electronic prescription—

(i) retain the electronic prescription or a computer-generated printed copy of it for at least 1 year; and

(ii) keep the electronic prescription or a computer-generated printed copy of it readily available for inspection by an authorised officer during that period.

Maximum penalty: $5 000.

(15) If a prescription has been issued in duplicate and the original is retained by the pharmacist or medical practitioner, it is sufficient compliance with this regulation if the required information is marked on the duplicate prescription.

(16) For the purposes of this regulation, a prescription for a drug is ***fully dispensed*** if—

(a) in the case of a prescription authorising dispensing of the drug once only—the drug has been dispensed on 1 occasion; or

(b) in the case of a prescription authorising dispensing of the drug more than once—the drug has been dispensed for the last time.

(17) This regulation (other than subregulations (2), (7)(a) and (7)(b)) does not apply to a pharmacist or medical practitioner who dispenses a drug on a prescription if—

(a) the prescription is a medication chart prescription; and

(b) the provisions of the Commonwealth Regulations applying to the sale or supply of a pharmaceutical benefit have been complied with in relation to the sale or supply of the drug (whether or not the drug is a pharmaceutical benefit).

39—Dispensing prescriptions for drugs of dependence and other monitored drugs—special provisions

(1) A pharmacist who dispenses a monitored drug on prescription must—

(a) each time that the drug is dispensed—make a record in electronic form that complies with regulation 38(2); and

(b) transmit that record electronically to a data source entity at, or immediately following, the time the record is created (unless subregulation (2) applies).

Maximum penalty: $5 000.

Expiation fee: $1 250.

(2) If a pharmacist is unable to transmit a record relating to a prescription in accordance with subregulation (1)(b) because the electronic system used to make the record is not compatible with the electronic system of a data source entity, the pharmacist must transmit the record electronically to the Chief Executive so that it is received no later than—

(a) if the drug is dispensed on a day falling within the first 14 days of a month—the 21st day of that month; or

(b) if the drug is dispensed on any other day—the 7th day of the month following the month in which the drug was dispensed; or

(c) in any case, such later day as the Chief Executive may, on application, authorise.

Maximum penalty: $5 000.

Expiation fee: $1 250.

(3) A pharmacist or medical practitioner who dispenses a drug of dependence on prescription must—

(a) in the case of a written prescription—

(i) retain the original prescription or a copy of the prescription for at least 2 years; and

(ii) keep it readily available for inspection by an authorised officer during that period; and

(iii) on request by an authorised officer—send a copy of the prescription to the authorised officer; or

(b) in the case of a prescription given by fax—

(i) retain the faxed copy of the prescription for at least 2 years; and

(ii) keep the faxed copy of the prescription readily available for inspection by an authorised officer during that period; and

(iii) on request by an authorised officer—send a copy of the faxed copy of the prescription to the authorised officer; or

(c) in the case of an electronic prescription—

(i) retain the electronic prescription or a computer-generated printed copy of it for at least 2 years; and

(ii) keep the electronic prescription or a computer-generated printed copy of it readily available for inspection by an authorised officer during that period; and

(iii) on request by an authorised officer—send a computer-generated printed copy of the electronic prescription to the authorised officer.

Maximum penalty: $5 000.

Expiation fee: $1 250.

(4) A pharmacist or medical practitioner must not—

(a) dispense a monitored drug unless the pharmacist or practitioner has taken all reasonable steps to check relevant information held in the monitored drugs database relating to the person for whom the drug is to be dispensed; or

(b) dispense more than 2 days supply of a drug of dependence unless at least 1 of the following applies:

(i) the person for whose use the drug is prescribed is known to the pharmacist or practitioner;

(ii) the pharmacist or practitioner recognises the signature on the prescription as that of the prescriber who purportedly gave the prescription;

(iii) the pharmacist or practitioner has verified with the prescriber who purportedly gave the prescription that the prescription was in fact given by that prescriber; or

(c) hand over a drug of dependence dispensed by the pharmacist or medical practitioner until—

(i) the person for whose use the drug is dispensed—

(A) has signed and dated the prescription or, if the prescription was given by fax, the faxed copy of the prescription; and

(B) has, unless the person is known to the pharmacist or practitioner, produced satisfactory evidence of their identity; or

(ii) the person for whose use the drug is dispensed—

(A) has signed a computer-generated printed copy of the prescription that includes all the information required to be provided on a written prescription; and

(B) has, unless known to the pharmacist or practitioner, produced satisfactory evidence of their identity; or

(iii) an agent acting on behalf of the person for whose use the drug is intended—

(A) has signed and dated the prescription or, if the prescription was given by fax, the faxed copy of the prescription; and

(B) has, unless the agent is known to the pharmacist or practitioner, produced satisfactory evidence of their identity; or

(iv) an agent acting on behalf of the person for whose use the drug is intended—

(A) has signed a computer-generated printed copy of the prescription that includes all the information required to be provided on a written prescription; and

(B) has, unless known to the pharmacist or practitioner, produced satisfactory evidence of their identity.

Maximum penalty: $5 000.

(5) This regulation (other than subregulation (3)) does not apply to a pharmacist or medical practitioner who dispenses a monitored drug on prescription if—

(a) the prescription is a medication chart prescription; and

(b) the provisions of the Commonwealth Regulations applying to the sale or supply of a pharmaceutical benefit have been complied with in relation to the sale or supply of the drug (whether or not the drug is a pharmaceutical benefit).

(6) Subregulation (4)(a) does not apply if—

(a) the drug (not being dextromoramide or pethidine) is for use by a person aged 70 years or more; or

(b) the prescription for the drug (not being dextromoramide or pethidine) is endorsed either "Notified Palliative Care Patient" or "NPCP"; or

(c) the drug is for use by a person who is receiving treatment in a hospital or a correctional institution and the duration of treatment of the person with the drug while the person is in the hospital or correctional institution does not exceed 14 days; or

(d) the drug is for use by a person who is being discharged from a hospital following treatment in the hospital and the duration of treatment of the person with the drug after discharge does not exceed 14 days.

40—Dispensing prescriptions—S4 drugs in serious shortage

(1) If—

(a) a prescription for an S4 drug is presented to a pharmacist for dispensing; and

(b) the drug to which the prescription relates is a medicine in respect of which a Serious Shortage Medicine Substitution Notice issued by the TGA under the Commonwealth Act is in force; and

(c) the pharmacist is unable to dispense the prescription because the pharmacist does not have, and cannot obtain, the drug to which the prescription relates in the strength, release form or dose form specified in the prescription; and

(d) the person for whom the prescription has been given consents to receiving the drug in a strength, release form or dose form specified in the Notice instead; and

(e) the pharmacist is of the opinion that it is appropriate to supply the person with the drug in a strength, release form or dose form specified in the Notice,

the pharmacist may supply the person with the drug in a strength, release form or dose form specified in the Notice in accordance with the conditions stated in the Notice.

(2) If a pharmacist supplies a drug as authorised by subregulation (1), the pharmacist must, as soon as practicable, give the prescriber of the drug notice in writing of the strength, release form and dose form in which the drug was supplied.

Maximum penalty: $3 000.

(3) For the purposes of these regulations, if a pharmacist supplies a drug as authorised by subregulation (1), the pharmacist will be taken to have dispensed the prescription for the drug presented to the pharmacist.

Part 5—Special provisions relating to monitored drugs

41—Supply or administration of monitored drugs—special provisions

(1) A registered health practitioner acting in the ordinary course of the practitioner's profession must not supply or administer a monitored drug to a person unless the practitioner has taken all reasonable steps to check relevant information held in the monitored drugs database relating to the person.

Maximum penalty: $5 000.

(2) A registered health practitioner acting in the ordinary course of the practitioner's profession must not instruct or otherwise cause another person to supply or administer a monitored drug unless the practitioner has taken all reasonable steps to check relevant information held in the monitored drugs database relating to the person to whom the drug is to be supplied or administered.

Maximum penalty: $5 000.

(3) This regulation does not apply—

(a) where a registered health practitioner (being a prescriber) gives a prescription for a monitored drug; or

(b) where a monitored drug is to be dispensed on prescription; or

(c) where a monitored drug (not being dextromoramide or pethidine) is to be supplied or administered to a person aged 70 years or more; or

(d) where a monitored drug (not being dextromoramide or pethidine) is to be supplied or administered to a person and the registered health practitioner principally responsible for treatment of the person—

(i) reasonably believes the person's life expectancy to be less than 12 months; and

(ii) has informed the Minister of that belief along with the person's name, address and date of birth; or

(e) where a monitored drug is to be supplied or administered to a person who is receiving treatment in a hospital or a correctional institution and the duration of treatment of the person with the drug while the person is in the hospital or correctional institution does not exceed 14 days; or

(f) where a monitored drug is to be supplied or administered to a person who is being discharged from a hospital following treatment in the hospital and the duration of treatment of the person with the drug after discharge does not exceed 14 days; or

(g) to a registered health practitioner supplying or administering a monitored drug on the lawful instruction of another person; or

(h) to a registered health practitioner who, on the lawful instruction of another person, instructs or otherwise causes another registered health practitioner to supply or administer a monitored drug; or

(i) to a person registered under the *Health Practitioner Regulation National Law* to practice in the paramedicine profession as a paramedic who is acting in the ordinary course of that profession.

(4) A reference in this regulation to a ***registered health practitioner*** will be taken to include a person, or person of a class, specified by the Minister by notice in the Gazette for the purposes of this subregulation.

Part 6—Special provisions relating to drugs of dependence

42—Interpretation

(1) In this Part, unless the contrary intention appears—

health service pharmacy means a pharmacy that is part of a health service facility;

order means an order other than a prescription;

supplier means—

(a) a pharmacist; or

(b) a person licensed under the Act to manufacture, sell by wholesale or supply drugs of dependence;

ward of a health service facility means a ward, clinic, unit, operating theatre or any other section of a health service facility in which persons receive medical or dental treatment.

(2) For the purposes of this Part—

(a) a reference to the administration of a drug is, if the drug is administered continuously over an extended period (for example, by means of an intravenous drip or pump) a reference to the commencement of administration by that means; and

(b) the registered health practitioner ***principally responsible*** for the treatment of a person is the practitioner having, for the time being, the greatest input in the determination of the course of treatment of the person.

43—Special restrictions on prescription or supply of drugs of dependence by registered health practitioners and veterinary surgeons

(1) A person must not prescribe or supply a drug of dependence for use by their spouse, domestic partner, parent, grandparent, child, grandchild, brother or sister unless—

(a) the prescription or supply is authorised by the Minister; or

(b) the prescription or supply is in circumstances of a verifiable emergency.

Maximum penalty: $5 000.

Expiation fee: $1 250.

(2) A registered health practitioner must not prescribe or supply a drug of dependence for use by themselves unless the prescription or supply is in circumstances of a verifiable emergency.

Maximum penalty: $5 000.

Expiation fee: $1 250.

(3) Subregulation (1) does not apply to the supply of a drug of dependence by a pharmacist if the pharmacist is dispensing a prescription for the drug.

(4) A veterinary surgeon must not prescribe, sell or supply a drug of dependence for an animal without having first examined the animal unless the prescription, sale or supply is in circumstances of a verifiable emergency.

Maximum penalty: $5 000.

44—Restriction on prescribing or supplying S2, S3 or S4 poisons containing S8 poisons

A prescriber must not prescribe or supply for use by a person who the prescriber knows or has reasonable cause to believe is dependent on drugs—

(a) an S2 poison or S3 poison that contains a poison listed in Schedule 8 of the Poisons Standard; or

(b) an S4 poison that contains a poison listed in Schedule 8 of the Poisons Standard,

for the purpose of maintaining or treating the person's dependence unless the prescriber prescribes or supplies the drug in accordance with an authority granted by the Minister.

Maximum penalty: $5 000.

45—Records to be kept by manufacturers of drugs of dependence

A person who manufactures a drug of dependence must—

(a) record the following details immediately after the drug is manufactured:

(i) the date of manufacture;

(ii) the trade name or the approved name of the drug or, if it does not have either a trade or approved name, its ingredients;

(iii) the amount and, if applicable, the strength of the drug manufactured;

(iv) the total amount of the drug now on the premises on which the drug was manufactured; and

(b) sign and date the record immediately after the record is made.

Maximum penalty: $5 000.

46—Records to be kept by sellers and suppliers of drugs of dependence

(1) A supplier who sells or supplies a drug of dependence must comply with the following provisions:

(a) the supplier must, immediately after selling or supplying the drug—

(i) make a record in electronic form of—

(A) their name and business address; and

(B) the name and address of the person to whom the drug was sold or supplied; and

(C) the date on which the drug was sold or supplied; and

(D) the trade or approved name of the drug or, if the drug does not have either a trade or approved name, the ingredients in the drug; and

(E) the amount and, if applicable, the strength of the drug; and

(F) if the drug was sold or supplied on order—the invoice number (if any) for the sale or supply of the drug;

(ii) make a record of the total amount of the drug now in stock on the premises from which the drug was sold or supplied and sign the record;

(b) if the drug is sold or supplied in accordance with an order, the supplier must, as soon as practicable after selling or supplying the drug, cancel the order by writing "CANCELLED" on the order or, if the order was given by fax endorsed with the name and address of a single pharmacy that may sell or supply the drug, on the faxed copy of the order;

(c) the supplier must transmit the record referred to in paragraph (a)(i) electronically to the Chief Executive so that it is received no later than the 7th day of the month following the month in which the drug was sold or supplied or such later date as the Chief Executive may, on application by the supplier, authorise.

Maximum penalty: $5 000.

(2) A supplier who sells or supplies a drug of dependence on an order must—

(a) retain the original order or a copy of the order for a period of at least 2 years; and

(b) keep it readily available for inspection by an authorised officer; and

(c) on request by an authorised officer—send a copy of the order to the authorised officer.

(3) Subregulation (1)(c) does not apply to—

(a) persons licensed under the Act to manufacture drugs of dependence or sell drugs of dependence by wholesale; or

(b) pharmacies (including health service pharmacies) in respect of the supply of drugs of dependence to a health service facility.

(4) A person who makes a record under subregulation (1) must ensure that the record is kept at all times on the premises from which the drug was supplied.

Maximum penalty: $5 000.

(5) A supplier must not supply a drug of dependence in accordance with an order—

(a) unless the supplier has reasonable cause to believe that the person who ordered the drug is lawfully authorised to do so; and

(b) unless the person receiving the drug—

(i) provides the supplier with a signed and dated receipt for the drug; and

(ii) is known to the supplier or produces satisfactory evidence of their identity.

Maximum penalty: $5 000.

(6) If a drug of dependence is authorised or required by the law of any place to be carried aboard a ship, a person must not supply that drug for carriage aboard a ship unless they have received a written order for the drug from the master or medical officer of the ship.

Maximum penalty: $5 000.

(7) The Minister may exempt a supplier, or a class of suppliers, from this regulation, or specified provisions of this regulation, if satisfied that the supplier, or class of suppliers, has adequate arrangements for the keeping of records.

47—Records to be kept by suppliers of drugs of dependence who receive such drugs

(1) If a supplier of drugs of dependence receives such a drug, or a person receives a drug of dependence from a supplier on order, the person receiving the drug must—

(a) give to the person who provided the drug a signed and dated receipt for the drug; and

(b) record the following details and sign the record:

(i) the name and address of the person who provided the drug;

(ii) the name and address of the person who took delivery of the drug;

(iii) the date on which the drug was received;

(iv) the trade or approved name of the drug or, if the drug does not have either a trade or approved name, the ingredients in the drug;

(v) the amount and, if applicable, the strength of the drug;

(vi) if the drug was provided on order—the invoice number (if any) for the supply of the drug;

(vii) the total amount of the drug now in stock on the premises at which the drug was received.

Maximum penalty: $5 000.

(2) A person who makes a record under this regulation must ensure that the record is kept at all times on the premises at which the drug was received.

Maximum penalty: $5 000.

(3) The Minister may exempt a person, or class of persons, from this regulation, or specified provisions of this regulation, if satisfied that the person, or class of persons, has adequate arrangements for the keeping of records and the security of drugs of dependence.

48—Supply or administration of drugs of dependence by registered health practitioner

(1) A registered health practitioner who supplies a drug of dependence for use by a person, or who administers a drug of dependence to a person, must, immediately after the drug is so supplied or administered, record the following details and sign the record:

(a) their name;

(b) the full name and address (or, in the case of a patient in a ward of a health service facility, the location of the ward) of the person to whom the drug is supplied or administered;

(c) in the case of the supply of the drug to a person acting on behalf of the person for whose use the drug is intended, the full name and address of the person for whose use the drug is intended;

(d) the trade name or approved name of the drug or, if it does not have either a trade or approved name, its ingredients;

(e) the amount and, if applicable, the strength of the drug supplied or administered;

(f) the date;

(g) the time at which the drug was supplied or administered;

(h) the amount of the drug (if any) now remaining—

(i) in stock on the premises at which the drug is supplied or administered; or

(ii) otherwise in the possession of the practitioner.

Maximum penalty: $5 000.

(2) Subregulation (1) does not apply to a pharmacist.

(3) If an error is discovered in a record made for the purposes of subregulation (1), the person authorised to make the record must correct it in the following way:

(a) it must not be deleted, whited out with correction fluid or erased;

(b) it must be ruled out or otherwise marked so as to still be clearly legible after it has been so ruled out or marked;

(c) a footnote or margin note reference must be made alongside the error;

(d) the footnote or margin note must—

(i) be made on the same page as the page on which the error occurs;

(ii) contain the correct information and the date of the correction;

(iii) be endorsed with the name and signature of the person making the correction.

Maximum penalty: $5 000.

(4) The Minister may exempt a registered health practitioner, or class of registered health practitioners, from this regulation, or specified provisions of this regulation, if satisfied that the registered health practitioner, or class of registered health practitioners, has adequate arrangements for the keeping of records.

49—Sale, supply or administration of drugs of dependence by veterinary surgeon

A veterinary surgeon who sells or supplies a drug of dependence for an animal or administers such a drug to an animal must, on the day on which the drug is so sold, supplied or administered, record the following details and sign the record:

(a) their name;

(b) the species of animal for which the drug is sold, supplied or administered, the name and address of the owner of the animal and the name (if any) of the animal;

(c) the trade name or approved name of the drug or, if it does not have either a trade or approved name, its ingredients;

(d) the amount and, if applicable, the strength of the drug sold, supplied or administered;

(e) the date;

(f) the time at which the drug was sold, supplied or administered;

(g) the amount of the drug (if any) now remaining—

(i) in stock on the premises at which the drug is sold, supplied or administered; or

(ii) otherwise in the possession of the veterinary surgeon.

Maximum penalty: $5 000.

50—Additional requirements for administration of drugs of dependence in health service facility

(1) The administration of a drug of dependence to a person in a health service facility must be carried out in accordance with the following additional provisions:

(a) the registered health practitioner principally responsible for the treatment of the person while in the health service facility, or a registered nurse or a midwife acting in accordance with a standing order prepared or endorsed by the health service facility and approved by the Minister must—

(i) ensure that the prescribed instructions in respect of the drug are entered in the person's medication record; and

(ii) endorse the relevant entries with their name and signature and the date of the making of the entries;

(b) the drug must be administered to the person by a registered health practitioner in accordance with all instructions in the person's medication record;

(c) the drug must not be administered to the person unless the administration is witnessed by a registered health practitioner, or, if a registered health practitioner is not reasonably available, by some other responsible person;

(d) the registered health practitioner who administers the drug must, immediately after doing so, ensure that the name and signature of the person who witnessed the administration of the drug is recorded;

(e) if a registered health practitioner gives prescribed instructions by telephone as to the administration of a drug of dependence to a person in a health service facility—

(i) the practitioner must give the instructions to—

(A) a registered health practitioner who is authorised to administer drugs of dependence; and

(B) another responsible person employed at the health service facility; and

(ii) the practitioner to whom the instructions are given must, immediately after receiving the instructions by that method, ensure that the following information is recorded in the person's medication record and sign the record:

(A) their full name;

(B) the prescribed instructions in respect of the drug;

(C) the words "by telephone";

(D) the date on which the telephone instructions were given;

(E) the name of the registered health practitioner who gave the telephone instructions;

(F) the name and signature of the other person to whom the instructions were given in accordance with subparagraph (i); and

(iii) the practitioner who gave the instructions must, within 48 hours of giving the instructions by that method, endorse the relevant entries in the medication record with their signature and the date.

Maximum penalty: $5 000.

(2) The designated nurse or designated midwife for a ward of a health service facility for a particular shift must ensure that the following additional record-keeping requirements are met in respect of drugs of dependence in the ward:

(a) all relevant records required to be kept under these regulations in respect of those drugs must be kept in the ward;

(b) all drugs of dependence must be counted at the end of the shift and—

(i) if the balance in respect of a particular drug is found to be correct, the word "correct", the time and date and the nurse's or midwife's name and signature must be recorded alongside the entry for the drug; and

(ii) if the balance in respect of a particular drug is found to be incorrect—

(A) the word "incorrect", a brief explanation of the discrepancy, if known, the time and date and the nurse's or midwife's name and signature must be recorded alongside the entry for the drug; and

(B) the Director of Nursing or manager of the health service facility, and the health service facility pharmacist, if any, must be notified, as soon as practicable, that an incorrect amount of drugs is stored in the ward;

(c) the drugs count and records made under paragraph (b)—

(i) must be witnessed by the designated nurse or designated midwife for the ward for the next shift and endorsed with their name and signature; or

(ii) must, if the next shift does not commence immediately after the previous shift—

(A) be witnessed by a nurse or midwife working on the same shift as the nurse or midwife who made the entry and be endorsed with the name and signature of the witnessing nurse or midwife; and

(B) be checked by the designated nurse or designated midwife for the ward for the next shift at the commencement of that shift and be endorsed with their name and signature.

Maximum penalty: $5 000.

(3) The Director of Nursing or, if there is no Director of Nursing, the manager of a health service facility must ensure that for each shift for each ward of the health service facility a nurse or midwife is designated as having responsibility for record keeping under subregulation (2).

Maximum penalty: $5 000.

(4) The nurse or midwife designated under subregulation (3) must be a nurse or midwife present on the ward during the shift and may only be an enrolled nurse if no registered nurse or midwife will be present.

(5) The manager of a health service facility must take all reasonable steps to ensure that—

(a) all drugs of dependence delivered to the health service facility or a ward of the health service facility are received by a registered health practitioner employed at the health service facility or, if such a practitioner is not reasonably available, by some other responsible person; and

(b) an accurate and up-to-date balance of stocks of all drugs of dependence in each ward of the health service facility is maintained at all times; and

(c) the requirements of this regulation are complied with.

Maximum penalty: $5 000.

(6) The Minister may exempt a health service facility, or class of health service facilities, from this regulation, or specified provisions of this regulation, if satisfied that the health service facility, or class of health service facilities, has adequate arrangements for the administration of drugs of dependence.

(7) In this regulation—

designated midwife for a ward of a health service facility for a shift means a midwife designated under subregulation (3) as having responsibility for record keeping under subregulation (2) for the ward for the shift;

designated nurse for a ward of a health service facility for a shift means a nurse designated under subregulation (3) as having responsibility for record keeping under subregulation (2) for the ward for the shift;

health service pharmacist means the pharmacist in charge of a health service pharmacy;

prescribed instructions, in respect of a drug, means the form and strength of the drug and the route, frequency and duration of administration of the drug.

51—Special provisions relating to the supply and administration of certain drugs of dependence

(1) A drug of dependence to which this regulation applies may only be supplied or administered to a person for whom it has been prescribed—

(a) by the medical practitioner who prescribed the drug of dependence for the person; and

(b) at a prescribed health service facility; and

(c) in accordance with an approved treatment protocol.

(2) A person who supplies or administers a drug in contravention of subregulation (1) commits an offence.

Maximum penalty: $5 000.

(3) This regulation applies in relation to the following drugs of dependence:

(a) N,α-dimethyl-3,4-(methylenedioxy)phenylethylamine (MDMA);

(b) Psilocybine (Psilocybin).

(4) In this regulation—

approved treatment protocol means a treatment protocol which has been approved by an ethics committee;

ethics committee has the same meaning as in the Commonwealth Act;

prescribed health service facility means—

(a) an approved treatment centre or authorised community mental health facility (both within the meaning of the *Mental Health Act 2009*); or

(b) any other health service facility, or health service facility of a class, determined by the Minister by notice in the Gazette to be a prescribed health service facility for the purposes of this regulation.

52—Destruction of drugs of dependence

(1) Subject to this regulation or any order of a court, a person must not destroy a drug of dependence unless—

(a) the destruction is witnessed by another person, being—

(i) an authorised officer; or

(ii) a police officer; or

(iii) a registered health practitioner; or

(iv) a veterinary surgeon; or

(v) a person who has been authorised in writing by the Chief Executive of the SA Ambulance Service to administer drugs of dependence; and

(b) the person destroying the drug ensures that the following information is recorded in respect of the drug immediately after its destruction:

(i) the full names and the signatures of the person and the witness to the destruction;

(ii) the trade name or approved name of the drug or, if it did not have either a trade or approved name, its ingredients;

(iii) the amount and, if applicable, the strength of the drug;

(iv) the date and time of the destruction;

(v) the amount of the drug (if any) now remaining in stock on the premises at which the destroyed drug was stored.

Maximum penalty: $5 000.

(2) This regulation does not apply to the destruction of a drug of dependence by—

(a) a person for whose use the drug was lawfully prescribed or supplied; or

(b) a police officer or an authorised officer.

Part 7—Special provisions relating to certain paints and tinters

53—Restrictions on manufacture, sale, supply and use of certain paints and tinters

(1) A person must not manufacture, sell, supply or use—

(a) a First Group Paint for application to—

(i) a roof or any surface to be used for the collection or storage of potable water; or

(ii) furniture; or

(iii) any fence, wall, post, gate or building (interior or exterior) other than a building which is used exclusively for industrial purposes or mining or any oil terminal; or

(iv) any premises used for the manufacture, processing, preparation, packing or serving of products intended for human or animal consumption; or

(b) an anti-fouling paint containing more than 0.1% Lead; or

(c) a paint (other than an anti-fouling paint) or tinter containing more than 0.009% Lead; or

(d) a paint for application to toys unless the paint complies with the specification for coating materials contained in AS/NZS ISO 8124.3:2012 *Safety of toys Part 3: Migration of certain elements (ISO 8124-03:2010, MOD)*, as in force from time to time; or

(e) a paint or tinter that contains a pesticide (other than a fungicide, algaecide, bactericide or antifouling agent).

Maximum penalty: $5 000.

(2) Subregulation (1) applies only in relation to a paint or tinter that contains a poison.

(3) For the purposes of this regulation, the proportion of Lead contained in a paint is calculated as a percentage of the element present in the non-volatile content of the paint.

(4) In this regulation, ***First Group Paint***, ***paint***, ***pesticide***, ***tinter*** and ***toy*** have the same respective meanings as in the Poisons Standard.

Part 8—Other offences

54—Prohibition on giving samples of S8 poisons

A person must not give another person a sample of an S8 poison.

Maximum penalty: $5 000.

55—Offences relating to sale or supply of poisons

(1) A person must not, in any residential premises, or from door-to-door, or in a public place, sell or supply—

(a) an S2 poison, S3 poison, S4 poison, S7 poison or S8 poison; or

(b) an S5 poison or S6 poison that is not a product sample.

Maximum penalty: $5 000.

(2) A person must not, in any residential premises, or from door-to-door, or in a public place, sell or supply an S5 poison or S6 poison that is a product sample except as permitted by Part 2 Section 61 of the Poisons Standard.

Maximum penalty: $5 000.

(3) For the purposes of subregulations (1) and (2), a poison is a product sample if—

(a) it is supplied directly to the consumer free of charge, or at a nominal charge, as a mechanism to promote the sale of the product; and

(b) it is supplied in a small pack produced specifically for the purposes of promotion, or packaged in a normal commercial pack that in other circumstances a consumer would need to purchase.

(4) A person must not sell or supply an S2 poison, S3 poison, S4 poison or S8 poison in a container that—

(a) is normally used for containing food or beverages; or

(b) is similar to a container that is normally used for containing food or beverages.

Maximum penalty: $5 000.

(5) A person must not sell any liquid preparation or admixture containing paraquat unless it is coloured blue or green and contains a stenching agent in sufficient quantity to produce an offensive odour.

Maximum penalty: $5 000.

(6) In this regulation—

public place includes—

(a) a place to which free access is permitted to the public, with the express or tacit consent of the owner or occupier of that place; and

(b) a place to which the public are admitted on payment of money, the test of admittance being the payment of money only; and

(c) a road, street, footway, court, alley or thoroughfare that the public are allowed to use, notwithstanding that the road, street, footway, court, alley or thoroughfare is on private property.

56—Offence relating to prescribing or supplying vaping substance to children

(1) A registered health practitioner must not prescribe or supply a vaping substance to a person under the age of 18 years.

Maximum penalty: $10 000.

(2) In this regulation—

vaping substance means nicotine (being an S4 poison) in solution in any concentration (including in a salt or base form).

Note—

See *Tobacco and E-Cigarette Products Act 1997* for further offences relating to the sale and supply of other e-cigarette products.

57—Offence to dispose of poison

A person must not—

(a) dispose of or use, or cause to be disposed of or used, an S5 poison, S6 poison or S7 poison except in accordance with the requirements of Part 2 Section 55 of the Poisons Standard; or

(b) dispose of or use, or cause to be disposed of or used, an S2 poison, S3 poison, S4 poison or S8 poison in any manner that constitutes, or is likely to constitute, a risk to public health or safety.

Maximum penalty: $5 000.

58—Keeping of records etc

(1) Subject to these regulations, a person who is required by these regulations to keep certain records must—

(a) in respect of any entry in the records, retain the records at the registered address of the business in this State for a period of—

(i) in the case of records relating to S7 poisons—for a period of 5 years; or

(ii) in any other case—for a period of 2 years,

from the day on which the entry was made; and

(b) have the records readily available for inspection at all reasonable times; and

(c) during that period, take all reasonable steps to ensure that the records are protected against deterioration, loss, theft and unauthorised access, modification or use.

Maximum penalty: $3 000.

(2) If the information contained in the records is available only after the record is subjected to an electronic or other process, it is sufficient for the purposes of subregulation (1)(b) for the person to produce for inspection a reproduction or computerised record of any entry in the records.

(3) Details that are required to be recorded under these regulations in respect of drugs of dependence must, unless otherwise specified, be recorded in a register of drugs of dependence (and any electronic register of drugs of dependence must be in a form approved by the Minister).

(4) A receipt required to be provided to a person under these regulations must be kept by that person in the manner set out in this regulation as if it were a record.

59—Vicarious liability

For the purposes of these regulations, an act or omission of an employee or agent will be taken to be the act or omission of the employer or principal unless it is proved that the act or omission did not occur in the course of the employment or agency.

Part 9—Miscellaneous

60—Personal identification code equivalent to signature

(1) If a provision of these regulations requires a person to sign a record or receipt that is in electronic form, evidence on the record or receipt that the person has entered their personal identification code will be taken to be sufficient compliance by that person with the requirement.

(2) To avoid doubt, nothing in subregulation (1) derogates from the application of the *Electronic Communications Act 2000* in relation to the manner in which a requirement to sign a record or receipt that is in electronic form may be met.

(3) In this regulation—

personal identification code means a code that—

(a) is allotted to a person by their employer for use by that person in connection with official duties; and

(b) is known only by that person and such other persons as may be authorised by the employer for management purposes.

61—Permits (section 56(1) of Act)

An application for a permit under section 56(1) of the Act must be made in writing to the Minister and signed by the applicant.

62—Prescribed professional associations (section 58(1a) of Act)

For the purposes of section 58(1a) of the Act, the following professional associations are prescribed:

(a) in the case of publishing information to medical practitioners—

(i) the Australian Medical Association; and

(ii) the Royal Australian College of General Practitioners;

(b) in the case of publishing information to pharmacists—

(i) the Friendly Society Medical Association; and

(ii) the Pharmaceutical Society of Australia; and

(iii) the Pharmacy Guild of Australia; and

(iv) the Society of Hospital Pharmacists of Australia.

63—Disclosure of confidential information contained in monitored drugs database (section 60A(1)(e) of Act)

(1) Information contained in the monitored drugs database relating to a particular person may be disclosed to a prescriber involved in the medical treatment or care of that person to enable that prescriber to access that information and disclose that information to—

(a) any registered health practitioner involved in the medical treatment or care of that person; and

(b) any pharmacist to whom a prescription for a monitored drug for that person has been presented.

(2) Information contained in the monitored drugs database relating to a particular person may be disclosed to a pharmacist to whom a prescription for a monitored drug for that person has been presented to enable that pharmacist to access that information and disclose that information to—

(a) any other pharmacist to whom a prescription for a monitored drug for that person has been presented; and

(b) any registered health practitioner involved in the medical treatment or care of that person.

(3) Information contained in the monitored drugs database may be disclosed to a health authority of an Australian jurisdiction responsible for the administration or enforcement of a law that regulates the sale, supply, prescription, administration and use of monitored drugs.

(4) Information contained in the monitored drugs database may be disclosed in accordance with an authorisation given by the Minister.

64—Corresponding laws (section 61(4) of Act)

For the purposes of the definition of ***corresponding law*** in section 61(4) of the Act, the following laws are prescribed:

(a) the *Drugs of Dependence Act 1989* of the Australian Capital Territory;

(b) the *Drugs Misuse and Trafficking Act 1985* of New South Wales;

(c) the *Misuse of Drugs Act 1990* of the Northern Territory;

(d) the *Drugs Misuse Act 1986* of Queensland;

(e) the *Medicines and Poisons Act 2019* of Queensland;

(f) the *Poisons Act 1971* of Tasmania;

(g) the *Drugs, Poisons and Controlled Substances Act 1981* of Victoria;

(h) the *Misuse of Drugs Act 1981* of Western Australia;

(i) the *Medicines and Poisons Act 2014* of Western Australia.

65—Place at which codes, standards and other documents must be kept for public inspection etc (section 63(5a)(a) of Act)

For the purposes of section 63(5a)(a) of the Act, the office of the Department at 11-13 Hindmarsh Square, Adelaide is prescribed.

66—Approvals, determinations and exemptions

(1) The Minister may, at any time, by notice in writing—

(a) impose such conditions as the Minister thinks fit on an approval or exemption granted by the Minister, or on a determination made by the Minister, under these regulations; or

(b) vary or revoke the conditions of such an approval, determination or exemption as the Minister thinks fit; or

(c) revoke, as the Minister thinks fit, an approval or exemption granted by the Minister, or a determination made by the Minister, under these regulations.

(2) A person must not contravene or fail to comply with a condition of an approval or exemption granted by the Minister, or a determination made by the Minister, under these regulations.

Maximum penalty: $3 000.

Schedule 1—Forms

End user statement

The chemical product I wish to purchase is classified as a possible illicit drug precursor or auxiliary reagent. I understand that to be supplied this product a signed end user declaration must be provided together with an order.

Catalogue No	Product Name	Quantity	Pack Size	Order No

Intended use: ☐Analytical ☐Research and Design ☐Manufacturing ☐Resale ☐Other

Please specify full details of assay, project, product customer etc

Purchaser details and declaration

I, *[insert full name]* being *[insert position]* on behalf of *[insert name of company or institution and ACN]*
Address:
Account No:
declare that the above chemical product will not be used for the manufacture of illicit drugs.

Signature:
Date:

Details of collecting agent's identification

Current Passport No:
Country of Issue:
Current Photograph Licence No:
Expiry date:
Photo Identification Card Type:

End user distributor/supplier details and declaration

I, *[insert full name]* being *[insert position]* on behalf of *[insert name of company or institution and ACN]*
Address:
Account No:
declare that the above chemical product will not be used for the manufacture of illicit drugs.

Signature:
Date:

Note—

1 Please attach a photocopy of current driver's licence bearing a photograph.

2 The form must be completed with all details.

Schedule 2—Repeal and transitional provisions

Part 1—Repeal of *Controlled Substances (Poisons) Regulations 2011*

1—Repeal of *Controlled Substances (Poisons) Regulations 2011*

The *Controlled Substances (Poisons) Regulations 2011* are repealed.

Part 2—Transitional provisions

2—Preliminary

In this Part—

repealed regulations means the *Controlled Substances (Poisons) Regulations 2011*.

3—Approvals, exemptions, authorisations and determinations

An approval, exemption, authorisation or determination made or given under or for the purposes of a provision of the repealed regulations in force immediately before the commencement of these regulations will, on that commencement, be taken to be an approval, exemption, authorisation or determination (as the case requires) made or given under or for the purposes of the corresponding provision of these regulations subject to such conditions (if any) as applied under the repealed regulations.

4—Requirements relating to administration of certain S4 drugs

A requirement that was, immediately before the commencement of these regulations, a requirement specified by the Minister under regulation 18(3)(d)(ii)(B) or (3a)(b)(ii) of the repealed regulations will, on that commencement, be taken to be a requirement specified by the Minister under regulation 19(3)(d)(ii)(B) or (4)(b)(ii) (as the case requires) of these regulations.

5—Requirements relating to giving prescription by approved electronic communication

A requirement that was, immediately before the commencement of these regulations, a requirement imposed by the Minister under regulation 33(10) of the repealed regulations will, on that commencement, be taken to be a requirement imposed by the Minister under regulation 35(12) of these regulations.

6—Specification of person or class of person

A person, or person of a class, that was, immediately before the commencement of these regulations, a person, or person of a class, specified by the Minister for the purposes of regulation 35C(4) of the repealed regulations will, on that commencement, be taken to be a person, or person of a class, specified by the Minister for the purposes of regulation 41(4) of these regulations (subject to any further Gazette notice made by the Minister for the purposes of that regulation).

Editorial note—

As required by section 10AA(2) of the *Legislative Instruments Act 1978*, the Minister has certified that, in the Minister's opinion, it is necessary or appropriate that these regulations come into operation as set out in these regulations.

Made by the Governor

on the recommendation of the Controlled Substances Advisory Council and with the advice and consent of the Executive Council
on 7 May 2026

No 15 of 2026

STATE GOVERNMENT INSTRUMENTS

EMPLOYMENT AGENTS REGISTRATION ACT 1993

Exemption

Notice is hereby given that, pursuant to Section 4(1) of the *Employment Agents Registration Act 1993*, I, Kyam Maher MLC, Deputy Premier, Minister for Industrial Relations, hereby exempt Connect Staffing Group East Pty Ltd of Victoria (Suite 5, A11/2A Westall Road, Clayton VIC 3168) from:

- Section 7(3) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement that, in making an application for a licence, the applicant must furnish the name of a natural person, who is a resident of the State and is to act as manager of the business;
- Section 11(1) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement that the business conducted in pursuance of the licence must be managed under the personal supervision of a natural person who is a resident of the State;
- Section 16(1) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement that the holder of a licence must not carry on the business of an employment agent except at premises registered under this Section;
- Section 17 of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement that a person carrying on business as an employment agent in pursuance of a licence must maintain in a conspicuous position at any registered premise a notice clearly displaying the name of the agent as it appears in the licence or a registered business name in which the agent carries on business as an agent and if a manager has been appointed, the name of the manager of the business; and
- Section 19(1) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement to display a notice clearly showing the scale of fees for the time being chargeable by the agent in respect of his or her business at the registered premises.
- Section 22(3) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement for records, accounts and documents to be kept at registered premises for the period of one year.

Dated: 3 May 2026

KYAM MAHER MLC
Deputy Premier
Minister for Industrial Relations

EMPLOYMENT AGENTS REGISTRATION ACT 1993

Exemption

Notice is hereby given that, pursuant to Section 4(1) of the *Employment Agents Registration Act 1993*, I, Kyam Maher MLC, Deputy Premier, Minister for Industrial Relations, hereby exempt Connect Staffing Group of Western Australia (Level 5, Suite 25, 92 Walters Drive, Osborne Park WA 6017) from:

- Section 7(3) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement that, in making an application for a licence, the applicant must furnish the name of a natural person, who is a resident of the State and is to act as manager of the business;
- Section 11(1) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement that the business conducted in pursuance of the licence must be managed under the personal supervision of a natural person who is a resident of the State;
- Section 16(1) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement that the holder of a licence must not carry on the business of an employment agent except at premises registered under this Section;
- Section 17 of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement that a person carrying on business as an employment agent in pursuance of a licence must maintain in a conspicuous position at any registered premise a notice clearly displaying the name of the agent as it appears in the licence or a registered business name in which the agent carries on business as an agent and if a manager has been appointed, the name of the manager of the business; and
- Section 19(1) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement to display a notice clearly showing the scale of fees for the time being chargeable by the agent in respect of his or her business at the registered premises.
- Section 22(3) of the *Employment Agents Registration Act 1993* (SA) in relation to the requirement for records, accounts and documents to be kept at registered premises for the period of one year.

Dated: 3 May 2026

KYAM MAHER MLC
Deputy Premier
Minister for Industrial Relations

HOUSING IMPROVEMENT ACT 2016

Rent Control

In the exercise of the powers conferred by the *Housing Improvement Act 2016*, the Delegate of the Minister for Housing and Urban Development hereby fixes the maximum rental amount per week that shall be payable subject to Section 55 of the *Residential Tenancies Act 1995*, in respect of each premises described in the following table. The amount shown in the said table shall come into force on the date of this publication in the Gazette.

Address of Premises	Allotment Section	Certificate of Title Volume/Folio	Maximum Rental per week payable
37 Vincent Street, Christies Beach SA 5165	Allotment 390 Deposited Plan 7460 Hundred of Noarlunga	CT5597/45	$442.50

Dated: 7 May 2026

CRAIG THOMPSON
Housing Regulator and Registrar
Housing Safety Authority
Delegate of the Minister for Housing and Urban Development

HOUSING IMPROVEMENT ACT 2016

Rent Control Revocations

In the exercise of the powers conferred by the *Housing Improvement Act 2016*, the Delegate of the Minister for Housing and Urban Development hereby revokes the maximum rental amount per week that shall be payable subject to Section 55 of the *Residential Tenancies Act 1995*, in respect of each premises described in the following table.

Address of Premises	Allotment Section	Certificate of Title Volume/Folio
16 Nari Drive, Sheidow Park SA 5158	Allotment 400 Deposited Plan 10122 Hundred of Noarlunga	CT5531/827
15 Duck Ponds Road, Stockwell SA 5355 (Rear of shop)	Allotment 53 Deposited Plan 135 Hundred of Moorooroo	CT6090/273

Dated: 7 May 2026

CRAIG THOMPSON
Housing Regulator and Registrar
Housing Safety Authority
Delegate of the Minister for Housing and Urban Development

HYDROGEN AND RENEWABLE ENERGY ACT 2023

Application for Grant of Associated Infrastructure Licence—AILA 15

Pursuant to Section 32(3) of the *Hydrogen and Renewable Energy Act 2023*, notice is hereby given that an application for an Associated Infrastructure Licence over area described below has been received from:

Utilacor Pty Ltd

Description of Application Area

All that part of the State of South Australia, bounded as follows:

All coordinates GDA2020, Zone 53

573744.05mE 6159855.67mN
573788.77mE 6159854.95mN
573788.58mE 6159876.30mN
574554.60mE 6159912.90mN
574576.35mE 6159882.26mN
574598.40mE 6159836.46mN
574635.94mE 6159597.53mN
574709.91mE 6159539.78mN
574697.60mE 6159524.02mN
574617.54mE 6159586.53mN
574585.11mE 6159770.12mN
573764.06mE 6159445.85mN
573744.05mE 6159855.67mN

AREA: **0.25** square kilometres approximately

The application may be inspected at the offices of the Department for Energy and Mining located at Level 4, 11 Waymouth Street, Adelaide SA 5000. To arrange an inspection, please contact the Department via email at DEM.ERDLicensing@sa.gov.au.

Dated: 29 April 2026

MICHAEL SMITH
Director, Regulatory Risk and Resource Tenure
Regulation and Compliance Division
Department for Energy and Mining
Delegate of the Minister for Energy and Mining

HYDROGEN AND RENEWABLE ENERGY ACT 2023

Application for Grant of Associated Infrastructure Licence—AILA 29

Pursuant to Section 32(3) of the *Hydrogen and Renewable Energy Act 2023*, notice is hereby given that an application for an associated infrastructure licence over area described below has been received from:

Regional Council of Goyder

Description of Application Area

All that part of the State of South Australia, bounded as follows:

All coordinates GDA2020, Zone 54

306845.83mE 6272881.96mN
306845.87mE 6272882.01mN
306960.06mE 6272784.99mN
306960.03mE 6272784.97mN
306946.53mE 6272769.27mN
306932.57mE 6272739.26mN
306934.85mE 6272727.10mN
306869.60mE 6272687.62mN
306829.38mE 6272704.00mN
306803.83mE 6272705.93mN
306759.89mE 6272713.70mN

306726.07mE 6272702.14mN
306711.16mE 6272691.97mN
306707.74mE 6272691.45mN
306695.73mE 6272688.66mN
306686.47mE 6272688.03mN
306675.79mE 6272688.59mN
306662.41mE 6272689.65mN
306654.04mE 6272690.81mN
306633.88mE 6272696.62mN
306604.94mE 6272714.67mN
306581.49mE 6272750.81mN
306571.64mE 6272765.26mN
306558.28mE 6272783.74mN
306583.35mE 6272805.77mN
306605.49mE 6272853.15mN
306643.07mE 6272866.22mN
306688.77mE 6272849.61mN
306693.91mE 6272847.72mN
306762.61mE 6272845.11mN
306775.39mE 6272846.59mN
306837.53mE 6272865.58mN
306845.83mE 6272881.96mN

AREA: **0.053** square kilometres approximately

The application may be inspected at the offices of the Department for Energy and Mining located at Level 4, 11 Waymouth Street, Adelaide SA 5000. To arrange an inspection, please contact the Department via email at DEM.ERDLicensing@sa.gov.au.

Dated: 30 April 2026

MICHAEL SMITH
Director, Regulatory Risk and Resource Tenure
Regulation and Compliance Division
Department for Energy and Mining
Delegate of the Minister for Energy and Mining

JUSTICES OF THE PEACE ACT 2005

SECTION 4

Notice of Appointment of Justices of the Peace for South Australia by the Commissioner for Consumer Affairs

I, Brett Humphrey, Commissioner for Consumer Affairs, delegate of the Attorney-General, pursuant to Section 4 of the *Justices of the Peace Act 2005*, do hereby appoint the people listed as Justices of the Peace for South Australia as set out below. It being a condition of appointment that the Justices of the Peace must take the oaths required of a justice under the *Oaths Act 1936* and return the oaths of office form to Justice of the Peace Services within three months after the date of appointment.

For a period of ten years for a term commencing on 14 May 2026 and expiring on 13 May 2036:

ZOLAISON
Deonne Krystal WYATT
Hayley Nicole WALMSLEY
Samantha Dharma Sri UDAKUMBURA
Beth Louise SZTEKEL
Fay Fadia SUMMERS-ROUDA
Jodie Corinna STONE
Amol Kumar PUNNI
Beverly POVEY
Diana Maria Costantina PANAZZOLO
Natasha Marie ODEGAARD
Amanda Paige O'MALLEY
Gillian Margaret NORRINGTON
Nicola NESBITT
Leonie Rachel LEIGHTON
Sourav Kumar KHURANA
Isabelle Anh Minh HUBCZENKO
Ronald Martin HORNE
David Andrew HEMER
Amy Caitlin HARVEY
Antony Alan HALL
Darryl DO
Andrew Robert Stanford COIDAN
Kamal Kishor BHAGAT
Roanne Selma BEREKMERI
Stephen David BANKS
Keyvan ABAK

Dated: 30 April 2026

BRETT HUMPHREY
Commissioner for Consumer Affairs
Delegate of the Attorney-General

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 1040 in D138091 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5119 Folio 766, expressly excluding right(s) of way over Allotments 285 and 286 in DP 1761 (T 497912) and expressly excluding the right(s) of way A in D138091 created by T 538592 (being one and the same as allotment 287 in DP 1761 created by T 538592).

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969*;
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969*.

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 4 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT: 2024/07411/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 1058 in D138101 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5119 Folio 946, expressly excluding right(s) of way over Allotments 285 and 286 in DP 1761 (T 497912) and expressly excluding free and unrestricted rights(s) of way A in D138101 (being one and the same as allotment 287 in DP 1761).

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969*;
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969*.

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 4 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT: 2024/07429/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 1052 in D138097 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5118 Folio 173, expressly excluding right(s) of way over Allotments 285 and 286 in DP 1761 (T 497912) and expressly excluding the right(s) of way A in D138097 created by T 506587 (being one and the same as allotment 287 in DP 1761 created by T 506587).

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969*;
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969*.

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 4 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT: 2024/07433/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 1042 in D138092 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5119 Folio 943, expressly excluding right(s) of way over Allotments 285 and 286 in DP 1761 (T 497912) and expressly excluding the right(s) of way A in D138092 created by T 535884 (being one and the same as allotment 287 in DP 1761 created by T 535884).

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969*;
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969*.

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 4 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT: 2024/07434/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 1001 in D138786 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5129 Folio 776.

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969*;
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969*.

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 5 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT 2024/08119/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 1001 in D139081 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5846 Folio 23, expressly excluding the free and unrestricted right(s) of way over the land marked A.

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969*;
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969*.

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 5 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT 2024/08121/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 1001 in D139338 lodged in the Lands Titles Office being:

First, portion of the land comprised in Certificate of Title Volume 5962 Folio 780.

Expressly excluding the:

- Party Wall right(s) over the land marked B and H on D139338 (T2929930)
- Party Wall right(s) over the land marked D on D139338 (T2929931)

Secondly, that portion of the easement over the land marked E on D139338 created by T2230276 comprised in Certificate of Title Volume 5222 Folio 765 appurtenant to Allotment 59 in FP 10391 (except those portions of Allotment 59 that are marked X on the said Certificate of Title), as is contained within and forms portion of the said Allotment 1001, to the intent that the easement will merge with and be extinguished in the fee simple in Allotment 1001.

This notice is given under Section 26F of the *Land Acquisition Act 1969.*

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969;*
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969.*

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 5 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT 2024/08168/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 1011 in D139082 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5676 Folio 400.

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969*;
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969*.

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 5 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT 2024/08169/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An unencumbered estate in fee simple in the whole of Allotment 2342 in D139130 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5881 Folio 38, expressly excluding the:

- Free and unrestricted right(s) of way over the land marked A appurtenant only to the land marked Y
- Free and unrestricted right(s) of way over the land marked C

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969*;
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969*.

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 5 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT: 2024/08200/01

LAND ACQUISITION ACT 1969

SECTION 26F

Form 6B—Notice of Acquisition of Underground Land

1. Notice of acquisition

The Commissioner of Highways (the Authority), of 83 Pirie Street, Adelaide SA 5000 acquires the following interests in the following land:

An estate in fee simple in the whole of Allotment 5021 in D139166 lodged in the Lands Titles Office, being portion of the land comprised in Certificate of Title Volume 5088 Folio 306, subject to service easement(s) over the land marked Z(T/F) for electricity supply purposes to Distribution Lessor Corporation (Subject to lease 8890000) (223LG RPA).

This notice is given under Section 26F of the *Land Acquisition Act 1969*.

2. Compensation not payable unless certain water infrastructure or rights are affected

You are not entitled to compensation in relation to the acquisition of the underground land to which this notice relates, unless the following conditions are satisfied:

- you held at least one of the following interests in relation to the underground land immediately before the notice of acquisition was published in relation to the land—
  - ownership of a lawful well that provides access to underground water in the underground land, and any underground infrastructure associated with the well; or
  - a right to take underground water from the underground land by means of such a well;
- you notified the Authority of your interest in response to a notice given under Section 26G of the *Land Acquisition Act 1969;*
- the acquisition of the underground land either—
  - involved the acquisition of your interest; or
  - resulted in the discharge of your interest; or
  - resulted in you being unable to take water by means of, or pursuant to, your interest;
- you make an application for compensation to the Authority under Section 26H of the *Land Acquisition Act 1969.*

3. Application for compensation under Section 26H

If you believe you are entitled to compensation, you must apply to the Authority for compensation within 6 months after the publication of the notice of acquisition in relation to the underground land to which this notice relates.

The application must be in the following manner and form:

"Application for Compensation for Acquisition of Underground Land" (enclosed) to be submitted by email to DIT.ULAapplications@sa.gov.au or by mail marked attention: Property Acquisition c/- GPO Box 1533, Adelaide SA 5001.

The application must be accompanied by the following information or documents:

Any relevant supporting documentation including, but not limited to water licences, bore licences etc.

After receiving your application, the Authority may (but is not required to) make you a written offer of compensation not exceeding $50,000.

See Section 26H(4) of the *Land Acquisition Act 1969* for further details on the payment of compensation.

4. Inquiries

Inquiries should be directed to: T2D Project Team
GPO Box 1533
Adelaide SA 5001
Telephone: 1800 572 414

Dated: 5 May 2026

The Common Seal of the COMMISSIONER OF HIGHWAYS was hereto affixed by authority of the Commissioner in the presence of:

ROCCO CARUSO
Director, Property Acquisition
(Authorised Officer)
Department for Infrastructure and Transport

DIT 2024/08248/01

South Australia

Legal Practitioners (Fees) Notice 2026

under the *Legal Practitioners Act 1981*

1—Short title

This notice may be cited as the *Legal Practitioners (Fees) Notice 2026*.

Note—

This is a fee notice made in accordance with the *Legislation (Fees) Act 2019*.

2—Commencement

This notice has effect on the day on which it is made.

3—Interpretation

In these Regulations, unless the contrary intention appears—

Act means the *Legal Practitioners Act 1981*.

4—Fees

The Fees specified in Schedule 1 are prescribed for the purposes of the Act.

Schedule 1—Fees

1	For the issue or renewal of a practising certificate (other than a volunteer practising certificate)—	
	(a) for more than 6 months	$770 fee $242 levy
	(b) for 6 months or less	$432 fee $121 levy
2	Fee for the issue or renewal of a volunteer practising certificate (see LPEAC rule 3B: category D practising certificate)	$103 fee $47 levy
3	Fee to accompany written notice provided under Section 23D of the Act	$34.50
4	Fee to accompany written notice provided under Schedule 1 Clauses 4(1) and 5(2) of the Act	$34.50

Made by the Attorney-General

Hon Kyam Maher MLC

On 4 May 2026

MINING ACT 1971

Application for a Mining Lease

Notice is hereby given in accordance with Section 56H of the *Mining Act 1971*, that an application for a Mining Lease over the undermentioned mineral claim has been received:

Applicant:	Adelaide Brighton Cement Limited (ACN 007 870 199)
Claim Number:	4581
Location:	CT5833/574, Wool Bay area—approximately 110km south of Kadina.
Area:	66.67 hectares approximately
Purpose:	Industrial Minerals (Limestone)
Reference:	MLA-01099

To arrange an inspection of the proposal at the Department for Energy and Mining, please call the Department on 08 8463 3103.

An electronic copy of the proposal can be found on the Department for Energy and Mining website:
https://www.energymining.sa.gov.au/industry/minerals-and-mining/mining/community-engagement-opportunities

Written submissions in relation to this application are invited to be received at the Department for Energy and Mining, Mining Regulation, Attn: Business Support Officer, GPO Box 618, Adelaide SA 5001 or dem.miningregrehab@sa.gov.au by no later than **4 June 2026**.

The Minister for Energy and Mining (or delegate) is required to have regard to submissions in determining whether to grant or refuse the application and, if granted, the terms and conditions on which it should be granted.

When you make a written submission, that submission becomes a public record. Your submission will be provided to the applicant and may be made available for public inspection.

Dated: 7 May 2026

C. ANDREWS
Mining Registrar
Delegate for the Minister for Energy and Mining
Department for Energy and Mining

MINING ACT 1971

Application for a Mining Lease

Notice is hereby given in accordance with Section 56H of the *Mining Act 1971*, that an application for a Mining Lease over the undermentioned Retention Lease has been received:

Applicant:	OneSteel Manufacturing Pty Limited operating as SIMEC Mining (Administrators Appointed) (ACN 004 651 325)
Retention Lease Number:	RL 133
Location:	CT6097/620, Ardrossan area, approx. 20km east of Maitland
Area:	38.95 hectares approximately
Purpose:	Industrial Minerals (Dolomite)
Reference:	MLA-01063

To arrange an inspection of the proposal at the Department for Energy and Mining, please call the Department on 08 8463 3103.

An electronic copy of the proposal can be found on the Department for Energy and Mining website:
https://www.energymining.sa.gov.au/industry/minerals-and-mining/mining/community-engagement-opportunities

Written submissions in relation to this application are invited to be received at the Department for Energy and Mining, Mining Regulation, Attn: Business Support Officer, GPO Box 618, Adelaide SA 5001 or dem.miningregrehab@sa.gov.au by no later than **2 June 2026**.

The delegate of the Minister for Energy and Mining is required to have regard to these submissions in determining whether to grant or refuse the application and, if granted, the terms and conditions on which it should be granted.

When you make a written submission, that submission becomes a public record. Your submission will be provided to the applicant and may be made available for public inspection.

Dated: 7 May 2026

G. HAMMOND
Acting Mining Registrar
Delegate for the Minister for Energy and Mining
Department for Energy and Mining

South Australia

Motor Vehicles (Approval of Motor Bikes and Motor Trikes) Notice 2026 No 3

under the *Motor Vehicles Act 1959*

1—Short title

This notice may be cited as the *Motor Vehicles (Approval of Motor Bikes and Motor Trikes) Notice 2026*.

2—Commencement

This notice will come into operation on the date of publication in this Gazette.

3—Approved motor bikes and motor trikes

For the purposes of Schedules 2 and 3 of the *Motor Vehicles Regulations 2025* the motor bikes and motor trikes specified in Schedule 1 are approved.

Schedule 1—Approved motor bikes and motor trikes

The following are approved:

- All motor bikes and motor trikes built before December 1960 with an engine capacity not exceeding 660ml
- All motor bikes and motor trikes with an engine capacity not exceeding 260 millilitres and a power to weight ratio not exceeding 150 kilowatts per tonne, except the following:

  Suzuki RGV250
  Kawasaki KR250 (KR-1 and KR1s models)
  Honda NSR250
  Yamaha TZR250
  Aprilia RS250
- All motor bikes and motor trikes with electric powered engines, with a power output not in excess of 25kw

Motor bikes and Motor trikes with electric powered engines listed in the table below are approved:

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
BRAAAP	MotoE	5000w	2022-current	Electric
	MotoE	8000w	2022-current	Electric
	MotoE	10000w	2022-current	Electric
EVOKE	URBAN S		2020-current	Electric
	URBAN CLASSIC		2020-current	Electric
FONZARELLI	125	125	2014-2015	Electric
KAWASAKI	NX011A	NR011A (Ninja e-1) NX011A (Ninja e-1)	2023-current	Electric
KYBURZ	DXP	KYBURZ	2017-current	Electric
UBCO	2018 2x2	UBCO	2018	Electric
	Duty	Duty	2024-current	Electric

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	2x2 ADV	2x2 ADV	2020-current	Electric
ZERO	DS	Zero DS	Unit 2015	Electric
	S	Zero S	Until 2015	Electric

Motor trikes with an engine capacity not less than 261ml and not exceeding 660ml listed in the table below are approved:

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
BRP	Can am Ryker	Rotax 600 ACE	2018	599
GILERA	FUOCO 500	FUOCO 500	2007-13	493
LAMBRETTA	All model	Lambretta	pre 2008	under 660
OZ TRIKE	FUN 500	FUN 500	pre 2008	500
METROPOLIS	AA	2018	399	METROPOLIS
PGO	All models	All models under 220	All	220
PIAGGIO	All Models	All models	2010-17	under 350

Motor bikes with an engine capacity not less than 261ml and not exceeding 660ml listed in the table below are approved:

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
AJP	PR7	PR7	2016-on	600
AJS	MODEL 18	MODEL 18	pre 1966	497
	MODEL 20	MODEL 20	1955-61	498
APRILIA	KV	RS 660 LAMS & Tuono 660 LAMS	2020-on	659
	Moto 6.5	Moto 6.5	1998-2000	649
	M35	SR MAX 300	2012-2018	278
	PEGASO 650	DUAL SPORTS	1994-01	652
	PEGASO 650	OUTBACK	2000-01	652
	PEGASO 650	Factory 650	2007-08	660
	PEGASO 650 I.E.	OUTBACK	2001-02	652
	PEGASO 650 I.E.	DUAL SPORTS	2001-06	652
	SCARABEO 300	VRG	2009-13	278
	SCARABEO 400	SCARABEO 400	2007	399
	SCARABEO 500	SCARABEO 500	2006-12	460
	SPORTCITY300	SPORTCITY300	2010-13	300
	STRADA 650	ROAD	2006-08	659
	STRADA 650	TRAIL	2006-08	659
	VP (RXV 450)	VPV-VPT-VPH 18.3kW	2006-10	449
	VP (RXV 550)	VPZ- VPX- VPL 20kW	2006-10	553
	VS (SXV 450)	SXV 450 (VSR-VSH) 14kW	2006-09	449
	VS (SXV 550)	SXV 550 (VSS-VSL) 14.5kW	2006-12	553
	XC	Tuareg 660 (LAMS)	2022-on	659
	XE (RS 457)	XEA	2024-on	457
	XH	XHB00, RS 660 Factory LAMS	2025-on	659
	XH	XHE00, Tuono 660 Factory LAMS	2025-on	659
	XH	XHL00, RS 660 LAMS	2025-on	659
	XM	XMA00, Tuareg 660 (LAMS)	2025-on	659
	XM	XMB00, Tuareg 660 Rally (LAMS)	2025-on	659
ASIAWING	LD450	ODES MCF450	2011-13	449
ATK	605	605	1995	598
BENELLI	LEON	Leoncino, Leoncino Trail	2018-on	500
	VELVET DUSK	Velvet 400 Dusk, Velvet Touring 400	2002-06	383
	P10	BN 302	2015-on	300

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	P16	TRK 502X	2018-on	500
	P18	LEONCINO 500	2017-on	500
	P18	LEONCINO 500 TRAIL	2018-on	500
	P18	BENELLI	2017	500
	P16	TRK502	2017	500
	P25	GT600 RESTRICTED	2014-16	600
	P25	BN 600 RESTRICTED	2013-on	600
	P36	502C	2019-on	500
	VELVET DUSK	VELVET DUSK	2002-06	383
BETA	BETA	FUPA RR E3	2018-on	293
	BMA RR	RR350 15	2018-on	349
	BMA RR	RR390 16	2018-on	386
	BMA RR	RR430 17	2018-on	431
	BMA RR	RR480 18	2018-on	478
	FUPA E5/Xtrainer	E5 00	2015-on	293
	FUPA E5/Xtrainer	E8/03	2016-on	293
	FUPA RR E3	RR 2T 300	2012-on	293
	FUPA RR E3	RR350 20 & RR350 15	2016-on	349
	FUPA RR E3	RR390 31 & RR390 16	2016-on	386
	FUPA RR E3	RR430 32 & RR430 17	2016-on	431
	FUPA RR E3	RR480 33 & RR480 18	2016-on	478
	RR E3	RR350	2011-on	349
	RR E3	RR400	2010-14	398
	RR E3	RR450	2010-14	449
	RR E3	RR520	2010-11	498
	RR300 2T	RR300 2T	2018-on	293
	RR350 4T	RR350 4T	2019-on	349
	RR390 4T	RR390 4T	2019-on	386
	RR430 4T	RR430 4T	2019-on	431
	RR450	RR450	2000-08	448
	RR480 4T	RR480 4T	2019-on	478
	RR525	RR525	2000-08	510
	XTRAINER 300 2T	XTRAINER 300 2T	2019-on	293
BMW	C400X	0C09/C400X	2018-on	350
	C400GT	0C06, C400GT	2018-on	350
	C650	C600 SPORT	All	647
	C650	C650 GT/Sport	All	647
	F650	FUNDURO	1995-00	652
	F650	G650 GS	2009-2016	652
	F650	G650 GS Sertao	2012-2016	652
	F650CS	SCARVER	2002-06	652
	F650CS	SE ROAD	2004-06	652
	F650GS	DAKAR	2000-08	652
	F650GS	F650GS	2000-10	652
	F650ST	F650ST	1998	652
	G 450 X	G 450 X	2008-10	450
	G310	G310R-0G01	2016-on	313
	G310GS	G310GS-0G02	2016-on	313
	G650GS	Sertao	All	650
	R45	R45	All	453
	R50	R50	All	499
	R60	R60	1960-78	590
	R65	R65	1978-88	650

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	R65LS	R65LS	1981-86	650
	R69	R69S	1961-67	600
BOLWELL	LM25W	FIRENZE	2009	263
BOLLINI	All models	All models under 250	All	250
BRAAAP	Moto4	Moto Range, Cruiser 400	2021-on	400
	ST	450	2016-on	450
	ST400	Shadow	2022-on	367
BSA	A50	A50	1964-70	500
	A65	A65	1966-69	650
	A7	A7	1961	500
	B40	B40	1969	350
	B44	B44	1967-71	440
	B50	B50	1971	495
	B50SS GOLDSTAR	B50SS GOLDSTAR	1971	498
	GOLD STAR	GOLD STAR	1938-63	500
	LIGHTNING	LIGHTNING	1964	654
	SPITFIRE MKIII	SPITFIRE MKIII	1967	650
	THUNDERBOLT	THUNDERBOLT	1968	499
BUELL	Blast	STREET FIGHTER	2002-07	491
BUG	SEE KYMCO			
BULTACO	ALPINA	ALPINA	1974	350
	FRONTERA	FRONTERA	1974	360
	SHERPA	SHERPA	1974	350
CAGIVA	360WR	360WR	1998-02	348
	410TE	410TE	1996	399
	610TEE	610TEE	1998	576
	650 ALAZZURA	650 ALAZZURA	1984-88	650
	650 ELFANT	650 ELFANT	1985-88	650
	CANYON 500	DUAL SPORTS	1999-06	498
	CANYON 600	DUAL SPORTS	1996-98	601
	RIVER 600	RIVER 600	1995-98	601
	W16 600	W16 600	1995-97	601
CAN-AM	CAN-AM RYKER	ROTAX 600 ACE	2019-on	599
CCM	GP Series	GP450-1(A1 30kW)	2015-16	450
	GP Series	GP450-2(A1 30kW)	2015-16	450
CFMOTO	CF300-7F	300CL-X	2023-on	292
	CF 400-6F	450SR	2022-on	449
	CF 400-10F	450CL-C	2024-on	449
	CF450MT	CF450MT	2024-on	449
	CF450NK	CF450NK	2024-on	449
	CF 650	CF650NK-LAM	2012-on	649
	CF 650	CF650TK-LAM	2013-17	649
	CF 650	650NK-LAM	2016-18	649
	CF 650	650MT	2016-on	649
	CF 650	650GT	2019-on	649
	CF 650 (400NK)	400NK	2016-18	400
	CF250-A	CF300, 300NK	2019-on	292
	CF250-A	CF300SR, 300SR	2020-on	292
CHANGJIANG	CJ650B with sidecar	Nomad, Tourer	2023-on	649
COSSCK	650	Ural	1974	649
DERBI	MULHACEN	MULHACEN	2008	659
	RAMBLA	RA 300	2010	278

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
DNEPR	K650	K650	1972	650
	K650	K650 DNEPR	1967-74	650
	MT9	MT9	1974	650
DUCATI	400 MONSTER	400 MONSTER	2002	398
	400 SIE	400 S I E monster		398
	400 SS JUNIOR	400 SS	1989-96	398
	400SS	400SS	1992-95	398
	500 DESMO	500 Sport Desmo	1977-83	497
	500GTL	500GTL	1975-77	497
	500SL	PANTAH	1984	499
	600 MONSTER	600 MONSTER	1994-01	583
	600 MONSTER	DARK	1998-01	583
	600 S	600 SUPERSPORT	1994-97	583
	600M	600M	1994-01	583
	600SL	PANTAH	1980-84	583
	600SS	600SS	1994-98	583
	620 MONSTER LITE	M620 LITE	2003-07	618
	620 MULTISTRADA LITE	MTS620 24.5Kw	2005-07	618
	659 Monster	Monster 659	All	659
	DM 350	350	pre 85	350
	DM 450	450	pre 85	448
	DM450	DM450	1972	450
	DM500	DM500	1981-84	498
	F3	350 F3	1986-1989	349
	F4	400 F4	1986	400
	KA (Scrambler)	00AA Sixty2	2015-on	399
	M4	M620ie LITE	2003-04	620
	M5	Monster 659	2011	659
	MD	02AU	2017	659
ELSTAR SHINERAY	XY400	CAFÉ RACER	2018	397
	XY400	CAFÉ RACER F	2018	397
	XY400	CLASSIC C	2018	397
	XY400	SCRAMBLER C	2018	397
	XY400	WB400 & WB400c	2015-17	397
ENFIELD	BULLET	CLASSIC	1993-08	499
	BULLET	DELUXE	1993-08	499
	BULLET	ELECTRA ROAD	2006-08	499
	BULLET 350	DELUXE	1988-01	346
	BULLET 350	SUPERSTAR	1988-95	346
	BULLET 350	CLASSIC	1993-01	346
	BULLET 500	500	1995-08	499
	BULLET 65	ROAD	2003-04	499
	LIGHTNING	ROAD	2000-08	499
	MILITARY	ROAD	2002-08	499
	TAURUS	DIESEL	2001	325
FANTIC	CA50	Flat Track	2020-on	449
	CA50	Scrambler	2020-on	449
	CA50	Rally	2020-on	449
	TZ	EC300	2011-12	300
	TZ	Gas Gas EC30	2012	300
GAS-GAS	4E	EC 30	2017	299
	4E	EC25	2017	299
	4E (IPA 48807)	EC 30	2018	299

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	CONTACT ES	280 ES	2018	272
	EC ENDURO	EC30	2016-17	299
	EC Series	EC300	2001-on	293
	EC Series	EC350F	2021-on	350
	EC300	SM SUPERMOTARD	2002	299
	EC300	ENDURO	2001-on	299
	EC300	EC300	2024-on	293
	EC350F	EC350F	2024-on	299
	EC400	FSE ENDURO	2002-03	399
	EC450	FSE ENDURO	2003-05	449
	EC450	FSE SUPERMOTARD	2003-08	449
	EC450	FSR ENDURO	2006-12	449
	EC450F	EC450F	2024-on	450
	EC500F	EC500F	2024-on	510
	FS 400	FS40A	2006	398
	FS 450	FS45	2006	443
	FS 500	FS50 (503)	2006-2009	503
	FSE 400	400	2002	398
	FSE 450	450	2003-08	398
	PAMPERA	320 TRAIL	1998-02	333
	PAMPERA	400 TRAIL	2006-08	399
	PAMPERA	450	2007-09	443
	SM400	SUPERMOTARD	2003-08	399
	SM450	SUPERMOTARD	2003-08	443
	TT300	EC300	1998-08	295
GILERA	FUOCO 500	FUOCO 500	2007-13	493
	NEXUS 500	NEXUS 500	2003-08	460
HARLEY DAVIDSON	LWZ Series	X350	2023-on	353
	LWZ Series	X500	2023-on	500
	SS350	Sprint	69-1974	350
	XGS SERIES	Street 500 -XG500 16MY	2014-15	494
	XGS SERIES	Street 500	2015-on	494
HONDA	600V TRANSALP	600V	1988	583
	BROS400	BROS400	1988	399
	C70	DREAM	pre 1970	305
	CB300 (FA)	CB300FA	2014-18	286
	CB300R	CBF300NA	2018-on	286
	CB350	CB350	1969-72	348
	CB350F	CB350F	1972-74	325
	CB360	CB360	1973-76	360
	CB400	CB400	1981-2014	395
	CB400 ABS	CB400 ABS	2008-2016	399
	CB400F	CB400F	1975-77	408
	CB400N	CB400N	1981	395
	CB400T	CB400T	1977	408
	CB450	CB450	1967-75	450
	CB500 FOUR	CB500-FOUR K, K1, K2	1971-73	498
	CB500 TWIN	CB500T	1974-78	498
	CB500F	CB500FA/F	2012-on	471
	CB500FA	CB500FA	2023-on	471
	CB500XA	CB500XA	2013-on	471
	CB500XK	CB500XK	2019-on	471
	CB550	CB550	1974-78	544

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	CB650	CB650	1979-85	627
	CB650F	CB650FA-LTD-16ym	2015-2017	649
	CB650F	CB650FL	2015-20	649
	CBR300R	CBR300R	2014-16	286
	CBR300R	CBR300RA	2014-18	286
	CBR500R	CBR500RA, CBR500R (ABS)	2012-on	471
	CBR650F	CBR650F LAMS CBR650F LAMS (CBR650FL)	2015-2019	649
	CBR650R	CBR650R ABS, CBR650R E-Clutch	2019-on	649
	CBX550	CBX550F	1982-85	572
	CJ360	CJ360	1976	356
	CL350	CL350 Scrambler	1971-73	325
	CL360	CL360 Scrambler	1974-76	356
	CL450	CL450 Scrambler	1965-77	444
	CL500	CL500	2023-on	471
	CL77	CL77 Scrambler	1965-68	305
	CMX500	CMX500 S ABS,CMX500	2016-on	471
	CMX500A	CMX500A	2016-20	471
	CRF300	CRF300 Rally	2020-on	286
	CRF300	CRF300L	2020-on	286
	CRF400R	CRF400R	2013	399
	CRF450L	CRF450L	2018-on	449
	CRF450L	CRF450L2019YM	2018-on	449
	CRF450X	CRF450X	2005-09	449
	CX500	CX500, CX500A, CX500C, CX500EC	1977-84	495
	DEAUVILLE	NT650V	2000-07	647
	FJS400	SILVERWING 400	2009-10	399
	FJS600D	SILVERWING 600	2006-09	582
	Fortza 300	NSS300 Forza	All	279
	FT500	FT500 Ascot	1982-84	498
	FTS600D	SILVERWING	2006-08	582
	GB350	GB350	2023-on	348
	GB350C	GB350C	2024-on	348
	GB350S	GB350S	2025-on	348
	GB400	GB400	All	400
	GB400TT	GB400 TT	All	399
	GB500	GB500 TT	1975-91	498
	GL400	GL400	1985	396
	GL500 S	Silverwing 500	1981-82	498
	NF02	SH300	2009	279
	NSS300	NSS300 Forza	2013-on	279
	NSS350	NSS350 Forza	2020-on	330
	NT400	NT400	1989-92	400
	NT650V	DEAUVILLE	2003-07	647
	NTV650	REVERE	1989-92	647
	NX500	NX500/NX500 E-Clutch	2024-on	471
	NX650	DOMINATOR	1988-00	644
	OBI RVF400 VFR400	OBI RVF400 Otobai import model only	All	400
	REVERE	REVERE	1990	647
	SH300i	SH300i	2009-11	297
	SL350	SL350	1970-72	348
	Steed	steed	2002	398
	VT400	VT 400	All	398

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	VT400C	SHADOW	All	399
	VT500E	Ascot (VT500E)	1983-87	491
	VT600C	VT600C	1993-00	583
	VT600C	SHADOW VLX	1988-2008	583
	XBR500	XBR500	1986-89	499
	XBR500S	XBR500SH	1986-89	499
	XL350	XL350	1973-74	339
	XL500	XL500	1979-84	498
	XL600R	XL600R	1983-88	589
	XL600RMG	XL600RMG	1986-88	591
	XL600VH	XL 600 TRANSALP	1983-89	583
	XL650	TRANSALP	2005	647
	XL650V	TRANSALP	2002-08	647
	XR350	XR350	1983	339
	XR350R	XR350R	1983-86	339
	XR350R	XR350R	1985-86	353
	XR400	XR400	1996-08	397
	XR400 MOTARD	XR400M	1996-08	397
	XR400R	XR400R	1996-08	397
	XR500R	XR500R	1983-84	498
	XR600R	XR600R	1985-00	591
	XR650L	XR650L/XR650R	2001-06	644
	XR650R	XR650R KSS and MSS (only)	2004-05	649
	XR650R	XR650R (Australian version only)	1999-2001	649
HUNTER	DD350E-2	BOBBER	2011-13	320
	DD350E-6C	DAYTONA	2010-13	320
	DD350E-6C	SPYDER	2010-13	320
HUSABERG	FE01	FE450 MY05 (Ab)	2004	449
	FE01	FS650 MY05 (Db)	2004	628
	FE350	ENDURO	All	350
	FE390	FE390	2009-12	393
	FE400	ENDURO	All	399
	FE450	ENDURO	2008-14	449
	FE501	ENDURO	2012-on	510
	FE501E	ENDURO	1997-12	501
	FE570	ENDURO	2008-10	565
	FE600E	ENDURO	1997-00	595
	FE650E	ENDURO	2004-08	628
	FE650E	ENDURO	2000-04	644
	FS450	SUPERMOTARD	2008-10	449
	FS450E	ENDURO	2004	449
	FS570	SUPERMOTARD	2009-10	565
	FS650C/E	SUPERMOTARD	2004-08	628
	FS650E	SUPERMOTARD	2002-04	644
	TE300	TE Series	2010-14	293
HUSQVARNA	300WR	WR300	2008-12	298
	310TE	TE310 A3	2009-13	303
	310TE	TE310 A2	2008-10	298
	350TE	TE350	1995	349
	400SM	SUPERMOTARD	2002-04	400
	400TE	ENDURO	2000-01	400
	410TE	ENDURO	1998-00	400
	410TE	ENDURO	1994-97	415

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	450SM/R/RR	SUPERMOTARD	2003-08	449
	450TC	MOTOCROSS	2001-08	449
	450TE	ENDURO	2001-10	449
	450TE-IE	ENDURO	2007-08	449
	450TXC	TRAIL	2007-08	449
	510SM	SUPERMOTARD	2006-10	501
	510TC	MOTOCROSS	2004-07	501
	510TE	ENDURO	1986-2008	510
	510TE-IE	TE510IE	2008	510
	570TE	570TE(RP)	2000	577
	610SM	SUPERMOTARD	2000-08	577
	A6 SMR 449	A600AB	2010-12	450
	A6 SMR 511	A601AB	2010-12	478
	A6 SMR 511	A602AB	2012	478
	A6 TE 449	A600AATE449	2010-13	450
	A6 TE 511	A601AATE511	2010-13	478
	A8	0H11B 35kW	2013	652
	AE430	ENDURO	1986-88	430
	FE	FE350	2014-on	350
	FE	FE450	2014 on	449
	FE	FE450	2016-on	450
	FE	FE501	2014 on	501
	FE	FE501	2016-on	510
	Pilen Series	VP 401	2018-on	373
	Pilen Series	SP 401	2018-on	373
	SMR449	SMR449	2011	449.6
	SMR511	SMR511	2012	447.5
	SMS630	A401AB SMS630	2010-on	600
	SVARTPILEN	SP 401	2023-on	399
	TE	TE300	2014 on	298
	TE	TE300	2016-17	293
	TE449	Enduro 2014	2013	449.6
	TE510 (A2)	Enduro 2013	2006-2013	477.5
	TE610	TE610(RP), dual sports	2000-on	577
	TE630	A401AA TE630	2010-on	600
	TR650	TR650 Terra	2013	652
	TR650 Strada	0H11F 35kW	2013-on	652
	TR650 Terra	0H11B 35kW and 0H11D 35kW	2013-on	652
	VITPILEN	VP 401	2023-on	399
	WR260	ENDURO	1990-91	260
	WR300	ENDURO	2010-13	293
	WR360	ENDURO	1991-03	349
	WR400	ENDURO	1984-88	396
	WR430	ENDURO	1988	430
HYOSUNG	GT650 EFI	GT650EFI Lams	All	647
	GT650R EFI	GT650R EFI Learner	All	647
	GV650C/S	Lams model	All	647
INDIAN	VELO	VELO	1969	500
JAWA	350	350	1974	350
	634 ROAD	634 ROAD	1984-85	343
	638 ROAD	638 ROAD	1985-86	343
JONWAY	MALIBU	MALIBU 320	2012	320

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
KAWASAKI	EL450A	EL450A L, EL450B L, Eliminator, Eliminator SE	2023-on	451
	EN400	Vulcan	1986	400
	EN450	450LTD	1985-87	454
	EN500	Vulcan	1990-02	500
	EN650B	Vulcan S ABS/ABS L	2014-current	649
	EN650B	EN650E ABS L 1 & 2	2016-17	649
	EN650C	VULCAN S, VULCAN S L	2016-on	649
	ER300B	ER300B (Z300 ABS)	2015-on	296
	ER-5	ER500	1999-06	498
	ER-650C	ER-6nL	2009-10	649
	ER-650C	ER-6nL ABS	2009-16	649
	ER650F	ER-6nl ABS learner model	2012-2016	649
	ER650H	ER650H LAMS (Z650L)	2016-on	649
	ER650H	ER659K LAM (Z650L)	2019-on	649
	ER650H	ER650M LAMS (Z650RS)	2021-on	649
	ER650H	ER650R L	2023-on	649
	ER650H	ER650S L	2021-on	649
	EX300A (Ninja 300)	EX300B Ninja/special (A&B)	2012-18	296
	EX300B	EX300B	2015-on	296
	EX400	GPX 400R	1987-94	399
	EX400G	Ninja 400 & EX400G	2018-on	399
	EX400G	KAWASAKI	2018	399
	EX400G	Z400 and ER400D	2019-on	399
	EX400G	Ninja 400 & EX400G	2018-on	399
	EX500G	Ninja 500 & Z500	2024-on	451
	EX500J	Ninja 500SE/KRT Edition	2025-on	451
	EX650F	Ninja 650L (2012)	2012	649
	EX650K	EX650S L	2021-on	649
	EX650K (LAMS)	Ninja 650 L	2016-on	649
	GPZ550	GPZ550	1981-90	553
	GT550	Z550	1983-88	553
	KL600	KLR600	1984-87	564
	KL650	KLR650	1987-99	651
	KL650E	KLR650, KLR650S	2013-on	652
	KL650E	KLR650 Adventure	2021-on	652
	KLE300C	KLE300C VERSYS-X 300	2017-on	295
	KLE500	DUAL SPORTS	1992-08	498
	KLE500	KLE500	1992-2008	498
	KLE650F	Versys 650L ABS	2014-on	649
	KLE650F	KLE650F ABS L & ABS L MY17	2016-on	649
	KLE650F	KLE650H L	2021-on	649
	KLR600	KL600	1984-87	564
	KLR650E	KL650E	1987-2012	651
	KLX300	KLX300	2025-on	292
	KLX300R	KLX300R	1996-on	292
	KLX400	KLX400	2002-05	400
	KLX450A	KLX450A, KLX450R	2007-on	449
	KLX650	KLX650	1989-95	651
	KLX650R	ENDURO	1993-04	651
	KZ400	KZ400	1974-84	398
	KZ440	KZ440D OR Z440	1980-85	443
	KZ500	KZ500	1979	497

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	KZ550	KZ550, ZX550A	1981-86	547
	LE650D	Versys 650L ABS	2010	649
	LE650D	Versys 650L ABS	2011-on	649
	LTD440	LTD440	1982	443
	LX400	LX400 Eliminator	1989	398
	Ninja 650	Ninja 650RL ABS	2009-11	649
	Ninja 650	Ninja 650L ABS	2011-16	649
	Ninja 650 L model	Ninja 650RL	2009	649
	S2	S2	1972	346
	S3	S3	1974	400
	W400	EJ400AE	2006-09	399
	Z400B2	KZ400B2	1979	398
	Z400D	KZ400D	1975	398
	Z500	Z500	1979-84	498
	Z500 SE	Z500 SE	2024-on	451
	Z500 (ER500)	Z500 (ER500)	2024-on	451
	ZR550	ZEPHYR 550, ZR550A	1983-99	553
	ZZ-R400	ZZR400	1984-92	399
KTM	2T-EXC	300 EXC	2012-on	293
	300 exc	300exc	All	300
	300EXC	ENDURO	1984-2011	293
	300EXC-E	ENDURO	2007-08	293
	300GS	ENDURO	1990-95	280
	350EXC Special-R	ENDURO	2005-06	350
	350EXC-F	ENDURO	2011-on	347
	360EXC	ENDURO	1996-98	360
	380EXC	ENDURO	2000	368
	390 Duke	390 Duke	All	390
	400EXC	ENDURO	2008-11	393
	400GS	ENDURO	1993-99	400
	400SC	400SC	1996-98	400
	400TE	400TE	2001	400
	450EXC	ENDURO	2002-07	448
	450EXC	ENDURO	2005-11	449
	450EXC	ENDURO	2011-on	449
	4T-EXC RACING	350 EXC-F	2012-on	350
	4T-EXC RACING	450 EXC	2012-on	449
	4T-EXC RACING	500 EXC	2012-on	510
	500EXC	ENDURO	2011-on	510
	500GS	ENDURO	1984-91	553
	510EXC	ENDURO	1999-02	510
	520EXC	ENDURO	2000-02	510
	525EXC	ENDURO	2002-05	510
	525EXC-R	ENDURO	2005-07	510
	530EXC	ENDURO	2008-11	510
	600 ENDURO	ENDURO	1987-93	553
	600 ENDURO INCAS	ENDURO	1989-90	553
	625SMC	625SMC	2004	609
	640 4T -EGS	640 LC4-EMY04	2004-05	625
	640 4T -EGS	640 LC4-MY05	2004-05	625
	660 SMC	4T-EGS	2004	654
	Adventure	390 Adventure	2020-on	373
	Freeride	Freeride (MY12 on)	2012	350

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	IS DUKE	390 DUKE (C3)	2013	373
	IS RC	RC 390	2016-on	373
	Rally	450 RALLY	2017-on	449
	Rally	690 RALLY	2017	654
	RC390	RC390	all	390
KYMCO	AGILITY 300	T4 (300)	2020-on	276
	AK Series	AK 550	2017-on	550
	Downtown Series	Downtown GT (E90000)	2024-on	321
	DT Series	DT X360/E70000	2021-on	321
	V2	Downtown 350i (V200010, V20020, V20030, V23010-V23000, C71100))	2015-on	321
	XCITING S 400	D62001 & D62000	2019-on	400
	X-Town	KS60A (300i)	2016-on	276
LAMBRETTA	All model	Lambretta	pre 2008	Under 660
	G350	G350	2025-on	330
	X300	X300A	2024-on	275
LARO	DD350E-6C	Pro Street 350	2011	320
	SPT series	SPT350	2011	320
LAVERDA	500	500	1979	497
LIFAN	All model	All models	2009-10	Under 300
LIFENG	Regal Raptor	CRUISER 350	2011	320
MAICO	Enduro	500E	1984-88	488
MATCHLESS	G12	G12	pre 1966	646
	G80	HARRIS	1988-90	494
	G80	G80	pre 1963	497
MBK	FALCONE	YAMAHA XT660R/X	2005-08	660
MONTESA	COTA 330	TRIAL	1985-86	328
	COTA 335	TRIAL	1986-88	327
	COTA 348T	TRIAL	1984-87	305
	COTA 350	TRIAL	1984-85	349
MOTO GUZZI	350 GT	350 GT	1992	350
	Falcone	Falcone	1972	498
	V35	V35	1977-90	346
	V50	V50	1977-79	490
	V50	Monza	1980-85	490
	V65	V65	1982-94	643
	V65	Lario	1984-89	643
MOTO MORINI	3.5 ROAD	3.5 ROAD	1984-85	344
	350 SPORT	350 SPORT	1974-85	344
	500 CAMEL	TRAIL	1984-86	479
	500 SEI	500 SEI	1984-85	479
	500 STRADA	500 STRADA	1977-85	479
MUZ	BAGHIRA	ENDURO	1999-02	660
	MASTIFF	SUPERMOTARD	1999-02	660
	SKORPION	REPLICA	1998-02	660
	SKORPION	SPORT	1998-02	660
	SKORPION	TRAVELLER	1998-02	660
	SKORPION	TOUR	1998-02	660
MV AGUSTA	350	350	1972-76	349
NORTON	650SS	650SS	1961-68	650
	ES2	ES2	pre 1963	490
	MANX	MANX	All	Under 660
	MODEL 50	MODEL 50	1933-63	348

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	MODEL 88	DOMINATOR	pre 1966	497
	NAVIGATOR	NAVIGATOR	1964	350
OZ TRAIL	FUN 500	FUN500	pre 2008	500
PANTHER	MODEL 100	600	pre 1963	598
	MODEL 120	650	pre 1966	645
PEUGEOT	GEOPOLIS	AEAA	2007-08	399
	METROPOLIS	AA	2018-on	399
	SATELIS	AEAA	2007-08	399
	SATELIS	AFAA	2007-08	493
	XP400 OR Y2	XP400	2023-on	399
PIAGGIO	All Models	All models	2010-17	Under 350
	PSI M59 (MP3 400)	M59101 (400ie RL)	2006-08	399
	PSI M52	M52101 XEVO 400ie	2006-08	399
QJ MOTORCYCLES	BJ60	BJ60	All	600
	P25	BJ600	All	600
RICKMAN	650	Triumph	1964	649
RIEJU	MR5E	MR300 ENDURO	2020-on	293
	MR5E	MR300 ENDURO PRO	2020-on	293
RIYA	RY300T (RY)	RY300T	2012-15	288
ROYAL ALLOY	GP300	GP300	2020-on	278
ROYAL ENFIELD	All models	All models under 660	till 2014	Under 660
	CLASSIC	Classic 350	2016-on	349
	CNEX	Bear 650	2024-on	648
	CNEX	CNEG	2018-on	648
	CNEX	CNEH	2018-on	648
	CNEX	CNEG (CONTINENTAL GT 650)	2018-on	648
	CNEX	CNEH (INTERCEPTOR GT 650)	2018-on	648
	CNEX	Shotgun 650	2024-on	648
	CNEX	Super Meteor 650	2023-on	648
	CNEX	Super Meteor 650 Touring	2023-on	648
	D4A5C	Himalayan	2016-on	411
	D4A5C EFI	Himalayan	2016-on	411
	GUERRILLA	450 G1	2024-on	450
	HIMALAYAN 450	G1	2024-on	452
	HUNTER	Hunter 350	2022-on	349
	J1	Bullet 350	2023-on	349
	Meteor	Meteor 350	2020-on	349
	UMI BULLET	U3S	2015-19	346
	UMI BULLET	BULLET 500 CKE	2015-19	499
	UMI CONTINENTAL	CONTINENTAL GT	2015	535
RS HONDA	XR400M	MOTARD	2005-08	397
RUDGE WHITWORTH	650	Rudge	pre 1961	650
SHERCO	S4	ENDURO 450	2007-2010	448
	S4	ENDURO 510	2007-2010	510
	S4	ENDURO 300	2010	290
	S6	300 2ST	2014-on	293
	S6	300 4ST	2015-on	303
	S6	450 4ST	2015-on	449
	S6	480ST	2021-on	479
	S6	500 4ST	2018-on	510

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
SUZUKI	AN400	AN 400	2004-on	400
	AN400	BURGMAN	2008-18	400
	AN400	AN400	2008-18	400
	AN650	BURGMAN	2008-18	638
	Burgman	Burgman 400ABS (AN400A)	2014-on	400
	DL650	DL650 AUE & DL650X AUE	2016-20	645
	DL650AUE	V Strom	2014-on	645
	DL650XAUE	V-Strom 650 XT learner approved	2015-on	645
	DR350	All	1991-98	349
	DR400	DR400	1999	400
	DR500	All	1981-84	498
	DR600	DR600	1984-86	589
	DR600R	DR600R	1985-90	598
	DR650	All	1990-08	644
	DR650SE	DR650SE	1995-on	644
	DR-Z400E	DR-Z400E	All	398
	DR-Z400E	DR-Z400 (2006 MY~)	All	398
	DR-Z400E	DR-Z400	All	
	DR-Z400S	DR-Z400S	2005-on	398
	DR-Z4S	DR-Z4S	2025-on	398
	DR-Z4SM	DR-Z4SM	2025-on	398
	DR-Z400SM	DR-Z400SM	2005-on	398
	Gladius 650 LAMS (SFV650U)	Gladius 650 LAMS	2009-17	645
	GN400	GN400	1980-81	400
	GR650	All	1983-88	651
	GS400	GS400	1976-82	400
	GS450	All	1981-89	450
	GS500	GS500	2000-13	487
	GS500E	GS500E	1976-99	492
	GS500F	GS500F	2003-13	487
	GS550	All	1977-82	549
	GSR400	GSR400	2006-08	398
	GSX400	F	1981-04	398
	GSX400	E	1981-84	398
	GSX650F	GSX650/FU	2008-12	656
	GT380	GT380	1973-78	380
	GT500	GT500	1976-78	500
	GT550	GT550	1973-78	550
	KATANA 550	KATANA 550	1981-83	550
	LS650S4	Boulevard S40	2014-on	652
	LS650	SAVAGE	1986-89	652
	LS650	LS650	2018	652
	PE400	PE400	1980-83	400
	RE5	ROTARY	1974	500
	RMX450 (market name RMX450Z)	RMX450	2014-15	449
	SFV650U	SFV650U	2009-17	645
	SP370	ENDURO	1978	370
	SV650-3	SV650 UA	2015-on	645
	SV650S LAMS	SV650SU LAMS Gladdius	2008/2013	645
	SV650SL	SV650SU LAMS	2008-13	645
	SV650UA	SV650 LAMS ABS	2016-on	645
	SV650	SV650UA LAMS	2016-on	645

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	SV650XAUE	SV650XA LAMS	2018-on	645
	SVF650 (Market name-Gladius)	SVF650 U/UA	2009-2014	645
	T350	T350 Rebel	1969-75	315
	T500	T500	1970-74	500
	TM400	TM400 Cyclone	1973	396
	TS400	TS400	1976	400
	XF650	FREEWIND	1997-01	644
SWM	A1	01/AA and 01/AB	2015-2017	600
	A2	01/AA	2016	300
	A2	03/AA and 03/AB	2016	500
	A3	00-01-02	2016	445
	B3	Silver Vase, Gran Milano	2016-on	445
SYM	All Models	All models under 400	2008-12	400
	LN	GTS 300i Sport	2015-16	278
	LX	MaxSYM 400i	2012-on	399
TGB	All Models	All models under 300	2012	300
TM	3002T	ENDURO	2010	297
	300E	ENDURO	2000-08	294
	400E	ENDURO	2002-03	400
	4504T	ENDURO	2010	450
	450E	ENDURO	2003-08	449
	450MX	450MX	2008	449
	5304T	ENDURO	2010	528
	530E	ENDURO	2003-08	528
	530MX	530MX	2008	528
TRIUMPH	21	21	1963	350
	DAYTONA 500	DAYTONA 500	1967-73	490
	HD Series	HD418MY	2017	660
	L Series	TRIDENT 660	2020-on	660
	L Series	TIGER SPORT 660	2022-on	660
	Street triple	LAMs Street Triple 659 L67Ls7	2014-on	659
	T010	Speed 400	2023-on	398
	T010	Scrambler 400 X	2023-on	398
	T010	Scrambler 400 XC	2025-on	398
	T100	TIGER	pre-1970	498
	T120	BONNEVILLE	1959-1974	649
	TR5	TROPHY	1969	449
	TR6	TROPHY	1961-73	649
	TR7	TIGER	1971	649
	Trident 660	Trident 660-LAMS	2024-on	660
	TRIBSA	TRIBSA	1960-70	650
	Z010	TF450-E	2025-on	450
UBCO	2018 2X2	UBCO	2018	
URAL	DNEPR	DNIEPNER	1974	650
	K650	K650	1967-74	650
	MT9	MT9	1974	650
	THRUXTON	THRUXTON	1965-67	499
VESPA	All Models	All models	until 1/09/2013	50-300
	GTS 300	GTS 300 (Super/Sport/Super Sport/Tech)	2008-on	278
	GTV 300	MD3109	All	278
	PSI M45	M45200 300 S/SS	2016-on	278
	PSI M45	M45202 300 ABS	2018-on	278
	PSI M45	M45710 300 S/SS	2018-on	278

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	PSI M45	M45715 300 S/TECH	2019-on	278
	PSI M45	M45710 300	2018	278
	PSI M45	M45719 GTS 300 SS HPE	2020-on	278
	PSI M45	M45724 GTS 300 SG	2020-on	278
	PSI M45	M45200 300 S/SS	2016-on	278
	PSI M45	M45202 300 ABS	2018-on	278
	PSI M45	M45710 300 S/SS	2018-on	278
	PSI MA3	MA330 300 E4 (GTS/SUPER/SS)	2016-17	278
VOR	400 ENDURO	400 ENDURO	2000	399
	450 ENDURO	450 ENDURO	2002	450
	500 ENDURO	500 ENDURO	2001	503
	530 ENDURO	530 ENDURO	2001	530
	VOR ENDURO	400SM	2000-01	399
	VOR ENDURO	500SM	2000-01	503
XINGYUE	XY400Y	XY400Y	2008-09	400
YAMAHA	CZD300 (X-Max300)	CZD300-A	2016-on	292
	DT400	DT400	1976-81	400
	FZ600	FZ600	All	600
	FZ6R	FZ6R	All	600
	IT426	IT426	1987	426
	IT465	IT465	1987	465
	IT490	IT490	1983-85	490
	MT 07	MT07 LAMS, MTN660-A	2015-on	655
	MT 07	MT07, MTN660	2015-on	655
	MT-03	MT03	2011 on	660
	MT-07	MT-07 LAMs	2015-on	655
	MTM660	XSR700	2016-on	655
	MTN320	MTN320-A	All	321
	MTT660-A	RM 161	2016-17	655
	MX400	MX400	1976	400
	RD350	RD350	to 1975	350
	RD400	RD400	1976-80	398
	RT2	RT2	1970	360
	RT350	RT350	1972	347
	SR400	SR400	All	400
	SR500	SR500	1978-1981	499
	SRX400	SRX400	1985-90	400
	SRX600	SRX600	1986-96	608
	SZR660	SZR660	1997	659
	T MAX	Tmax 530	All	530
	Tenere	Tener	All	660
	Tricity 300 (MWD300)	Tricity 300 (MWD300)	2020-on	292
	TT350	TT350	1986-01	346
	TT500	TT500	1975	500
	TT600	TT600	All	595
	TT600E	TT600E	All	595
	TT600R	TT600R	All	595
	TX650	TX650	1976	653
	WR400F	WR400F	1998-2000	399
	WR426F	Belgarda import ONLY	2001	426
	WR450F	WR450F	All	450
	WR450F	WR450F (2GC)	All	449
	XJ550	XJ550	1981-83	528

MAKE	MODEL	VARIANT NAME	YEAR(S)	CAPACITY
	XJ6	XJ6FL/NL (25kW & 35kW)	All	600
	XJ6	XJ6SL (25kW)	All	600
	XJR400	XJR400	1999	400
	XJR400	4HM	2003	399
	XP500	TMAX (XP500)	All	499
	XP530	TMAX 530 (XP530)	All	530
	XP560D	TMAX (XP560D)	2022-on	562
	XS360	XS360	All	359
	XS400	XS400	All	391
	XS650	XS650	1972-1984	653
	XSR700	RM131	2015-17	655
	XT350	XT350	All	346
	XT500	XT500	All	499
	XT550	XT550	All	552
	XT600	XT600	All	590
	XT660R	XT660R	All	659
	XT660X	XT660X	All	659
	XT660Z T N R	XT660Z	All	660
	XTZ660	XT660Z Tenere	All	659
	XV400	XV400 Virago	1983	399
	XV535	XV535 Virago	All	535
	XVS400	XVS400 Dragstar	2001-08	399
	XVS650A/custom	XVS650 custom and classic	All	649
	XZ400	XZ400	1982	399
	XZ550	XZ550	1982-84	550
	YP400	MAJESTY	All	395
	YZF R3	YZF R3A	All	321
	YZF320-A	YZF320-A	2022-on	321
	YZF660	YZF-R7 LAMS	2021-on	649
ZHEJIANG	HT300T	Base	2015	275

An approved motor bike and motor trike must:

- Be the standard model and variant as specified on the above list; and
- Not be modified in any way that increases its power-to-weight ratio.

Schedule 2—Revocation

The *Motor Vehicles (Approval of Motor Bikes and Motor Trikes) Notice 2026 No 2* made on 26 March 2026.

(Gazette No.18, p.764) is revoked.

Stuart Gilbert
Deputy Registrar of Motor Vehicles

On 5 May 2025

RETAIL AND COMMERCIAL LEASES ACT 1995

Exemption

Pursuant to Section 77(2) of the *Retail and Commercial Leases Act 1995* (SA) I, Hon. Daniel van Holst Pellekaan, Small Business Commissioner for the State of South Australia, exempt the Lease agreement, which commenced on 1 July 2025, between the Minister for Climate, Environment and Water and Lutheran Church of Australia, SA & NT District Holdings Ltd (ABN 22 122 101 160), in relation to the whole of the land comprised in Certificate of Title Volume 6318 Folio 631, commonly known as Camp Kedron, from the entirety of the *Retail and Commercial Leases Act 1995*.

Dated: 30 April 2026

HON. DANIEL VAN HOLST PELLEKAAN
Small Business Commissioner

ROADS (OPENING AND CLOSING) ACT 1991

SECTION 24

NOTICE OF CONFIRMATION OF ROAD PROCESS ORDER

Road Closure—Portion of Public Road (Mount Magnificent Road), Mount Magnificent

By Road Process Order made on 2 July 2024, the Alexandrina Council ordered that:

1. Portion of Public Road (Mount Magnificent Road), Mount Magnificent, (being portion of allotment 41 in Deposited Plan 114700 and portion of allotment 33 in Deposited Plan 96312) situated adjoining Allotments 41 and 42 in Deposited Plan 114700 and allotment 31 in Deposited Plan 96312, more particularly lettered 'A' and 'B' in Preliminary Plan 21/0014 be closed.
2. Transfer the whole of the land subject to closure to Corey David Ross Duke and Joanna Duke in accordance with the Agreement for Transfer dated 19 January 2024 entered into between the Alexandrina Council and Corey David Ross Duke and Joanna Duke.

On 4 May 2026 that order was confirmed by the Minister for Planning conditionally upon the deposit by the Registrar-General of Deposited Plan 138842 being the authority for the new boundaries.

Pursuant to Section 24(5) of the *Roads (Opening and Closing) Act 1991*, notice of the Order referred to above and its confirmation is hereby given.

Dated: 7 May 2026

B. J. SLAPE
Surveyor-General

2021/09603/01

ROADS (OPENING AND CLOSING) ACT 1991

SECTION 24

NOTICE OF CONFIRMATION OF ROAD PROCESS ORDER

Road Closure—Unmade Road (Roehr Road), Ebenezer and Stockwell

By Road Process Order made on 26 August 2025, the Light Regional Council ordered that:

1. Portion of Unmade Road (Roehr Road), Ebenezer and Stockwell, situated adjoining Allotment 8 in Filed Plan 156177 and Section 3010, Hundred of Belvidere, more particularly delineated and lettered 'A' in Preliminary Plan 24/0041 be closed.
2. Issue a Certificate of Title to the Light Regional Council for the whole of the land subject to closure in accordance with the Application for Document of Title dated 26 August 2025.

On 4 May 2026 that order was confirmed by the Minister for Planning conditionally upon the deposit by the Registrar-General of Deposited Plan 139035 being the authority for the new boundaries.

Pursuant to Section 24(5) of the *Roads (Opening and Closing) Act 1991*, Notice of the Order referred to above and its confirmation is hereby given.

Dated: 7 May 2026

B. J. SLAPE
Surveyor-General

2024/08663/01

PUBLIC NOTICES

NATIONAL GAS LAW

Notice of Draft Determination and Draft Rule

The Australian Energy Market Commission (AEMC) gives notice under the National Gas Law as follows:

Under s 308, the making of a draft determination and related draft rule on the *Gas cyber security roles and responsibilities for AEMO* (Ref. GRC0091) proposal. Written requests for a pre-determination hearing must be received by **14 May 2026**. Submissions must be received by **18 June 2026**.

Submissions can be made via the AEMC's website. Before making a submission, please review the AEMC's privacy statement on its website, and consider the AEMC's Tips for making a submission. The AEMC publishes submissions on its website, subject to confidentiality and other considerations.

Written requests should be sent to submissions@aemc.gov.au and cite the reference in the title. Before sending a request, please review the AEMC's privacy statement on its website.

Documents referred to above are available on the AEMC's website and are available for inspection at the AEMC's office.

Australian Energy Market Commission
Level 15, 60 Castlereagh St
Sydney NSW 2000
Telephone: (02) 8296 7800
www.aemc.gov.au

Dated: 7 May 2026

TRUSTEE ACT 1936

PUBLIC TRUSTEE

Estates of Deceased Persons

In the matter of the estates of the undermentioned deceased persons:

CARROLL Rita Ruta late of 29-31 Austral Terrace Morphettville Retired Carer who died 10 April 2024
COAD Mervyn Alan late of 1 Steel Street Campbelltown Retired Leading Hand/Toolmaker who died 23 September 2025
FLORES Don late of 550 Portrush Road Glen Osmond of no occupation who died 31 October 2025
HASSAN Mustaque late of 390 Regency Road Prospect Chef who died 30 May 2024
LAWS Alan Clifford late of 26 Mark Road Elizabeth South Retired Tool Maker/Machinist who died 17 August 2024
LAWS Gisela late of 80 Moseley Street Glenelg South Retired Care Attendant who died 8 April 2025
PILKINGTON Leo John late of 39 Fisher Street Myrtle Bank Retired Clerk who died 12 June 2025
SCHWALBACH Bruno late of 1 Parkwood Grove Klemzig Retired Toolmaker who died 5 December 2025
THOROUGHGOOD Joan Margaret late of 206 Sir Donald Bradman Drive Cowandilla Retired Clinical Officer who died 16 August 2025
VERCO Winifred Glen Trevalvwyn late of 5-11 Sirius Avenue Hope Valley Retired Clerical Officer who died 18 December 2025
VINEY Edith Lorraine late of 151 The Terrace Port Pirie of no occupation who died 19 September 2025

Notice is hereby given pursuant to the *Trustee Act 1936* (SA), the *Succession Act 2023* (SA) and the *Family Relationships Act 1975* (SA) that all creditors, beneficiaries, and other persons having claims against the said estates are required to send, in writing, to the office of Public Trustee at GPO Box 1338, Adelaide 5001, full particulars and proof of such claims, on or before the 5 June 2026 otherwise they will be excluded from the distribution of the said estate; and notice is also hereby given that all persons indebted to the said estates are required to pay the amount of their debts to the Public Trustee or proceedings will be taken for the recovery thereof; and all persons having any property belonging to the said estates are forthwith to deliver same to the Public Trustee.

Dated: 7 May 2026

T. BRUMFIELD
Public Trustee

UNCLAIMED MONEY ACT 2021 (SA)

Unclaimed Moneys Held by Uniting Funds South Australia

Notice is hereby given pursuant to Section 5 of the *Unclaimed Money Act 2021* that the persons listed below are entitled to amounts of money which have remained unclaimed for a period of more than six (6) years and are held with Uniting Funds South Australia.

The unclaimed amounts relate to closed accounts. A full list of the names and last known addresses of persons entitled to the unclaimed monies, together with the corresponding amounts, may be inspected at:

Uniting Church SA
Level 2, 212 Pirie Street Adelaide SA 5000
during normal business hours (09:00—16:30) or refer to Schedule 1.

Persons having a claim to any of the listed amounts should submit written application to: Manager, Investment Services
GPO Box 2145 Adelaide SA 5001 or info@ucinvest.com.au

Applications must include sufficient proof of identity and any documentation supporting the claim to the funds.

Dated: 30 April 2026

TROY HAMPTON
Manager, Investment Services
Uniting Church SA

SCHEDULE 1—UNCLAIMED MONIES

Name	Amount $	Last Known Address
Miss Ethel Freda Smith	$50.00	Address Unknown
Miss Ashlee Jayne Thackeray	$274.60	11 Garnet Court, Highbury SA 5089
Miss Megan Elyse Thackeray	$327.86	11 Garnet Court, Highbury SA 5089
Mr E M Farleigh	$400.00	Address Unknown
Mr Benjamin James Niejalke	$500.83	6 Quigley Court, Aberfoyle Park SA 5159
Josephine To	$838.90	Unit 6/3 Julia Court, Collinswood SA 5081
Rev Mosese Vakasiuola Latu	$1,044.74	27 Orpington Street, Ashfield NSW 2131
Mr Phillip James Bayly	$1,330.61	Address Unknown
Miss Emily Judith Harwood	$2,143.18	38 Gleneagles Road, Mount Osmond SA 5064
Mrs Inga Ruth Lempp and Mr Constantino Pinto	$3,099.57	c/- CTID PO Box 234, Dilli
Mr Bruce and Mrs Margaret Napier	$12,243.98	8 High Street, Unley Park SA 5061

NOTICE SUBMISSION

The South Australian Government Gazette is published each Thursday afternoon.

Notices must be emailed by 4 p.m. Tuesday, the week of publication.

Submissions are formatted per the gazette style and a proof will be supplied prior to publication, along with a quote if applicable. Please allow one day for processing notices.

Alterations to the proof must be returned by 4 p.m. Wednesday.

Gazette notices must be submitted as Word files, in the following format:

- Title—the governing legislation
- Subtitle—a summary of the notice content
- Body—structured text, which can include numbered lists, tables, and images
- Date—day, month, and year of authorisation
- Signature block—name, role, and department/organisation authorising the notice

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