Government Gazette

OF

WESTERN AUSTRALIA

(Published by Authority at 3.30 pm)

No. 78] PERTH: WEDNESDAY, 5 AUGUST [1987

POISONS ACT 1964

POISONS REGULATIONS

1965

Reprinted under the *Reprints Act 1984* as at 22 July 1987.

WESTERN AUSTRALIA

POISONS ACT 1964

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Citation

1. These regulations may be cited as the *Poisons Regulations 1965*[1].

[*Regulation 1 amended in Gazette 12 October 1984 p. 3267.*]

Interpretation

2. In these regulations unless the context requires otherwise—

"Animal" includes bees, birds, cetaceans, crustaceans, fish, molluscs and reptiles;

"Approved Name" means—

(a) the common name given to any substance by the British Standards Institution or the Australian Standards Association; or

(b) the English name by which any poison or substance is described in the British Pharmacopoeia, the Australian Pharmaceutical Formulary, or the British Pharmaceutical Codex; or

(c) if the substance is not described in the British Pharmacopoeia, the Australian Pharmaceutical Formulary, or the British Pharmaceutical Codex, the approved name as published by the General Medical Council of Great Britain; or

(d) if the substance is not described in the British Pharmacopoeia, the Australian Pharmaceutical Formulary, or the British Pharmaceutical Codex, and has not been given an approved name by the General Medical Council of Great Britain, the name given to the substance in any standard book approved by the Commissioner for the purpose of these regulations; or

(e) the systematic chemical name using the English system of nomenclature;

"Child" means a person under the age of 12 years;

"Direction" means regular and frequent supervision but does not necessarily imply continuous personal supervision;

"Dispense" in relation to a medicine or a poison means supplying the medicine or poison on and in accordance with a prescription duly given by a medical practitioner, a dentist or a veterinary surgeon;

"Experienced Person" means a person who for at least 5 years has been employed in the manufacture, handling or selling of poisons;

"Fourth Schedule drug" means any substance included in the Fourth Schedule to the Act;

"Internal Use" means a substance which is given parenterally, or orally, or a substance which is administered by way of a body orifice for the purpose of absorption and the production of a systemic effect;

"Manufacture" includes the processes of packing and repacking, refining manipulating and mixing any poison or hazardous substance (including such substance in the raw state);

"New Drug" has the same meaning as that term has in and for the purposes of section 37 of the Act;

"Permit" means a permit granted pursuant to the Act;

"Personal Supervision" means close and continuous control requiring the actual presence of the person exercising the supervision;

"Poisons Cupboard" means a substantially made cupboard provided with an effective locking device, and except in the case of a cupboard used for the storage of an Eighth Schedule poison having the word "Poison" conspicuously painted on the outside of the cupboard;

"Qualified Person" means—

(a) a medical practitioner, pharmaceutical chemist, dentist, veterinary surgeon;

(b) a person who is the holder of a degree approved by the Permanent Head, conferred by a University of the British Commonwealth;

(c) a person who is eligible to be—

(i) a Fellow or Associate of the Royal Australian Chemical Institute; or

(ii) a Fellow, Associate or Licentiate of the Royal Institute of Chemistry; or

(d) any other person approved of by the Permanent Head;

"Quarter" means any one of the three-monthly periods of any year ending on 31 March, 30 June, 30 September or 31 December;

"Sale" includes exposing or offering for sale or having in possession for sale, whether by wholesale or retail, and also delivery with or without consideration, in any shop or store or premises appurtenant thereto by the keeper thereof or by his servant or agent; and the verb "to sell" has a corresponding meaning;

"Supply" includes "distribute" and "sell" but the administration to a patient of any substance specified in any of the Schedules to the Act by a medical practitioner or dentist, or by a nurse when acting under the direction of a medical practitioner, or the administration of any substance specified in any of the Schedules to the Act to an animal under the direct personal supervision or under the direction of a veterinary surgeon, shall not be deemed to be supplying within the meaning of these regulations;

"the Act"means the *Poisons Act 1964*.

[*Regulation 2 amended in Gazettes 23 September 1983 p. 3803; 29 June 1984 p. 1784; 28 February 1986 p. 618; 5 December 1986 p. 4467.*]

Licences and Permits

3. A licence to procure, manufacture and supply by wholesale dealing poisons (other than drugs of addiction) shall authorize the licensee to procure, manufacture and supply (according to the endorsement thereon) by wholesale dealing such substances as are specified in the licence, and shall be in the Form 1 in Appendix A.

4. (1) A licence to procure, manufacture and supply by wholesale dealing drugs of addiction shall authorize the licensee to procure, manufacture, and supply by wholesale dealing drugs of addiction on or from the premises described in the licence, and shall be in the Form 2 in Appendix A.

(2) In addition to any other conditions required by these regulations the licence shall be subject to the following conditions:—

(a) The manufacture shall be carried out—
  (i) by a qualified person whose name appears on the licence; or
  (ii) by an experienced person acting under the personal supervision of a qualified person whose name appears on the licence;

(b) the supply shall be carried out—
  (i) by a qualified person whose name appears on the licence; or
  (ii) by an experienced person whose name appears on the licence;

but should the person whose name appears on the licence cease employment or be unable to exercise the necessary supervision, the Permanent Head may authorize, in writing, some other person having the required qualification to act in his stead.

[*Regulation 4 amended in Gazette 29 June 1984 p. 1784.*]

Pharmaceutical Chemist's Licence to Sell Poisons

5. A pharmaceutical chemist shall not sell or supply any poison except at or from a pharmacy registered under the *Pharmacy Act 1964*, and described in the licence issued under these regulations as provided in Form 3 in Appendix A.

Retailer's Licence to Sell Poisons Specified in the Sixth Schedule to the Act

6. This licence shall authorize the licensee to procure, and to sell by retail, the poisons specified in the Sixth Schedule to the Act, at the premises described in the licence, and shall be in the Form 4 in Appendix A.

Retailer's Licence to Sell Poisons Specified in the Second or Sixth Schedules to the Act

[*Heading amended in Gazette 23 May 1986 p. 1716.*]

7. This licence shall authorize the licensee to procure, and to sell by retail, poisons specified in either the Second or Sixth Schedules to the Act at the premises described in the licence, and shall be in the Form 5 in Appendix A.

[*Regulation 7 amended in Gazette 23 May 1986 p. 1716.*]

Retailer's Licence to Sell Poisons Specified in the Seventh Schedule to the Act

8. This licence shall authorize the licensee to procure, and to sell by retail, poisons specified in the Seventh Schedule to the Act at the premises described in the licence, and shall be in the Form 6 in Appendix A.

Poisons Permit (Distribution of Samples)

8A. (1) This permit shall, subject to the succeeding provisions of this regulation, authorize the holder to procure from any manufacturer or wholesale dealer specified therein and to supply to certain persons, samples of poisons specified in the First, Second, Third or Fourth Schedules to the Act and the permit shall be in the Form 6B in Appendix A.

(2) A permit under this regulation may be granted only to a person who is—

(a) a representative of a person—

(i) licensed to manufacture poisons or to supply poisons by wholesale dealing; or

(ii) licensed under the provisions of the laws of any other State or territory of the Commonwealth to manufacture poisons or to supply poisons by wholesale dealing;

(b) not less than 21 years of age; and

(c) of good character,

and the holder of a permit shall, for the purposes of these regulations be known as a detailer.

(3) A permit under this regulation shall contain the name and address of the detailer and the name of each manufacturer or wholesale dealer whom he represents.

(4) Where a detailer ceases to represent a manufacturer or wholesale dealer named in his permit—

(a) the permit shall thereupon cease to authorize the detailer to procure samples from that manufacturer or wholesale dealer or to supply to any person samples procured at any time from that manufacturer or wholesale dealer;

(b) the detailer shall return to the manufacturer or wholesale dealer any samples that were procured from the manufacturer or wholesale dealer and that are still in the possession or control of the detailer; and

(c) within 7 days of ceasing to represent the manufacturer or wholesale dealer, the detailer shall advise the Permanent Head in writing of the fact and deliver up therewith his permit to the Permanent Head, and the Permanent Head shall delete from the permit the name of the manufacturer or wholesale dealer or shall cancel the permit, as the case requires.

(5) A detailer shall not supply a sample to any person who is not—

(a) a medical practitioner;

(b) a veterinary surgeon;

(c) a dentist; or

(d) a pharmacist.

(6) A detailer shall not procure, carry or supply a sample that is larger than is required to provide 7 days of therapeutic treatment, according to the directions for maximum dosage supplied with the sample, except for the purpose of satisfying a prior request for a larger sample that has been made in writing by the person seeking to be supplied with such a sample to the manufacturer or wholesale dealer whom the detailer represents.

(7) The provisions of subregulation (6) do not apply to a sample of a proprietary preparation where—

(a) that sample is of a size not exceeding the smallest size manufactured for sale of that proprietary preparation; and

(b) the Permanent Head on the recommendation of the Poisons Advisory Committee has declared such a sample to be a sample to which subregulation (6) does not apply, notwithstanding that the sample may be larger than is required to provide 7 days of therapeutic treatment.

(8) A detailer shall not carry more than—

(a) 25 samples of any single proprietary preparation; or

(b) samples of more than 5 different proprietary preparations,

in a vehicle at any one time.

(9) Where the proper storage of a poison requires that poison to be stored under special conditions or at specific temperatures, a detailer shall not store or transport that poison except in a manner which maintains those conditions or temperatures.

(10) Subject to subregulation (11), a detailer shall not cause or permit samples in his possession or control to be stored other than—

(a) on the premises of the manufacturer or wholesale dealer whom he represents; or

(b) at his address as specified in his permit.

(11) A detailer may keep samples in a vehicle while he is actually using that vehicle in the course of his business, but at no other time.

(12) Where pursuant to this regulation samples are stored at an address specified in a detailer's permit which is not a wholesaler's premises, the detailer shall cause those samples to be stored in a locked cupboard or locked refrigerator and a detailer shall not cause or permit—

(a) more than 100 samples of any single proprietary preparation; or

(b) samples of more than 5 different proprietary preparations,

to be kept at that address at any one time.

(13) A detailer shall not supply a sample unless—

(a) he has received a signed request from a person to whom he is authorized in accordance with subregulation (5) to supply the sample; and

(b) immediately upon supplying the sample, he signs and dates the request form to certify that the sample has been delivered.

(14) A detailer shall keep a record of every sample received or supplied by him and shall preserve all records so kept together with consignment notes, invoices, advice notes and request forms relating thereto, for not less than 2 years.

(15) Upon receiving a written request from the Permanent Head, a detailer shall submit all records of samples received and delivered and shall make an account of those samples to the Permanent Head or a person authorized in accordance with section 54 of the Act.

(16) For the purposes of this regulation—

"proprietary preparation" means one or more forms of a poison intended for therapeutic use boxed or wrapped in a single sample package;

"sample" means a sample package containing a poison intended for therapeutic use specified in the First, Second, Third or Fourth Schedule.

[*Regulation 8A inserted in Gazette 22 September 1969 pp. 2874-76; amended in Gazette 29 June 1984 p. 1784.*]

Poisons Permit (Industrial)

9. This permit shall authorize the holder to purchase from a manufacturer or wholesale dealer such poisons as are specified in the permit, which shall be in the Form 7 in Appendix A.

Poisons Permit (Educational, Advisory or Research)

10. This permit shall authorize the holder to purchase from a manufacturer or wholesale dealer such poisons as are specified in the permit, which shall be in the Form 8 in Appendix A.

Poisons Permit (Departmental and Hospital)

10A. (1) This permit shall authorize the holder to purchase from a manufacturer or wholesale dealer such poisons as are specified in the permit which shall be in the Form 13 in Appendix A and shall not, except in the case of a permit held by a public hospital, authorize the sale of any poison obtained by the permit holder under the authority of the permit.

(2) This permit may be granted only to—

(a) a Department or instrumentality of the State or of the Commonwealth; and

(b) a public hospital within the meaning of the *Hospitals Act 1927*.

[*Regulation 10A inserted in Gazette 14 June 1967 p. 1582*]

Licence to Hawk, Peddle or Distribute Poisons

11. This licence issued under section 48 of the Act shall authorize the licensee to sell or distribute in the areas specified in the licence such poisons as are included in the licence, subject to the conditions, limitations and restrictions specified therein, and such licence shall be in the Form 9 in Appendix A.

Application for Licences or Permits

12. (1) A person desirous of obtaining a licence or permit pursuant to these regulations shall lodge with the Permanent Head an application in or to the effect of such of the Forms 1A to 13A in Appendix A as is appropriate in the particular case, together with the appropriate fee prescribed in Appendix G.

(2) Where the applicant for a licence or permit applies on behalf of a corporate body or firm the application shall contain the name of a natural person who, in respect of the premises named in the application, shall be responsible for carrying out the provisions of the Act and these regulations.

[*Regulation 12 amended in Gazette 5 October 1979 p. 3085; 6 November 1981 p. 4527; 29 June 1984 p. 1784.*]

Licences and Permits—General Conditions

13. Every licence or permit issued pursuant to these regulations shall be subject to these regulations and the conditions, limitations and restrictions set out in the licence or permit.

14. Every licence or permit issued pursuant to these regulations shall be valid until 30 June next following the day of issue, unless sooner cancelled, suspended or revoked, and may thereafter be renewed annually at the discretion of the Permanent Head on payment of the prescribed fee (if any).

[*Regulation 14 amended in Gazette 29 June 1984 p. 1784.*]

15. A licence or permit shall not be issued to any person under the age of 21 years, unless he is a qualified person approved by the Permanent Head.

[*Regulation 15 amended in Gazette 29 June 1984 p. 1784.*]

16. A sale of any poison shall not be made by any person other than the licensee or a person, not less than 16 years of age, acting on his behalf.

[*Regulation 16 amended in Gazette 21 November 1986 p. 4270.*]

17. A licence or permit is not transferable from one person to another.

Provided that—

(a) a licence or permit held in the name of a person on behalf of a firm or corporate body may, on endorsement by the Permanent Head, be transferred into the name of another person on behalf of the firm or corporate body;

(b) the holder of a licence or permit who ceases to carry on or conduct the business or practice to which the licence or permit relates shall within 14 days surrender such licence or permit to the Permanent Head.

[*Regulation 17 amended in Gazette 29 June 1984 p. 1784.*]

18. The holder of a licence shall keep such licence displayed in a conspicuous place within the premises specified in the licence.

Containers

19. (1) Subregulations (2) to (9) do not apply in respect of a vessel containing a medicine made up ready for human internal use or for animal internal use.

(2) Except as provided by these regulations, the immediate container in which any poison or hazardous substance is stored, sold, supplied or transported—

(a) shall be impervious to its contents;

(b) shall not be capable of reacting with its contents;

(c) shall be of sufficient strength and capacity to withstand the ordinary risks of breakage and expansion during storage, handling or transport without leakage; and

(d) shall be securely closed and, except where it contains a preparation packed for use on one occasion only, shall be capable of being securely reclosed.

(3) An immediate container on which the name of any poison or hazardous substance is embossed or otherwise permanently marked shall not be used except to contain that poison or hazardous substance.

(4) Subject to the provisions of subregulation (6), a hazardous substance shall not be sold unless the vessel immediately containing it—

(a) is embossed with or has indelibly written thereon the words "Not to be used as a food container" or the words "Not to be taken"; and

(b) is readily distinguishable from any type of container in which food, wine or other beverage is ordinarily sold.

(5) Subject to the provisions of subregulation (6) and subregulation (7) no hazardous substance which is referred to in the Fifth Schedule to the Act under any of the following descriptions,

HYDROCARBONS, LIQUID.

KEROSENE.

METHYLATED SPIRIT.

MINERAL TURPENTINE.

OIL OF TURPENTINE.

PETROL.

WHITE SPIRIT,

shall be sold unless the vessel immediately containing it complies with the requirements of subregulation (8) or subregulation (9).

(6) A vessel containing a poison or hazardous substance made up as a medicament for eye, ear or nose in the form of drops or a spray and which contains 15 millilitres or less of medicament is not required to comply with the requirements of subregulation (4), (5) or (9).

(7) A vessel containing a poison or hazardous substance made up as a medicament for the eye in the form of drops—

(a) must be capable of being sterilized;

(b) must have a locking cap secured with a screw or bayonet type fitting and adapted for the proper delivery of drops;

(c) must comply with the requirements of subregulation (9) (b) where the vessel has a capacity exceeding 15 millilitres;

(d) is not required to be of any particular colour.

(8) A poison shall not be sold in any immediate container having a capacity exceeding 2 litres unless the word "POISON" appears in letters which are—

(a) not less than 12 millimetres in height; or

(b) of a height which is not less than one-thirty second of the depth or width of the container,

whichever measure gives the greatest dimension, and is embossed, or indelibly written in colour contrast to the ground colour, on the side of that container.

(9) Subject to subregulation (6) and subregulation (7), a poison shall not be sold in any immediate container having a capacity of 2 litres or less unless that container complies with the following conditions—

(a) A bottle or jar shall have the word "POISON" or the words "not to be taken" appearing in raised lettering on the outer surface.

(b) A bottle or jar shall be provided with prominent vertical ribs or grooves, or prominent points or stars, which shall be of sufficient number to render the vessel distinguishable by sight and touch—

(i) from bottles or jars ordinarily used as containers for any food, drink or condiment;

(ii) from a vessel ordinarily used as a container for medicine for internal use; and

(iii) as a vessel reserved to contain poison.

(c) A bottle or jar shall be provided with a panel, or panels free from ribs, grooves points or stars of sufficient area for the purposes of labelling.

(d) A bottle or jar made of glass shall be colourless or brown in colour.

(e) A bottle or jar made of plastic shall comply with the Australian Standard Specification for Plastic Containers for Poisonous Substances as published by the Standards Association of Australia.

(10) A paper or plastic bag or envelope, or a cardboard box shall not be used as a container for a Second, Third, Fourth or Eighth Schedule poison whether dispensed or not, unless the poison is also presented to the purchaser in foil or in individually sealed, measured amounts, commonly described as strip packaging, or unless the container is approved by the Permanent Head.

(11) A paper bag shall not be used as the sole container of any poison or hazardous substance unless it has been approved by the Permanent Head for that purpose.

[*Regulation 19 substituted in Gazette 26 May 1971 pp. 1771-73; amended in Gazettes 3 May 1974 p. 1434; 29 June 1984 p. 1784; 5 December 1986 p. 4467.*]

19A. A person shall not sell any food, drink, or condiment, or any drug or medicine for internal use, in a container—

(a) of a description which is not readily distinguishable by sight and touch from a container in which a poison or hazardous substance intended for external use may be sold; or

(b) of a like description to that prescribed for a container in which a poison or hazardous substance intended for external use may be sold.

[*Regulation 19A inserted in Gazette 26 May 1971 p. 1773.*]

Labels

20. (1) Except as provided by these regulations, a person shall not sell, supply or distribute a poison or hazardous substance unless the container immediately containing the poison or hazardous substance bears, or has securely affixed to it a label bearing—

(a) the name and address of the manufacturer, wholesaler or retailer;

(b) the approved name of the poison or hazardous substance;

(c) the proportion or percentage of the poison or hazardous substance in relation to the whole of the contents of the container;

(d) where the poison or hazardous substance, not being a poison specified in the Fourth Schedule to the Act or the Eighth Schedule to the Act, is made up for a specific purpose, directions for the use of that poison or hazardous substance;

(e) where a poison or hazardous substance is set out in Appendix C, the first aid measures to be followed if poisoning occurs;

(f) where a poison or hazardous substance is set out in Appendix D, the relevant warning statement set out in that Appendix;

(g) where a poison is set out in Appendix E, the warning statement and first aid statement set out in respect of that poison in that Appendix; and

(h) the particulars specified in respect of that poison or hazardous substance in subregulation (1a).

(1a) The particulars referred to in subregulation (1) (h)—

(a) in relation to a poison specified in the First Schedule to the Act, are—
  (i) POISON;
  (ii) NOT TO BE TAKEN; and
  (iii) KEEP OUT OF REACH OF CHILDREN;

(b) in relation to a poison specified in the Second Schedule to the Act and prepared and packaged for internal use, are—
  (i) CAUTION;
  (ii) USE STRICTLY AS DIRECTED; and
  (iii) KEEP OUT OF REACH OF CHILDREN;

(c) in relation to a poison specified in the Second Schedule to the Act, other than a poison prepared and packaged for internal use, are—
  (i) POISON;
  (ii) NOT TO BE TAKEN; and
  (iii) KEEP OUT OF REACH OF CHILDREN;

(d) in relation to a poison specified in the Third Schedule to the Act, are—
  (i) CAUTION;
  (ii) USE STRICTLY AS DIRECTED; and
  (iii) KEEP OUT OF REACH OF CHILDREN;

(e) in relation to a poison specified in the Fourth Schedule to the Act, are—
  (i) CAUTION;
  (ii) SUPPLY WITHOUT PRESCRIPTION ILLEGAL; and
  (iii) KEEP OUT OF REACH OF CHILDREN;

(f) in relation to hazardous substance specified in the Fifth Schedule to the Act, are—
  (i) WARNING; and
  (ii) KEEP OUT OF REACH OF CHILDREN;

(g) in relation to a poison specified in the Sixth Schedule to the Act, and prepared and packaged for internal use in animals, are—
  (i) CAUTION;
  (ii) USE STRICTLY AS DIRECTED; and
  (iii) KEEP OUT OF REACH OF CHILDREN;

(h) in relation to a poison specified in the Sixth Schedule to the Act, other than a poison prepared and packaged for internal use in animals, are—
  (i) POISON;
  (ii) NOT TO BE TAKEN;
  (iii) KEEP OUT OF REACH OF CHILDREN; and
  (iv) READ SAFETY DIRECTIONS BEFORE OPENING.

(i) in relation to a poison specified in the Seventh Schedule to the Act, are—
  (i) POISON;
  (ii) KEEP OUT OF REACH OF CHILDREN; and
  (iii) READ SAFETY DIRECTIONS BEFORE OPENING;
  and

(j) in relation to a poison specified in the Eighth Schedule to the Act, are—
  (i) CAUTION;
  (ii) KEEP OUT OF REACH OF CHILDREN; and
  (iii) SUPPLY WITHOUT PRESCRIPTION OR POSSESSION WITHOUT AUTHORITY IS ILLEGAL.

(2) The provisions of subregulation (1) shall not apply—

(a) to persons licensed pursuant to section 24 (1) (a) of the Act; and

(b) with respect to the supply by a medical practitioner of any poison or substance containing a poison or hazardous substance for the purposes of therapeutic treatment to a patient for a period not exceeding 5 days at any one time.

[*Regulation 20 amended in Gazettes 20 October 1978 p. 3760; 28 February 1986 pp. 616-17.*]

21. Notwithstanding the provisions of regulation 20 a medicine containing any poison dispensed or supplied—

(a) by a pharmaceutical chemist for human internal use shall comply with that regulation if it is labelled according to the instructions given on the prescription together with the identifying number of the prescription and the name and address of the pharmacy at which it is supplied;

(b) by a pharmaceutical chemist for external therapeutic use shall comply with that regulation if it is labelled with the words "not to be taken" together with the directions, prescription identification number and the name and address of the pharmacy at which it is supplied;

(c) for use in or on an animal shall comply with that regulation if it is labelled—
  (i) with the owner's surname and species of animal;
  (ii) with instructions for use;
  (iii) with the date of dispensing or the identifying number for that prescription or supply; and
  (iv) with the name and address of the pharmacy or veterinary practice at which it is supplied.

[*Regulation 21 substituted in Gazette 5 December 1986 p. 4467.*]

21A. (1) A person, whether a pharmaceutical chemist or otherwise, shall not sell, supply, distribute or dispense a poison set out in Appendix K unless the container immediately containing the poison bears, or has securely affixed to it a label bearing either of the following statements—

"This medicine may cause drowsiness. If affected do not drive a motor vehicle or operate machinery. Avoid alcohol."; or

"This medicine may cause drowsiness and may increase the effects of alcohol. If affected do not drive a motor vehicle or operate machinery.".

(2) Subregulation (1) shall not apply to—

(a) a person licensed pursuant to section 24 (1) (a) of the Act; and

(b) the supply by a medical practitioner of any poison or substance containing a poison for the purposes of therapeutic treatment to a patient while that patient is hospitalized.

[*Regulation 21A inserted in Gazette 11 July 1986 p. 2339.*]

22. Wherever the word "Poison" or "Caution" is required to be shown on a label it shall appear in red on a white background and be surrounded by a red frame. Such word shall form the first line of the principal label and no other word or words shall appear on the same line. The word "Poison" or "Caution" as the case may be shall be in bold face sans serif capital letters of a size not less than half the size of the largest lettering on the label and in any case not less than 6 points face measurement.

23. A label shall not be attached or affixed to any bottle containing any poison or hazardous substance in such a manner that the embossed points, ridges, flutes, stars, name of the article or the prescribed words blown thereon are covered or obliterated: Provided that the label may cover the front panel of the bottle and extend around the adjacent sides, if the embossed matter on the back panel of the bottle and the name of the article and the prescribed words blown thereon are not covered or obliterated.

24. Any preparation containing any poison or hazardous substance which is supplied for veterinary use, whether in pursuance of a prescription or otherwise, shall be labelled "For veterinary use only" or "For animal treatment only".

Containers and Labels—General

25. The Permanent Head may approve, in writing, a container or label which does not comply with these regulations if, having regard to the nature of the poison and the purpose for which it is to be used, it is unlikely that the interests of safety will be adversely affected by the use of such container or label.

[*Regulation 25 amended in Gazette 29 June 1984 p. 1784.*]

26. The Permanent Head may, in the interest of safety, suspend or prohibit the use of any form of container or label for the packing or labelling of any poison.

[*Regulation 26 amended in Gazette 29 June 1984 p. 1784.*]

27. Wherever it is required that the words—

"Keep out of reach of children" or

"First aid measures" or

the approved name of the poison or poisons

shall appear on a label, such words or particulars shall be shown—

(a) in bold face san serif capitals of not less than 6 point face measurement; and

(b) in such colour or colours as to afford a distinct contrast to the background colour.

27AA. Whenever it is required that a warning statement referred to in regulation 21A shall appear on a container or a label, the words in that statement shall be not less than 4 point face measurement.

[*Regulation 27AA inserted in Gazette 11 July 1986 p. 2339.*]

27A. Every preparation containing a poison dispensed by count by a pharmaceutical chemist on the prescription of a medical practitioner, dentist or veterinary surgeon or supplied by a veterinary surgeon to the custodian of an animal, shall be labelled—

(a) with the name and strength or amount of each poison in the preparations; or

(b) with the trade name and strength of the preparation, unless the trade name also uniquely identifies the strength, in which case only the trade name need be given.

[*Regulation 27A substituted in Gazette 5 December 1986 p. 4467.]*

Calculation or Percentages

28. Where required, percentages may be expressed in units other than those in section 51 of the Act.

Storage

29. [*(1), (2) repealed*]

(3) Hydrocyanic acid or cyanides, in bulk, shall, subject to regulation 29B, be stored on premises which are specified in a licence or permit—

(a) in a shipping container which is raised off the ground;

(b) raised off the ground on a well drained site—

(i) surrounded by a security fence with barbed wire and a security gate with barbed extension both being not less than 2 metres in height; and

(ii) where less than 5 kilometres from a town, roofed so as to protect the hydrocyanic acid or cyanides from exposure to rain; or

(c) being a warehouse ventilated in accordance with regulation 7 or 8 of the Factories (Health and Safety) Regulations; or

(d) being premises approved by the Permanent Head.

[*Regulation 29 substituted in Gazette 28 February 1986 p. 618; amended in Gazette, 1 August 1986 p. 2739.*]

29A. Hydrocyanic acid or cyanides stored in accordance with regulation 29(3)—

(a) shall be adequately and properly ventilated to prevent the accumulation of hydrogen cyanide gas;

(b) shall be secured in a lockable container or enclosure which is kept locked when the premises are unattended; and

(c) shall not be stored in the same premises as the liquid form of any acid other than hydrocyanic acid, and

no other items shall be stored in the same premises as the hydrocyanic acid or cyanides unless they are physically divided and easily and readily distinguishable from the hydrocyanic acid or cyanides.

[*Regulation 29A inserted in Gazette 28 February 1986 p. 618.*]

29B. Where hydrocyanic acid or cyanide is in the process of being transported it may be stored on other than licensed premises for a period of not more than 24 hours.

[*Regulation 29B inserted in Gazette 28 February 1986 p. 618.*]

30. Any person having a hazardous substance or a poison, other than those specified in regulations 29 or 56, in or on any premises for the purpose of sale or use in his profession, business, trade or industry shall keep that poison or hazardous substance in such a manner as to preclude contamination of any food, drink or condiment by the poison or hazardous substance; and to preclude access to the poison or hazardous substance by children.

[*Regulation 30 amended in Gazette 1 August 1986 p. 2739.*]

Disposal of Poisons

31. A person shall not dispose of any poison in any place or manner likely to constitute a risk to the public.

Notification of Loss or Theft of Poison

32. Every person who loses any poison or from whom any poison is stolen shall immediately notify a member of the Police Force of such loss or theft.

Poison Not to be Sold to Persons Under 16 years

33. A person who sells or supplies any poison to any person who is apparently under 16 years of age commits an offence against these regulations, but this regulation does not apply to sales of pharmaceutical preparations or medicines by persons licensed to sell poisons by retail.

[*34., and Heading. Regulation 34 and heading thereto repealed in Gazette 23 May 1986 p. 1716.*]

[*34A. 34B., 34C., and Heading. Regulations 34A, 34B, 34C and the heading before regulation 34A repealed in Gazette 23 May 1986 p. 1716.*]

New Drugs

34D. (1) Before any new drug is first offered for sale to the public the manufacturer, importer or distributor, as the case may require, shall make application to the Permanent Head to classify the new drug by determining the Schedule (if any) to the Act in which it is to be included.

(2) An application made under this regulation shall be in Form No. 10 of Appendix A.

[*Regulation 34D was previously numbered regulation 35 and while it was numbered 35 it was amended in Gazette 29 June 1984 p. 1784; renumbered as 34D in Gazette 8 February 1985 p. 521.*]

Restrictions on Retail Sale of Second and Third Schedule Poisons

[*Heading to regulation 35A deleted and regulation 35 and heading substituted in Gazette 8 February 1985 p. 521.*]

35. A substance referred to in the Second Schedule shall not be stored for retail sale in any area or in any manner that allows physical access to that substance by any person other than—

(a) the owner of the business carried on; or

(b) a person employed,

on the premises where it is stored.

[*Regulation 35: See note under heading.*]

35A. (1) A pharmaceutical chemist shall not sell any of the following substances namely,

Butyl Nitrite;

Amyl Nitrite;

Substances containing butyl nitrite or amyl nitrite;

Chloral hydrate when included in the Third Schedule;

or

Substances containing chloral hydrate when contained in the Third Schedule,

to any person who is apparently under the age of 21 years.

(1a) A substance referred to in the Third Schedule shall not be sold or supplied by retail except under the personal supervision of a pharmaceutical chemist.

(1b) A pharmaceutical chemist shall not store in any part of the retail area of premises any of the substances referred to in Appendix J.

(1c) A substance referred to in Appendix J shall only be sold or supplied by direct, personal sale by a pharmaceutical chemist or by a graduate trainee in pharmacy under the personal supervision of a pharmaceutical chemist.

(2) Before a substance referred to in Appendix J is delivered to a purchaser on a sale by retail, the pharmaceutical chemist or graduate trainee in pharmacy making the sale shall—

(a) record, in ink, in the prescription book referred to in regulation 36 (3) (c), the following particulars—

(i) the date of sale;

(ii) the name and address of the purchaser and, where the person for whom the substance is intended is not the purchaser, the name and address of the person for whom the substance is intended; and

(iii) the name and quantity of the substance supplied,

and the entry in the prescription book shall be given a unique identification number or letter;

(b) label the product with—

(i) the name and address of the pharmacy; and

(ii) the unique identifying number or letter allocated in accordance with paragraph (a).

(3) The prescription book referred to in this regulation shall be available for inspection upon request by any inspector appointed under the Health Act 1911 or to a person authorized in that behalf by the Minister.

(4) The seller shall retain the records required to be made under this regulation for a period of at least 2 years.

[*Regulation 35A inserted in Gazette 28 November 1968 p. 3458; amended in Gazette 29 August 1980 p. 3028; 20 September 1985 p. 3743.*]

35B. A substance referred to in the Third Schedule shall not be stored in a pharmacy in any area or in any manner that allows physical access to that substance by any person other than a person who is a member of the staff of the pharmacy.

[*Regulation 35B inserted in Gazette 29 August 1980 p. 3028.*]

35C. A substance referred to in the Third Schedule shall not be advertised except in a *bona fide* professional or trade journal or other publication intended for circulation only within the medical, veterinary, dental or pharmaceutical professions or the wholesale and manufacturing drug trade.

[*Regulation 35C inserted in Gazette 23 September 1983 p. 3803.*]

Advertising, storage and display of Fourth Schedule substances

35D. A substance referred to in the Fourth Schedule—

(a) shall not be advertised except in a publication that is normally sold or intended for sale or circulation only among—

(i) persons of the kind referred to in section 23 (2) of the Act; or

(ii) persons who are holders of licences granted under section 24 (1) (a), (b) or (c) of the Act;

and

(b) shall not be held, stored or exposed or offered for sale in any portion of a pharmacy to which persons other than members of the staff of the pharmacy have access.

[*Regulation 35D inserted in Gazette 23 January 1987 p. 187.*]

Supply of Fourth Schedule Drugs

36. (1) Subject to the Act and these regulations a Fourth Schedule drug shall not be sold or supplied to any person unless—

(a) he is authorized under regulation 40 to procure the drug and, if he is a person referred to in regulation 40 (1) (b) or 40 (1) (g), he provides satisfactory evidence that he is so authorized;

(b) he is the holder of a prescription written by a medical practitioner, dentist or veterinary surgeon, prescribing the drug according to the requirements of these regulations; or

(c) the pharmaceutical chemist supplying the drug is satisfied that the person in respect of whom the drug is to be sold or supplied is under medical treatment with the drug and requires emergency treatment with the drug and supplies only a maximum of 3 days medication of the drug or where the drug is supplied to the pharmacist in prepacked individual packs then only one individual standard pack.

(2) A medical practitioner, pharmaceutical chemist, or veterinary surgeon or an assistant under the direct personal supervision of a medical practitioner, pharmaceutical chemist, or veterinary surgeon shall be the only person who shall dispense a Fourth Schedule drug.

(3) The following conditions shall be observed by persons dispensing such prescriptions:—

(a) The prescription shall not be dispensed more than the maximum number of times indicated thereon, and on each occasion upon which it is dispensed the prescription shall be stamped or marked to show clearly the date upon which it is dispensed and the name and address of the pharmacy at which it is dispensed.

(b) The person who dispenses a prescription which does not clearly indicate the maximum number of times it is to be dispensed, or which has reached the last occasion upon which it may be dispensed according to the maximum indicated thereon, shall write in ink, stamp or mark in legible letters across such prescription the word "cancelled".

(c) (i) For the purpose of this paragraph any card system photographic system, or other reference system, of recording the details of prescriptions required by this paragraph and which is approved by the Permanent Head shall be deemed to be the Prescription Book;

(ii) before the drug is handed to the purchaser the following details from the prescription shall be entered into the prescription book—

the name and quantity of the drug, the direction for use (if applicable), the date of issue of the prescription, the name and address of the patient, the name and address, or the name and identifying initials, of the prescriber, the date of dispensing the prescription, and the entry shall be given an identifying letter or number or combination of letter and number;

(iii) in the event of the dispensing of a repeated prescription an annotation of this fact showing the date of the repeat on the original entry in the Prescription Book shall be sufficient compliance with this regulation;

(iv) the label on the bottle or package containing the drug shall be marked with the identifying letter or number of the prescription as appearing in the Prescription Book; and

(v) the Prescription Book shall be kept at the place at which the Fourth Schedule drug was dispensed and shall be produced on demand to any person authorized in that behalf under the Act or these regulations.

(d) A prescription marked "cancelled" or that is more than 6 months old shall not be dispensed.

(e) A prescription which is illegible or defaced or which appears to be for the purpose of enabling some unauthorized person to obtain a Fourth Schedule drug, or which does not appear to be genuine, shall not be dispensed.

(f) A pharmaceutical chemist to whom a prescription referred to in paragraph (e) is presented shall retain the prescription and forthwith inform the Permanent Head of the relevant circumstances and the reasons for his refusal to dispense the prescription.

(g) A pharmaceutical chemist may dispense a prescription not bearing the address of the patient or the prescriber or both if he keeps a record of the prescription and notifies the Permanent Head concerning it.

[*Regulation 36 amended in Gazettes 19 February 1971 pp. 518-19; 29 August 1980 p. 3028; 29 June 1984 p. 1784; 5 July 1985 p. 2392.*]

37. A prescription for a Fourth Schedule drug shall comply with the following conditions:—

(a) it shall show in a clearly legible and indelible manner—

(i) the name and address of prescriber;

(ii) the address of the patient;

(b) there shall be written in ink in the prescriber's own handwriting—

(i) the name of the patient;

(ii) the name and quantity of the substance;

(iii) direction for use, if necessary;

(iv) the date on which it is written;

(v) the maximum number of times it may be repeated, if any, and (where applicable) the intervals at which it may be repeated; and

(vi) the signature of the prescriber;

(ba) notwithstanding paragraph (b), a prescription shall not be required to be in the prescriber's own handwriting where that prescription is processed on a computer the use of which is approved, in writing, by the Poisons Advisory Committee and the Executive Director, Public Health, for that purpose, but subparagraphs (i) to (vi) of paragraph (b) still apply to that prescription.

(c) a prescription written by a dentist shall be for dental purposes only and shall be marked as such and a prescription written by a veterinary surgeon shall be for veterinary use only and shall be marked "For veterinary use only" or "For animal treatment only";

(d) if a prescription contains an unusual dose the prescriber shall indicate that such a dose is intended by underlining that part of the prescription and initialling the same in margin;

(e) a prescription shall not bear the impression of a rubber stamp or other such contrivance in lieu of the written signature of the medical practitioner, dentist or veterinary surgeon by whom it has been written; and

(f) a prescription shall not be written in cipher.

[*Regulation 37 substituted in Gazette 19 February 1971 p. 519; amended in Gazette 21 November 1986 p. 4269; 5 December 1986 p. 4467.*]

Dispensing Fourth Schedule Drugs in Emergency Cases

38. Where a medical practitioner, dentist or veterinary surgeon in a case of emergency orally or by telephone or telegram directs the dispensing of a Fourth Schedule drug, he shall forthwith write a prescription complying with the conditions prescribed in regulation 37, mark such prescription to show that it has been given as a confirmation of instructions given by him orally or by telephone or telegram, and despatch such prescription within 24 hours to the person to whom the instructions were given.

Silver Sulphadiazine

38A. (1) Silver sulphadiazine or a substance containing silver sulphadiazine shall not be used except in a hospital for the purpose of the treatment of major burns or the treatment of conditions where full thickness skin loss has occurred.

(2) Notwithstanding any other provision of these regulations, silver sulphadiazine shall not be sold or supplied, except—

(a) for use in the treatment of major burns or the treatment of conditions where full thickness skin loss has occurred, to—

(i) a medical practitioner, for treatment of a patient at a hospital;

(ii) a matron of a hospital;

(iii) a patient in a hospital;

(iv) the Royal Flying Doctor Service for inclusion in the Royal Flying Doctor Service Kits for use only under the direction of a medical practitioner;

(b) to a pharmaceutical chemist for the purpose of sale or supply to a person referred to in paragraph (a);

(c) to a licensed wholesaler for the purpose of sale or supply to a person referred to in paragraph (a) (i), (a) (ii) or (b);

(d) to a licensed manufacturer for the purpose of sale or supply to a person referred to in paragraph (a) (i), (a) (ii), (b) or (c).

[*Regulation 38A and heading inserted in Gazette 24 December 1982 p. 4904; amended in Gazette 23 May 1986 p. 1716.*]

Buprenorphine

38B. Notwithstanding any other provision of these regulations, regulations 51C, 51D and 51E apply to the prescribing, use, sale and supply of buprenorphine as if references in those regulations to methadone were references to buprenorphine.

[*Regulation 38B inserted in Gazette 6 April 1984 p. 928; erratum in Gazette 13 April 1984 p. 1020.*]

Clomiphene and Cyclofenil

38C. Clomiphene or Cyclofenil or a substance containing clomiphene or cyclofenil and other substances specifically prepared to stimulate ovulation shall not be supplied except on the prescription or order of a qualified specialist practising in the field of gynaecology or obstetrics or both, or for the purpose of the conduct of medical or scientific research including veterinary trials under the direction of veterinary surgeons.

[*Regulation 38C inserted in Gazette 8 February 1985 p. 519; amended in Gazettes 31 May 1985 p. 1882; 23 May 1986 p. 1716.*]

Etretinate

38D. Etretinate or a substance containing etretinate shall not be supplied except on the prescription or order of specialist physicians or dermatologists and the supplier—

(a) shall, when supplying the patient or the agent of the patient with the substance, provide that person with a leaflet approved by the Executive Director setting out the hazards associated with the use of the substance; and

(b) shall ensure that the container in which the substance is supplied is labelled with a statement as follows—

"WARNING—CAUSES BIRTH DEFECTS".

[*Regulation 38D inserted in Gazette 8 February 1985 p. 519; amended in Gazettes 31 May 1985 p. 1882; 23 May 1986 p. 1716.*]

Prostaglandins

38E. Prostaglandins or substances containing prostaglandins, except where separately specified in the Fourth Schedule, shall not be supplied except on the prescription of a specialist medical practitioner practicing in general medicine, obstetrics and gynaecology, or of a veterinary surgeon, or unless approved in writing by the Permanent Head for research or other purposes.

[*Regulation 38E substituted in Gazette 5 December 1986 p. 4467.*]

Isotretinoin

38F. Isotretinoin or a substance containing isotretinoin shall not be supplied except on the prescription or order of a specialist physician or dermatologist and the supplier—

(a) shall, when supplying the substance to the patient or the agent of the patient, provide that person with a leaflet approved by the Executive Director setting out the hazards associated with the use of the substance; and

(b) shall ensure that the container in which the substance is supplied is labelled with a statement as follows—

"WARNING—CAUSES BIRTH DEFECTS".

[*Regulation 38F inserted in Gazette 31 May 1985 p. 1882; amended in Gazette 23 May 1986 p. 1716.*]

Thalidomide

38G. Thalidomide or substances containing thalidomide shall not be supplied except on the prescription or order of a leprologist for the treatment of erythema nodosum leprosum.

[*Regulation 38G inserted in Gazette 31 May 1985 p. 1882; amended in Gazette 23 May 1986 p. 1716.*]

Fourth Schedule Drugs for Veterinary Use

39. (1) Notwithstanding the provisions of regulation 36 a pharmaceutical chemist is authorized to supply for veterinary use of Fourth Schedule drug listed in Appendix "H" without a prescription where—

(a) the purchaser satisfies such pharmaceutical chemist that it is not reasonably practicable for him to obtain such a prescription;

(b) the name and address of the purchaser, date of supply, form and quantity of drug supplied, species of animal and number of animals to be treated, and a descriptive name of the disease for which the animals are to be treated, are entered in a register of poisons;

(c) the quantity of drugs supplied is not greater than is required to provide 72 hours of therapeutic treatment according to the directions for normal dosage with the drug, or in the case of a pre-packed proprietary brand the smallest size manufactured for sale of the proprietary brand which provides 72 hours treatment; and

(d) the pharmaceutical chemist provides adequate written instructions for the use of the drug.

(2) Any preparation for veterinary use containing a Fourth Schedule poison registered in Register 1 under the *Veterinary Preparations and Animal Feeding Stuffs Act 1976* and labelled in accordance with the requirements of that Act may be sold subject to such conditions, restrictions and limitations as are prescribed by that Act and its regulations, by a pharmaceutical chemist or by a person holding a permit as provided in Form 11, Appendix A.

[*Regulation 39 substituted in Gazette 26 August 1977 p. 2966.*]

39A. (1) Notwithstanding any other provision of these regulations a stockfeed manufacturer holding an appropriate permit under subregulation (3) may sell by retail to any person producing a written order therefor signed by a veterinary surgeon, a mixture of stockfeed with any Fourth Schedule drug being an antibiotic or sulphonamide, in such quantity and of such composition as is specified in the order.

(2) The signed order shall be cancelled by the stockfeed manufacturer and retained by him for not less than 2 years after the sale, and upon request shall be produced for inspection to an officer authorized in that behalf by the Permanent Head.

(3) A stockfeed manufacturer who wishes to sell by retail mixtures pursuant to subregulation (1) may apply to the Permanent Head for, and at the discretion of the Permanent Head be granted, a permit in Form 11AA in Appendix A, specifying the Fourth Schedule drugs that may be contained in such mixtures, and any limits as to the quantity or composition of such mixtures that may be sold.

(4) The provisions of section 23 of the Act do not apply to a sale by a stockfeed manufacturer pursuant to and in accordance with this regulation or to the preparation of a mixture of stockfeed for the purposes of such sale.

[*Regulation 39A inserted in Gazette 5 October 1979 p. 3085; amended in Gazette 29 June 1984 p. 1784.*]

Special Authority to Purchase Fourth Schedule Drugs

40. (1) Until in any particular case such authority is withdrawn—

(a) a medical practitioner;

(b) a pharmaceutical chemist;

(c) a dentist;

(d) a veterinary surgeon;

(e) an analyst appointed under the *Health Act 1911;*

(f) a matron of a hospital registered under the *Hospitals Act 1927;*

(g) any other person authorized in writing by the Permanent Head,

is authorized to procure a Fourth Schedule drug to the extent that it is required for the purpose of his profession or employment, but such authority does not entitle any person to have in his possession any Fourth Schedule drug other than in accordance with these regulations or in any quantity greater than is permitted by the Permanent Head.

(2) A person who wishes to use any Fourth Schedule drug being an antibiotic or sulphonamide for the preparation of mixtures for sale pursuant to regulation 39A, and who holds a permit under that regulation to sell such mixtures, is authorized to procure, use and be in possession of such Fourth Schedule drug for the preparation of such mixtures.

[*Regulation 40 amended in Gazettes 5 October 1979 p. 3085; 29 June 1984 p. 1784; 8 February 1985 p. 519.*]

Delivery of a Fourth Schedule Drug on Order

41. (1) A Fourth Schedule drug shall not be delivered to any person other than a person authorized by these regulations to purchase the drug or except on the authority of a written order signed by such authorized person, and the person supplying the drug shall satisfy himself that the authority is geniune.

(2) This regulation shall not be deemed to apply to medicines dispensed in pursuance of the foregoing regulations relating to the dispensing of Fourth Schedule drugs.

[*Regulation 41 amended in Gazette 9 February 1970 p. 370.*]

Sale of certain Sixth and Seventh Schedule poisons

41A. (1) A person who sells any poison included in the Seventh Schedule, or 2 or more kilograms of a poison included in the Sixth Schedule and listed in the Table to this regulation—

(a) to a *bona fide* primary producer;

(b) to a licensed pesticide operator;

(c) to a holder of an industrial poisons permit specifying that poison; or

(d) by retail,

shall, in addition to any conditions, restrictions and limitations imposed by notice under section 24 (5) or by licence issued in accordance with these regulations, keep a record of sale in accordance with this regulation.

(2) A person recording a sale for the purposes of subregulation (1) shall—

(a) record in ink in a register kept for that purpose particulars of—

(i) the date of sale;

(ii) the name, occupation and address of the purchaser;

(iii) the nature and quantity of the poison sold; and

(iv) the place and purpose of intended use,

and obtain the signature of the purchaser to the entry in the register; or

(b) keep a record in the form of an invoice showing particulars of—

(i) the date of sale;

(ii) the name and address of the purchaser;

(iii) the nature and quantity of the poison sold; and

(iv) the address to which the poison is to be delivered if that address differs from the address recorded under subparagraph (ii).

(3) A record of sale made for the purposes of this regulation shall be—

(a) made before delivery of the poison;

(b) kept for a period of at least 2 years; and

(c) produced for inspection on demand by a person appointed or authorized by or under the Act to inspect those records.

TABLE

Aldrin
Chlordane
Dieldrin
Heptachlor

[*Regulation 41A substituted in Gazette 23 May 1986 p. 1721; erratum in Gazette 30 May 1986 p. 1769.*]

Record of Third, Fourth and Seventh Schedule Poisons

41B. (1) Every person who holds a licence to procure, manufacture, or supply poisons referred to in the Third, Fourth, or Seventh Schedule by wholesale dealing shall, in relation to the supply of any such poison, keep an accurate record of—

(a) the day on which the poison was supplied;

(b) the quantity, form and strength of the poison supplied;

(c) the name and address of the person to whom it was supplied; and

(d) the reference number on the invoice or other document evidencing the supply,

and the record shall be made on the day of supply and shall be kept for not less than 2 years after that day.

(2) A person referred to in subregulation (1) shall send to the Executive Director particulars in writing of any of the information required to be recorded and kept by that person under subregulation (1)—

(a) within 7 days of being requested to do so where the information has been recorded within 2 months immediately before the request; and

(b) otherwise within 28 days of being requested to do so.

[*Regulation 41B inserted in Gazette 19 December 1986 pp. 4874-75.*]

DRUGS OF ADDICTION

Authority for Prescribed Persons to Procure and have Drugs of Addiction

42. (1) Until in any particular case such authority is withdrawn—

(a) a medical practitioner;

(b) a pharmaceutical chemist employed in dispensing medicines at any public hospital or at a pharmacy for which a licence is held under regulation 5;

[*(c) deleted*]

(d) a veterinary surgeon;

(e) an analyst registered under the *Health Act 1911;*

(f) a nurse employed in a public hospital (so far as the possession or use of such drug of addiction is required in connection with its administration to a patient under the instruction of a medical practitioner); and

(g) a person in possession of a permit granted by the Permanent Head under these regulations,

is, subject to these regulations, hereby authorized to procure and be in possession of any drug of addiction for the purpose of his profession or employment.

(2) A person to whom a prescription for a drug of addiction has been given is hereby authorized to procure and have possession of the drug of addiction to the extent specified in the prescription.

(3) The authority under this regulation to procure and be in the possession of any drug of addiction does not entitle the holder to procure or have in his possession any drug of addiction in any quantity greater than is permitted by the Permanent Head.

(4) Until in any particular case such authority is withdrawn a dentist is, subject to these regulations and for the purpose of his profession, hereby authorized to procure and be in possession of the following drugs of addiction in quantities not greater than those set out hereunder—

PETHIDINE, in a form prepared for injection with a total pethidine content of 600 milligrams

PAPAVERETUM, in tablet form, with a total papaveretum content of 240 milligrams

CODEINE PHOSPHATE, in tablet form, with a total codeine phosphate content of 900 milligrams

METHADONE, in tablet form, with a total methadone content of 240 milligrams

MORPHINE, in a form prepared for injection, with a total morphine content of 180 milligrams

OXYCODONE, in tablet form, with a total oxycodone content of 120 milligrams

PENTAZOCINE, in a form prepared for injection, with a total pentazocine content of 360 milligrams.

[*Regulation 42 amended in Gazette 9 February 1970 p. 370; 29 June 1984 p. 1784; 8 February 1985 p. 520.*]

Authority for Pharmacists to Retail, Compound and Dispense Drugs of Addiction

43. (1) Until in any particular case such authority is withdrawn, every pharmaceutical chemist holding a Pharmaceutical Chemist's licence to sell poisons under these regulations is hereby authorized, subject to the conditions, limitations and restrictions imposed by the Permanent Head, to procure and to manufacture at his registered premises in the ordinary course of his retail business any preparation, admixture, or extract of any drug of addiction, and to carry on at his registered premises the business of dispensing or compounding any drug of addiction, and also of retailing and supplying a drug of addiction, but only to persons licensed or authorized under these regulations to be in possession of or to procure the drug of addiction.

(2) The authority under this regulation does not in any way entitle the holder to procure, manufacture, sell, distribute, supply, or have in his possession any drug of addiction in any quantity greater than is permitted by the Permanent Head.

[*Regulation 43 amended in Gazette 29 June 1984 p. 1784.*]

Authority to procure, possess, etc. Drugs of Addiction and Specified Drugs may be revoked, etc.

[*Heading inserted in Gazette 9 February 1970 p. 370.*]

43A. The Permanent Head may by notice given to any such person as is referred to in subsection (2) of section 23 of the Act, revoke, in whole or in part, the authority conferred by that subsection on that person in relation to drugs of addiction and specified drugs.

[*Regulation 43A substituted in Gazette 29 August 1980 p. 3028; amended in Gazette 29 June 1984 p. 1784.*]

Register of Drugs of Addiction

44. (1) Any person authorized to manufacture, distribute, sell or possess any drug of addiction (other than a person having possession by the authority of a prescription from a medical practitioner or veterinary surgeon to the extent shown in the prescription) shall keep or cause to be kept a Register in the form or to the effect of Appendix B or in any other form required by the Commissioner in writing, and shall enter or cause to be entered in such Register accurate records of the drugs of addiction manufactured, procured, used, supplied or kept by him or on his behalf.

(2) The entries in such Register shall be written in ink on the day of the transaction, and such Register shall be kept on the premises at which the drugs of addiction are kept, manufactured, or disposed of; and where the holder of a licence or other authorized person has drugs of addiction on other premises, he shall keep, or cause to be kept, such Register on those premises also.

(3) All such Registers shall be at all times available for inspection by persons authorized by or under the Act or the regulations to inspect such Registers.

(4) Alterations, obliterations or cancellations shall not be made in any Register, but any mistake made in any entry may be corrected by a marginal or foot note, initialled and dated.

(5) Every person required by these regulations to keep a Register of Drugs of Addiction shall enter in the Register—

(a) the name, quantity and form of every drug of addiction received, manufactured, sold, used or otherwise disposed of;

(b) the date of each transaction;

(c) the name and address of the person or firm concerned in each transaction;

(ca) the name of the person who wrote the prescription or order;

(d) the balance remaining on hand after each transaction;

(e) in the case of a pharmaceutical chemist the identifying number of the prescription;

(f) in the case of a manufacturer or distributor an identifying number of the order or other authority on which the drug of addiction is supplied.

(6) A separate page of the Register shall be used for each drug of addiction recorded, so that the balance on hand at any time in respect of each drug of addiction will be clearly apparent.

[*Regulation 44 amended in Gazette 23 September 1983 p. 3803.*]

Inventory of Drugs of Addiction

45. (1) An inventory of drugs of addiction held in stock shall be made—

(a) at intervals of not more than one month by every person required to keep a Register of Drugs of Addiction; and

(b) by a person who is about to relinquish control of drugs of addiction; and

(c) forthwith on assuming control by any person who assumes control of drugs of addiction.

(2) If such inventory of drugs of addiction in stock does not agree with the balance recorded in the Register, the person required to keep the Register shall immediately notify the Permanent Head in writing of the discrepancy.

[*Regulation 45 amended in Gazette 29 June 1984 p. 1784.*]

46. (1) Where a person authorized to have drugs of addiction in his possession for the purpose of his profession or employment does not manufacture, retail, dispense or compound drugs of addiction, or where such dispensing or compounding is done by a medical practitioner, dentist, or veterinary surgeon for the purpose of treatment under his instructions, or his direct personal supervision, it shall be a sufficient compliance with regulation 44 if such person keeps a record of—

(a) the drugs of addiction obtained by him and the quantities of each;

(b) the person or firm from whom he obtained such drugs of addiction;

(c) the drugs of addiction disposed of or used by him, the quantities of each, and the date of such disposal or use;

(d) the manner in which such drugs of addiction were disposed of or used; and

(e) the drugs of addiction remaining in his possession and the quantities of each.

(2) The records required to be kept pursuant to this regulation shall be in a book, either written in ink, or gummed or pasted on the pages when the invoices or other documents are used for the purpose, and shall together with the drugs of addiction then in the possession of the authorized person, be produced for inspection on demand by a person appointed or authorized by or under the Act to inspect those records.

Records to be Retained for 7 years and Available on Demand

47. (1) All records, registers, prescription books, invoices and other documents relating to drugs of addiction, and transactions in regard thereto shall be kept by the person licensed or authorized to have drugs of addiction in his possession for not less than 7 years from the latest date on which such record, register, prescription book, invoice or other document was made or acted upon.

(2) The records, registers, prescription books, invoices or other documents and stocks of drugs of addiction on hand shall be made available for inspection on demand by a person authorized by or under the Act or regulations or by a member of the Police Force and shall be accounted for, during the inspection, by the person licensed or authorized to have drugs of addiction in his possession.

(3) In the event of a register being lost or destroyed the person to whom such register belongs shall upon becoming aware of the loss or destruction make and forward to the Permanent Head a statutory declaration concerning that loss or destruction and shall immediately take stock of all drugs of addiction in his possession and enter particulars of those stocks in a new register in accordance with the requirements of these regulations.

(4) A person authorized or licensed to procure and be in possession of a drug of addiction, on ceasing to be so authorized or licensed shall, if requested by the Permanent Head, surrender any records, registers, prescription books, invoices or other documents and stocks of drugs of addiction that are in his possession to the Permanent Head.

(5) A person required by these regulations to make and keep records, registers, returns, prescription books, invoices and other documents relating to drugs of addiction shall not make an entry therein which is false or untrue in any particular.

(6) The duplicate copy of the form approved by the Permanent Head for the purposes of regulations 52 (3) (h) and 52A is a record to be retained for the purposes of this regulation.

[*Regulation 47 amended in Gazette 23 September 1983 p. 3804; 29 June 1984 p. 1784; 31 January 1986 p. 332.*]

Returns from Manufacturers and Wholesalers

[*Heading substituted in Gazette 23 September 1983 p. 3804.*]

48. (1) Every person who holds a licence to manufacture, distribute or sell drugs of addiction by wholesale shall complete and forward to the Permanent Head, every 7 days, a form approved by the Permanent Head for that purpose, reporting all transactions in drugs of addiction made by him during that week.

(2) The form referred to in subregulation (1) may be required by the Permanent Head to be in a code approved by him from time to time and shall describe the composition, form, strength, size and quality of each drug of addiction and identify the person from whom, or to whom a drug of addiction has been obtained or supplied.

[*Regulation 48 substituted in Gazette 23 September 1983 p. 3804; amended in Gazette 29 June 1984 p. 1784.*]

Drugs of Addiction for Use on Ships and Aircraft

49. (a) Ships.—

(1) The master of a ship is authorized to procure and be in possession of any drug of addiction which is necessary to complete the equipment of the ship in order to comply with the requirements of the Navigation Act.

(2) The holder of a licence or other authorized person may supply such drug on receipt of a written order signed by the master of the ship and endorsed by the manager, or a person authorized in writing by him, of the ship's agents in Western Australia certifying that the drug of addiction is necessary to complete the equipment of the ship in order to comply with the requirements of the Navigation Act.

(b) Aircraft.—

(1) The person in charge of an aircraft is authorized to be in possession of a drug of addiction for the purpose of medical treatment on such aircraft but only in such quantity as does not exceed the quantity required by the Department of Civil Aviation to be carried on the aircraft.

(2) The holder of a licence or other authorized person may supply such drug of addiction on receipt of a written order signed by the manager, or a person authorized in writing by him, of the airline company or firm responsible for the operation of the aircraft in Western Australia.

(c) Any person who supplies a drug of addiction pursuant to this regulation shall within 24 hours of such supply report the details to the Permanent Head or the officer in charge of the nearest Police Station.

[*Regulation 49 amended in Gazette 29 June 1984 p. 1784.*]

Drugs of Addiction at Hospitals

50. (a) Where a Pharmaceutical Chemist is Employed.—The pharmaceutical chemist in charge of the pharmacy department of a hospital shall be responsible for ordering, storing and issuing all drugs of addiction in such hospital and for keeping records of drugs of addiction as required by these regulations.

(b) Where a Pharmaceutical Chemist is not Employed.—The matron of a hospital or other person authorized by the Permanent Head shall be responsible for ordering, issuing and storing all drugs of addiction in such hospital and for keeping records of drugs of addiction as required by these regulations.

[*Regulation 50 amended in Gazette 29 June 1984 p. 1784.*]

Prescriptions

51. (1) A prescription for the supply of a drug of addiction shall comply with the following conditions—

(a) the prescription shall bear on its face, legibly printed or written in ink the name and address of the prescriber; and

(b) the prescription shall bear on its face, legibly written in ink in the handwriting of the prescriber—

(i) the date when it is written;

(ii) the signature of the prescriber;

(iii) the name and full address of the patient or, in the case of a prescription for veterinary use, the name and full address of the person having the care of the animal for which it is intended;

(iv) the description and quantity of the drug or preparation containing the drug to be dispensed;

(v) adequate directions for use, including where applicable, the dose to be taken or administered;

(vi) where it is to be dispensed more than once, the maximum number of times it may be repeated and the intervals at which it may be dispensed;

(vii) the words 'for animal treatment only' if it is written by a veterinary surgeon;

(viii) the words 'for dental treatment only' if it is written by a dentist;

(ix) where it contains an unusual dose, an indication that such is intended, by underlining that part of the prescription and initialling the same in the margin;

(c) the prescription shall not bear the impression of a rubber stamp or other such contrivance in lieu of the written signature of the medical practitioner, dentist or veterinary surgeon by whom it has been issued.

(2) With the written approval of the Permanent Head a person authorized to prescribe drugs of addiction may issue a typewritten prescription where the Permanent Head is satisfied that by reason of physical infirmity the prescriber is unable to write legibly in his own handwriting but in that case the prescriber shall sign the prescription with his usual signature.

[*Regulation 51 substituted in Gazette 23 September 1983 p. 3804; amended in Gazette 29 June 1984 p. 1784; 31 January 1986 p. 332.*]

51A. For the purposes of regulations 51AA to 51G—

"drug addict" means a person who is—

(a) under a state of periodic or chronic intoxication produced by consumption of a drug of addiction or any substitute therefor;

(b) under a desire or craving to take a drug of addiction or any substitute therefor until he has so satisfied that desire or craving; or

(c) under a psychic or physical dependance to take a drug of addiction or any substitute therefor.

[*Regulation 51A substituted in Gazette 29 August 1980 p. 3028; amended in Gazette 12 October 1984 p. 3267.*]

51AA. A drug addict shall, when seeking to obtain from a medical practitioner—

(a) a drug of addiction; or

(b) a prescription or document prescribing the use, sale or supply of a drug of addiction,

disclose to the medical practitioner the fact that he is a drug addict.

[*Regulation 51AA inserted in Gazette 12 October 1984 p. 3267.*]

51B. A medical practitioner shall not write, issue or authorize a prescription or document prescribing the use, sale or supply of a drug of addiction, other than methadone or supply a drug of addiction, other than methadone for the treatment of a person who is—

(a) a drug addict; or

(b) a person who has been named as a drug addict by the Permanent Head by notice forwarded to the medical practitioner,

unless he has first obtained written authorization to do so from the Permanent Head.

[*Regulation 51B inserted in Gazette 29 August 1980 p. 3028; amended in Gazettes 29 June 1984 p. 1784; 23 May 1986 p. 1716.*]

51C. A medical practitioner shall not write, issue or authorize a prescription or document prescribing the use, sale or supply of methadone or supply methadone for the treatment of a person who is—

(a) a drug addict; or

(b) a person who has been named as a drug addict by the Permanent Head by notice forwarded to the medical practitioner,

unless the medical practitioner has—

(c) notified the Permanent Head of the condition of health of that person in accordance with the *Drugs of Addiction Notification Regulations 1980* as in force under the *Health Act 1911* from time to time; and

(d) received written authorization to do so from the Permanent Head.

[*Regulation 51C inserted in Gazette 29 August 1980 p. 3029; amended in Gazette 29 June 1984 p. 1784.*]

51D. (1) Before an authorization is issued by the Permanent Head for the treatment of a person with methadone the person in relation to whom the treatment is to be authorized, prescribed or used shall be assessed for such treatment by—

(a) a medical practitioner employed by the Alcohol and Drug Authority established under the *Alcohol and Drug Authority Act 1974;*

(b) a medical practitioner selected by the Authority referred to in paragraph (a) and approved of by the Permanent Head;

(c) a psychiatrist employed by the Mental Health Services of the State;

(d) a psychiatrist in the course of treating that person at a psychiatric unit of a hospital that is approved of as a teaching hospital as defined in the *Hospitals Act 1927;* or

(e) a medical officer attached to a regional hospital established under the *Hospitals Act 1927* who is approved of by the Permanent Head.

(2) A person who makes an assessment for the purposes of subregulation (1) shall specify in the assessment—

(a) the maximum daily dose not to be exceeded in the treatment in relation to the person with respect to whom the assessment is made; and

(b) the maximum period of the treatment,

and shall sign the assessment in his usual signature.

(3) Regulation 51C does not apply to or in relation to the carrying out of an assessment by a person referred to in subregulation (1) (a) or (b).

[*Regulation 51D inserted in Gazette 29 August 1980 p. 3029; amended in Gazette 29 June 1984 p. 1784.]*

51E. (1) In an authorization given with respect to the treatment of a particular drug addict with methadone the Permanent Head may specify that any one or more of the conditions and restrictions set out below apply, namely—

(a) that the prescription be issued by, or treatment administered by, a specified medical practitioner;

(b) that the type of methadone prescribed or administered be of the type specified;

(c) that the amount to be prescribed or used for treatment shall not exceed the amount specified;

(d) that the amount to be prescribed or used on any one day shall not exceed the amount specified;

(e) that the concentration to be prescribed or used shall not exceed the concentration specified;

(f) that the intervals between the issue of prescriptions or the administration of the treatment shall be such as are specified;

(g) that the prescription be supplied at the pharmacy or institution specified;

(h) that the amount dispensed on a single prescription form shall not exceed such amount as is specified;

(i) the amount that may be supplied on any one day shall not exceed such amount as is specified.

(2) Subject to subregulation (3) an authorization under subregulation (1) is valid for a period of 3 months from the date of its issue or such earlier date (if any) as is specified.

(3) The Permanent Head may at any time revoke an authorization or if the period has not expired vary the period for which the authorization is valid.

(4) A medical practitioner shall not write, issue or authorize a prescription or document prescribing the use, sale or supply of methadone or supply methadone contrary to such conditions and restrictions as are specified.

(5) A pharmaceutical chemist shall not sell or supply methadone otherwise than in accordance with such conditions and restrictions as are specified.

(6) An authorization issued under these regulations prior to 1 October 1980 is valid until it is revoked by the Permanent Head or until it expires whichever first occurs.

(7) In this regulation "specified" means by the Permanent Head in an authorization issued by him under this regulation in relation to a drug addict.

[*Regulation 51E. inserted in Gazette 29 August 1980 p. 3029; amended in Gazette 29 June 1984 p. 1784.*]

51F. (1) A medical practitioner shall not write, issue or authorize a prescription or document or supply a drug of addiction for the treatment of a person, other than a drug addict, for a period in excess of 30 days unless he has first obtained written authorization to do so from the Permanent Head.

(2) Where a medical practitioner has written, issued or authorized a prescription or document prescribing the use, sale or supply of a drug of addiction, for the treatment of a person other than a drug addict, or supplied a drug of addiction for the treatment of a person, other than a drug addict, for a period of 30 days the medical practitioner shall not thereafter write, issue or authorize a prescription or document prescribing the use, sale or supply of a drug of addiction in relation to that person or supply a drug of addiction in relation to that person unless—

(a) the medical practitioner has first obtained written authorization under this regulation to do so from the Permanent Head; or

(b) the Permanent Head has issued an authorization under this regulation to do so in relation to that person and the authorization is current.

(3) Notwithstanding any authorization referred to in subregulation (1) or (2) a medical practitioner shall not write, issue or authorize a prescription or document prescribing the use, sale or supply of methadone for the treatment of a person or supply methadone for the treatment of a person except for a person suffering from intractable pain arising from a condition of health other than addiction to drugs.

(4) In any authorization issued for the purposes of subregulation (1) or (2) given with respect to a particular person the Permanent Head may specify that any one or more of the conditions and restrictions set out below apply, namely—

(a) that the prescription be issued by or the treatment be administered by a specified medical practitioner;

(b) that only a specified drug of addiction be prescribed or used for treatment;

(c) that the type of the drug of addiction specified be of the type specified;

(d) that the amount to be prescribed or used for treatment shall not exceed the amount specified;

(e) that the amount to be prescribed or used on any one day shall not exceed the amount specified;

(f) that the concentration to be prescribed shall not exceed the concentration specified;

(g) that the intervals between the issue of prescriptions or the administration of the drug of addiction shall be such as are specified;

(h) that the prescription be supplied at the pharmacy specified;

(i) that the amount dispensed on a single prescription not exceed such amount as is specified;

(j) that the amount that may be supplied on any one day shall not exceed such amount as is specified;

(5) An authorization issued for the purposes of subregulation (1) or (2) is valid for such period as is specified unless revoked by the Permanent Head before the expiration of that period.

(6) A medical practitioner shall not write, issue or authorize a description or document prescribing the use, sale or supply of a drug of addiction or supply a drug of addiction otherwise than in accordance with such conditions and restrictions as are specified.

(7) A pharmaceutical chemist shall not sell or supply a drug of addiction otherwise than in accordance with such conditions and restrictions as are specified pursuant to this regulation.

(8) An authorization issued prior to 1 October 1980 is valid until revoked by the Permanent Head or until it expires whichever first occurs.

(9) In this regulation "specified" means by the Permanent Head in an authorization issued by him under this regulation in relation to a person other than a drug addict.

[*Regulation 51F inserted in Gazette 29 August 1980 pp. 3030-31; amended in Gazettes 23 September 1983 p. 3805; 29 June 1984 p. 1784; 8 February 1985 p. 520.*]

51G. (1) Notwithstanding regulations 51B to 51F (inclusive) a medical practitioner shall not write, issue or authorize a prescription or document prescribing the use, sale or supply of any of the following drugs of addiction, namely,

Amphetamine

Dexamphetamine

Methyl amphetamine

Phenmetrazine

or the salts of any of those substances and any preparation or admixture containing any of those substances, or the salts of any of those substances,

or supply any such drug of addiction in relation to any person unless he is authorized to do so by the Permanent Head.

(2) An authorization by the Permanent Head referred to in subregulation (1) may be varied or revoked by the Permanent Head at any time.

(3) A medical practitioner who receives an authorization given pursuant to subregulation (1), whether varied under subregulation (2) or not, shall not, in any event, write, issue or authorize a prescription or document authorizing the use, sale or supply of any of the drugs of addiction referred to in subregulation (1) or supply any such drug of addiction in relation to any person other than for the treatment of a person suffering from narcolepsy or a person suffering from brain damage.

[*Regulation 51G inserted in Gazette 29 August 1980 p. 3031; amended in Gazettes 23 September 1983 p. 3805; 29 June 1984 p. 1784; 8 February 1985 p. 520.*]

51H. (1) A dentist shall not write, issue or authorize a prescription or document for a drug of addiction other than—

(a) codeine;

(b) a preparation of codeine which does not contain any other drug of addiction; or

(c) pentazocine,

and the prescription or document shall not authorize treatment of a person for a period in excess of 5 days.

(2) A dentist may supply a patient with codeine and preparations of codeine not containing any other drug of addiction for a period of medication not exceeding 5 days.

[*Regulation 51H inserted in Gazette 8 February 1985 p. 520.*]

51J. (1) Notwithstanding regulations 51B to 51F a medical practitioner shall not write, issue or authorize a prescription or document prescribing the use, sale or supply of methylphenidate for oral use or the salts, preparations or admixture of methylphenidate for oral use, or supply such a drug of addiction in relation to any person unless he is authorized to do so by the Executive Director.

(2) Subregulation (1) does not apply to the oral use of methylphenidate for therapeutic trials of up to 30 days when initiated in a teaching hospital as defined in the *Hospitals Act 1927* or at the Psychiatric Services Branch of the Health Department of Western Australia.

(3) An authorization by the Executive Director referred to in subregulation (1) may be varied or revoked by the Executive Director.

(4) A medical practitioner who receives an authorization given pursuant to subregulation (1), whether varied under subregulation (3) of this regulation or not, shall not in any event, write, issue or authorize a prescription or document authorizing the use, sale or supply of the drug of addiction referred to in subregulation (1) or supply any such drug of addiction in relation to any person other than for the treatment of a person suffering from narcolepsy, brain damage, behavioural disorders in children or for the continued treatment of a person who previously had authorization for the supply of this drug of addiction approved by the Executive Director.

[*Regulation 51J inserted in Gazette 8 February 1985 p. 521 as regulation 51H; renumbered as 51J in Gazette 7 June 1985 p. 1941.*]

Dispensing Drugs of Addiction

52. (1) Subject to the Act and these regulations a drug of addiction shall be dispensed or supplied only in accordance with a prescription complying with the requirements of the regulations.

(2) A medical practitioner, pharmaceutical chemist, or veterinary surgeon or an assistant under the direct personal supervision of a medical practitioner, pharmaceutical chemist, or veterinary surgeon shall be the only person who shall dispense a drug of addiction.

(3) The following conditions shall be observed by persons dispensing prescriptions referred to in this regulation—

(a) the person to whom a prescription is submitted for dispensing shall satisfy himself—

(i) that the prescription is in accordance with the requirements of the Act; and

(ii) that the person who signed the prescription is a medical practitioner, a dentist or a veterinary surgeon duly registered within the State;

(b) the prescription shall not be dispensed more than the maximum number of times indicated thereon, or at intervals less than those indicated therein, and on each occasion at the time at which it is dispensed it shall be marked in ink, by writing in the hand of the dispensing pharmacist, to show clearly his usual signature, the date upon which it is dispensed, and marked or stamped with the name and address of the pharmacy at which it is dispensed;

(c) a prescription written by a veterinary surgeon shall be dispensed once only;

(d) a prescription shall be dispensed once only where the prescription directs the dispensing of 2 or more items as separate articles;

(e) subject to subregulation (7)—

(i) where a prescription is produced to him and the prescription prescribes no more than one occasion on which it is to be dispensed, the person dispensing the prescription shall retain the prescription in safe custody after having dispensed it;

(ii) where a prescription is produced to him and the prescription prescribes more than one occasion on which it is to be dispensed, the person dispensing the prescription shall, after having dispensed it as directed in the prescription, mark the prescription with the number of occasions remaining to be dispensed and return it to the person producing it but if there remain no more occasions on which it is to be dispensed shall, after having dispensed it as directed in the prescription, retain the prescription in safe custody:

(f) the person who dispenses a prescription which does not clearly indicate the maximum number of times such prescription is to be dispensed or the intervals at which it is to be repeated, or which has reached the last occasion upon which it can be lawfully dispensed according to the maximum indicated thereon, shall write in ink, stamp, or mark in legible letters across such prescription the word "cancelled";

(g) the person who dispenses a prescription shall enter, or cause to be entered, in the book prescribed by regulation 44, a proper record of the transaction which record shall be made in such a way as to be easily understood;

(h) before the drug of addiction is handed to the purchaser, details of the transaction, including—

(i) prescription number;

(ii) name and address of patient;

(iii) drug description;

(iv) quantity of the drug;

(v) directions for use;

(vi) date of the prescription; and

(vii) the name and address of the person who wrote the prescription,

shall be entered on a duplicate form approved by the Permanent Head and shall be signed and dated by the person who actually dispensed the drug of addiction;

(j) in the case of a repeat prescription, an entry in the Prescription Book of the fact of the repeat and the repeat number signed and dated shall be sufficient compliance with this regulation;

(k) the label on the bottle or package containing the drug of addiction shall be marked with the identifying letter or number of the prescription as appearing in the Prescription Book;

(l) the Prescription Book shall be kept for a period of one year from the date of the last dispensing recorded therein at the place at which the drug of addiction was dispensed and shall be produced on demand to any person authorized in that behalf under the Act; and

(m) the drug of addiction so dispensed shall conform in quantity, description, composition, strength, form and every other material particular to the directions of the prescriber,

and in respect of a prescription prescribing a drug of addiction issued under the *National Health Act 1953* or the *Repatriation Act 1920* of the Commonwealth a copy of such a prescription is deemed to be a prescription for the purposes of this regulation.

(4) A prescription marked "cancelled" or that is more than 6 months old shall not be dispensed.

(5) A prescription which is illegible or defaced or which appears to be for the purpose of enabling some unauthorized person to obtain a drug of addiction, or which does not appear to be genuine, shall not be dispensed.

(6) A pharmaceutical chemist shall retain possession of any prescription prescribing a drug of addiction presented to him for dispensing and which he suspects of being false in any particular and hold the prescription for such reasonable period as will enable him to satisfy himself as to its genuineness and make enquiries concerning the *bona fides* of the person by whom it is presented or the identity of the individual by whom such prescription purports to have been written.

(6a) When a pharmaceutical chemist to whom a prescription is submitted for dispensing is satisfied that the prescription is not in accordance with the requirements of the Act or regulations, he shall cancel it and endorse upon its face in ink in his own handwriting the date and his usual signature, together with the address of the dispensary either stamped or written and forward it to the Permanent Head and inform the Permanent Head of the relevant circumstances and the reasons for his refusal to dispense the prescription.

(7) A pharmaceutical chemist shall forward to the Permanant Head not later than 21 days after the last day of each month every form of prescription of a drug of addiction (whether original or copy thereof) retained at the pharmacy pursuant to this regulation during that month unless he is required to do otherwise by a law of the Parliament of the Commonwealth.

(8) A pharmaceutical chemist shall deliver up any document, prescription, authorization or record relating to the sale or supply of a drug of addiction upon request made by any inspector appointed under the *Health Act 1911* or to any other person authorized in that behalf by the Minister.

[*Regulation 52 amended in Gazettes 29 August 1980 p. 3031; 23 September 1983 pp. 3805-6; 29 June 1984 p. 1784; 31 January 1986 pp. 332-33.*]

Movement of Drugs of Addiction in other circumstances

52A. Any movement of stocks of drugs of addiction other than by prescription and other than supplies received from wholesalers shall be recorded on a duplicate form approved by the Permanent Head.

[*Regulation 52A and heading inserted in Gazette 31 January 1986 p. 333.*]

Documentation of Drugs of Addiction

52B. Every owner of a pharmacy which dispenses drugs of addiction shall return the original of the completed, approved form referred to in regulations 52 (3) (h) and 52A to the Department monthly, by the 7th day of the following month, and where there have been no transactions during the previous month, the form shall be returned to the Department completed as to the name and address of the pharmacy and otherwise marked "NIL".

[*Regulation 52B and heading inserted in Gazette 31 January 1986 p. 333.*]

Dispensing Drugs of Addiction in Case of Emergency

53. (1) Where a medical practitioner or veterinary surgeon in a case of emergency orally or by telephone or telegram directs the dispensing of a drug of addiction, he shall forthwith write a prescription complying with the conditions prescribed in regulation 51 mark such prescription so as to show clearly that it is in confirmation of the directions given by him orally or by telephone or telegram, and despatch such prescription within 24 hours to the person by whom the drug of addiction was dispensed.

(2) A person by whom a drug of addiction was dispensed in accordance with subregulation (1), shall where the prescription is not received by him within 72 hours, immediately report the circumstances to the Permanent Head.

[*Regulation 53 amended in Gazette 23 September 1983 p. 3806.*]

Dispensing Certain Drugs of Addiction

[*Heading and regulation inserted in Gazette 23 September 1983 p. 3806*]

53A. (1) A person shall not dispense a prescription for or supply upon a prescription any of the following drugs of addiction, namely,

Dextromoramide
Hydromorphone
Methadone
Morphine
Pethidine

or the salts of any of those substances and any preparation or admixture containing any of those substances, or the salts of any of those substances, unless—

(a) he is familiar with the prescriber's handwriting; or

(b) he has verified with the purported prescriber that the prescription was written by him.

(2) Where a person cannot comply with subregulation (1), for good cause, he may dispense the prescription for, or supply upon the prescription, a quantity of the drug of addiction sufficient to enable treatment at the rate prescribed for no more than 2 days.

[*Regulation 53A: See note under heading. R.53A amended in Gazette 31 January 1986 p. 333.*]

Delivery of Drugs of Addiction on Order

54. (1) Subject to regulation 53 and to subregulation (3) a drug of addiction shall not be delivered to a person except—

(a) on the authority of a written order—

(i) legibly written in ink;

(ii) bearing on the face thereof—

(I) the date when it is written;

(II) the name and address of the person requiring it to be supplied;

(III) the quantity and description of the drug of dependence to be supplied; and

(iii) signed by a person licensed or otherwise authorized to procure or be in possession of the drug of addiction; or

(b) on the authority of an order placed by telephone or telex, but such an order shall be confirmed by the person requiring the drugs to be supplied, by the signing and and dating of the despatch note or invoice delivered with the goods and the despatch note or invoice shall be returned to the supplier within 7 days of the delivery or the supplier shall notify the Permanent Head of the default.

(2) A drug of addiction shall not be delivered to any person not licensed, or otherwise authorized to be in possession of the drug of addiction, who purports to be sent by or on behalf of the person so licensed or authorized, unless the firstmentioned person produced an authority in writing signed by the person so licensed or authorized to receive the drug of addiction on his behalf, and unless the person supplying the drug of addition is satisfied that the authority is genuine.

(3) This regulation does not apply to medicines dispensed in pursuance of the foregoing regulations.

[*Regulation 54 amended in Gazette 23 September 1983 p. 3807.*]

54A. A person forwarding for delivery a drug of addiction shall enclose the drug separately from goods of any other kind in a secure and sturdy package without exterior writing which might indicate the contents of the package and clearly address the package to the authorized person.

[*Regulation 54A inserted in Gazette 23 September 1983 p. 3807.*]

Common Carrier Protected

55. A common carrier or his employee is hereby authorized to be in possession of any drug of addiction so far only as the possession is necessary for the transport of the drug of addiction in the ordinary course of business.

Safe Custody of Drugs of Addiction

56. (1) Subject to regulation 56A any person licensed or authorized to have a drug of addiction in his possession shall store such drug of addiction in a poisons cupboard, and such person shall ensure the safekeeping of the key to the poisons cupboard and shall also keep the poisons cupboard locked at all times except when drugs of addiction are being placed into or removed from it.

(2) This regulation does not apply to—

(a) a person who has been supplied with the drug of addiction pursuant to a prescription from a medical practitioner or veterinary surgeon; or

(b) a medical practitioner, dentist or veterinary surgeon when transporting the drug of addiction for the purpose of his profession or practice, if such medical practitioner, dentist or veterinary surgeon takes reasonable precautions to protect such drug of addiction against theft or loss.

[*Regulation 56 amended in Gazettes 7 September 1971 p. 3278; 23 September 1983 p. 3807.*]

56A (1) Where a pharmacist is in possession of Eighth Schedule poisons for the purposes of his profession or employment, he shall store those poisons in the type of safe prescribed by this regulation or in similar storage accommodation approved by the Permanent Head for this purpose.

(2) The safe required by subregulation (1) shall be in a portion of the premises not accessible to the public and shall be—

(a) constructed of black mild steel plate not less than 9.5 millimetres thick;

(b) constructed with continuous welding of all edges; and fitted with a solid mild steel bar of not less than eight millimetres at its smallest diameter, situated not more than 100 millimetres above or below the lock, fixed to both sides of the safe by drilling and backwelding;

(c) fitted with a door constructed of black mild steel plate not less than 9.5 millimetres thick, the door being flush fitting with a clearance around the door of not more than 1.5 millimetres;

(d) fitted with 2 or more fixed locking bars welded to the inside face of the door near the hinge edge at not greater distances than 300 millimetres apart from centre of locking bar to centre of locking bar; one fixed locking bar to be not further than

150 millimetres from the top of the safe door, and one fixed locking bar to be not further than 150 millimetres from the bottom of the safe door; each locking bar engaging in a rebate in the cupboard body when the door is closed.

(e) fitted with a 5 lever keylock, or locking mechanism providing at least equivalent security, securely affixed to the rear face of the door; when the height of the safe door exceeds 610 millimetres but does not exceed 915 millimetres a second 5 lever keylock or locking mechanism providing at least equivalent security shall be securely affixed to the rear face of the door, and this lock shall be keyed alike to the first lock.

(f) securely attached to the wall or floor in the following manner—

(i) Where the wall and the floor are constructed of brick or concrete the safe shall be attached to the wall or the floor by means of suitably sized expanding bolts through holes 9.5 millimetres in diameter drilled in the rear or floor of the safe.

(ii) Where the wall only is constructed of brick or concrete the safe shall be attached to the wall by means of suitably sized expanding bolts through holes 9.5 millimetres in diameter drilled in the rear of the safe.

(iii) Where the floor is constructed of brick or concrete, but the wall is of timber construction, the safe shall be attached to the floor by means of suitably sized expanding bolts through holes 9.5 millimetres in diameter drilled in the bottom of the safe.

(iv) Where neither a floor nor a wall constructed of brick or concrete is available, the safe shall be attached to the wall or floor by a method that will ensure that the safe cannot be easily removed.

(3) Notwithstanding subregulation (2) a safe built or placed under the floor, shall be deemed to have met the security specifications of that subregulation it it meets the following requirements—

(a) the container and neck of the safe shall be constructed of black mild steel plate;

(b) the container and neck of the safe shall be embedded in reinforced concrete; and

(c) the safe shall have a substantial closure fitted with a 5 lever keylock or other locking mechanism providing at least equal security, or alternatively a keyless combination lock.

(4) A pharmacist shall keep in his immediate and personal possession the key to any such safe referred to in subregulation (1) and the safe shall be locked at all times except when items are being placed into or removed from it.

[*Regulation 56A inserted in Gazette 7 September 1971 p.3278; amended in Gazettes 3 May 1974 pp. 1434-35; 15 April 1976 p. 1183; 29 June 1984 p. 1784.*]

56B. All Eighth Schedule poisons—

(a) stored in the pharmacy department of a hospital which employs a pharmacist, shall be stored in a locked safe kept solely for that purpose or in similar storage accommodation approved by the Permanent Head and the key shall be kept in the possession of the pharmacist-in-charge and not left on the premises where the Eighth Schedule poisons are stored except when it is given into the possession of another pharmacist, medical practitioner or dentist;

(b) in a hospital which does not employ a pharmacist, shall be stored in the hospital in locked storage accommodation approved by the Permanent Head prior to the distribution of supplies to wards, and ward supplies shall be stored in locked cupboards in wards or in locked portions of cupboards kept solely for the storage of Eighth Schedule poisons;

(c) kept by persons licensed to procure, manufacture or supply drugs of addiction by wholesale dealing, shall be stored in a locked storage accommodation approved by the Permanent Head and the key shall be in the possession of the person so licensed or in the possession of some other person authorized by the Permanent Head.

[*Regulation 56B inserted in Gazette 7 September 1971 p. 3279; amended in Gazette 29 June 1984 p. 1784.*]

56C. The exterior surface of the cupboard or safe in which an Eighth Schedule drug is stored shall not bear the word "poison".

[*Regulation 56C inserted in Gazette 28 February 1986 p. 618.*]

56D. The cupboard or safe in which drugs of addiction are kept in accordance with regulations 56, 56A or 56B shall not be used for any purpose other than the storage of poison.

[*Regulation 56D inserted in Gazette 1 August 1986 p. 2739.*]

Labelling

57. (1) A person shall not supply any drug of addiction unless the package or bottle containing the drug of addiction is plainly labelled or marked to show the quantity of such drug of addiction contained therein.

(2) A person shall not supply any preparation or admixture containing any drug of addiction, unless the package or bottle containing that preparation or admixture is plainly labelled or marked to show the total quantity of such preparation or admixture in the package or bottle and the percentage or quantity of the drug of addiction contained therein, or in the case of tablets or other articles, the number of such tablets or articles in the package or bottle and the percentage or quantity of the drug of addiction contained in each tablet or article.

(3) This regulation does not apply to any drug of addiction, preparation, or admixture dispensed in accordance with these regulations.

Improper Prescribing or Use of Drugs of Addiction

58. (1) A medical practitioner, dentist or veterinary surgeon shall not knowingly give a prescription for a drug of addiction merely for purposes of addiction.

(2) A medical practitioner, dentist or veterinary surgeon shall not knowingly supply or administer a drug of addiction merely for purposes of addiction.

[*Regulation 58 amended in Gazettes 23 September 1983 p. 3807; 20 March 1987 p. 954.*]

Names of Persons from whom Licence or Authority Withdrawn to be Published

59. A decision of the Permanent Head cancelling, suspending or revoking an authorization, licence or permit conferred or issued under the Act or these regulations or any other decision of the Permanent Head may be published in the *Government Gazette.*

[*Regulation 59 substituted in Gazette 29 August 1980 p. 3031; amended in Gazette 29 June 1984 p. 1784.*]

Appeals

60. (1) Any person desirous of appealing under the provisions of section 29 of the Act shall lodge with the Clerk of Petty Sessions of the Court of Petty Sessions held nearest to the place of business of the appellant notice of appeal in the Form 12 in Appendix A.

(2) A copy of the notice shall be served on the Permanent Head within 7 days after lodging the notice with the Clerk of Petty Sessions.

(3) On proof by affidavit to the satisfaction of the magistrate that the notice has been duly served, the magistrate shall cause to be sent to the parties written notice of a date and time for the hearing of the appeal, which date shall not be less than 14 clear days from the service of the notice.

[*Regulation 60 amended in Gazette 29 June 1984 p. 1784.*]

61. If either party to the appeal neglects to appear personally or by counsel or solicitor on the date and at the time fixed for hearing, the magistrate may—

(a) where the appellant fails to appear, dismiss the appeal;

(b) where the Permanent Head fails to appear, hear the appeal or adjourn it to some other date.

[*Regulation 61 amended in Gazette 29 June 1984 p. 1784.*]

62. On the hearing and determination of the appeal the magistrate may make such order as to costs to be paid by either party to the appeal as he may think just.

63. On the hearing of the appeal, it shall proceed according to the procedure and rules of evidence applicable in the Court of Petty Sessions.

APPENDIX A

Form 1

Poisons Act 1964

LICENCE TO PROCURE, MANUFACTURE AND SUPPLY POISONS (OTHER THAN DRUGS OF ADDICTION) BY WHOLESALE DEALING

This licence is granted to .....and authorizes him to procure, manufacture and supply by wholesale dealing on behalf of.....the poisons specified in the *1st, 2nd, 3rd, 4th, 6th, 7th Schedules to the Poisons Act.

Subject to the following conditions:—

1. The poisons will be manufactured at premises situated at .....
.....

(a) under the personal supervision of .....
who holds the qualification.....(or)

(b) under the direction of.....who
holds the qualification.....and
under the personal supervision of .....
who is an experienced person within the meaning of the regulations
(or)

(c) .....
.....

2. The poisons will be supplied from premises situated at.....

(a) under the personal supervision of .....who holds the qualification.....(or)

(b) under the direction of.....who holds the qualification.....and under the personal supervision of ..... who is an experienced person within the meaning of the regulations (or)

(c) .....

3. (a) .....

(b) .....

Dated at Perth.....19.....

Valid until 30 June, 19.....

.....

Executive Director, Public Health and Scientific Support Services.

---

*Strike out whichever is not applicable.

Form 1A

Poisons Act 1964

APPLICATION FOR LICENCE TO PROCURE, MANUFACTURE AND SUPPLY POISONS (OTHER THAN DRUGS OF ADDICTION) BY WHOLESALE DEALING

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia.
‡Perth.

I, Mr. / Mrs. / Miss .....
(Full Name)

hereby apply for a licence to procure, manufacture and supply by wholesale dealing on behalf of .....the poisons specified in the *1st, 2nd, 3rd, 4th, 6th, 7th Schedules to the Poisons Act.

In support of this application I declare that—

1. The poisons will be manufactured at premises situated at .....

(a) under the personal supervision of.....who holds the qualification.....(or)

(b) under the direction of.....who holds the qualification .....and under the personal supervision of .....who is an experienced person within the meaning of the regulations (or)

(c) .....

2. The poisons will be supplied from premises situated at .....
.....

(a) under the personal supervision of .....
who holds the qualification.....(or)

(b) under the direction of.....who holds the qualification .....and under the personal supervision of .....who is an experienced person within the meaning of the regulations (or)

(c) .....
.....

Date..... .....
Signature of Applicant.

Fee $.....with application

---

*Strike out whichever is not applicable.
‡The present address is 60 Beaufort Street, Perth.

Form 2

Poisons Act 1964

LICENCE TO PROCURE, MANUFACTURE AND SUPPLY BY WHOLESALE DEALING DRUGS OF ADDICTION

This licence is granted to .....and authorizes him to procure, manufacture and supply by wholesale dealing on behalf of .....
.....the following drugs of addiction .....
.....

Subject to the following conditions:—

1. The drugs of addiction will be manufactured at premises situated at.....
.....

(a) under the personal supervision of .....
who holds the qualification.....(or)

(b) under the direction of.....who holds the qualification .....and under the personal supervision of.....who is an experienced person within the meaning of the regulations.

2. The drugs of addiction will be supplied from premises situated at .....
.....

(a) under the personal supervision of .....
who holds the qualification.....(or)

(b) under the direction of.....who holds the qualification .....and under,the personal supervision of.....who is an experienced person within the meaning of the regulations.

3. (a) .....

(b) .....

Dated at Perth.....19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 2A

Poisons Act 1964

APPLICATION FOR LICENCE TO PROCURE, MANUFACTURE AND SUPPLY BY WHOLESALE DEALING DRUGS OF ADDICTION

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
*Perth.

I, Mr. / Mrs. / Miss .....hereby apply for
(Full Name)

a licence to procure, manufacture and supply by wholesale dealing the following drugs of addiction .....
.....

In support of this application I declare that:—

1. The drugs of addiction will be manufactured at premises situated at .....
.....

   (a) under the personal supervision of .....
   who holds the qualification.....(or)

   (b) under the direction of.....who holds
   the qualification .....and under the
   personal supervision of.....who is an
   experienced person within the meaning of the regulations.

2. The drugs of addiction will be supplied from premises situated at .....
.....

   (a) under the personal supervision of .....
   who holds the qualification.....(or)

   (b) under the direction of.....who holds
   the qualification .....and under the
   personal supervision of.....who is an
   experienced person within the meaning of the regulations.

Date.....

.....
Signature of Applicant.

Fee $.....with application

---

*The present address is 60 Beaufort Street, Perth.

Form 3

Poisons Act 1964

PHARMACEUTICAL CHEMIST'S LICENCE TO SELL POISONS

This licence is granted to.....
and authorizes him to sell poisons at premises situated at .....
.....

Dated at Perth .....19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 3A

Poisons Act 1964

APPLICATION FOR PHARMACEUTICAL LICENCE TO SELL POISONS

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
*Perth.

Mr.
I, Mrs. .....
Miss (Full Name)

a pharmaceutical chemist registered to practice in Western Australia, hereby apply for a licence to sell poisons at premises situated at.....
.....which premises are registered as a Pharmacy under the *Pharmacy Act 1964*, the Registration Certificate in respect of which is No.....valid until 30 June, 19.....

Date.....

.....
Signature of Applicant.

Fee $.....with application

---

* The present address is 60 Beaufort Street, Perth.

Form 4

Poisons Act 1964

LICENCES TO SELL BY RETAIL POISONS SPECIFIED IN THE 6TH SCHEDULE

This licence is granted to.....
and authorizes him to procure, and to sell by retail, on behalf of.....
.....the poisons specified in the 6th Schedule to the *Poisons Act 1964*, at premises situated at.....
.....

Dated at Perth.....19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 4A

Poisons Act 1964

APPLICATION FOR LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 6TH SCHEDULE

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
* Perth.

Mr.
I, Mrs.....
Miss (Full Name)

hereby apply for a licence to sell, by retail, on behalf of.....
.....the poisons specified in the 6th Schedule to the *Poisons Act 1964*.

I declare that—

(a) I have attained the age of 21 years.

(b) The poisons will be sold only at premises situated at .....
.....

(c) The poisons will not be sold by an assistant under 16 years of age.

(d) The poisons will not be sold to anyone who is apparently under 16 years of age.

Date .....

.....
Signature of Applicant.

* The present address is 60 Beaufort Street, Perth.

Form 5

Poisons Act 1964

LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 2ND OR 6TH SCHEDULES

This licence is granted to.....
and authorizes him to procure, and to sell by retail, on behalf of.....
.....the poisons specified in the 2nd or 6th Schedules to the *Poisons Act 1964,* at premises situated at .....
.....

Dated at Perth.....19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 5A

Poisons Act 1964

APPLICATION FOR LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 2ND OR 6TH SCHEDULES

To the Executive Director, Public Health and Scientifc Support Services,
Health Department of Western Australia,
* Perth.

I, Mr. / Mrs. / Miss .....
(Full Name)

hereby apply for a licence to sell, by retail, on behalf of.....
.....the poisons specified in the 2nd or 6th Schedules of the *Poisons Act 1964,* at premises situated at.....
.....

I declare that—

(a) I have attained the age of 21 years.

(b) These premises are distant at least 8 kilometres from the nearest place at which a pharmaceutical chemist conducts a pharmacy.

(c) The poisons will not be sold by an assistant under 16 years of age.

(d) The poisons will not be sold to anyone who is apparently under 16 years of age.

Date.....

.....
Signature of Applicant.

Fee $.....with application

* The present address is 60 Beaufort Street, Perth.

Form 6

Poisons Act 1964

LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 7TH SCHEDULE

This licence is granted to.....
and authorizes him to procure, and sell by retail, on behalf of.....
....., at premises situated at
....., the following poisons specified in the 7th Schedule:—

.....
.....

Subject to the following conditions:—

.....
.....

Dated at Perth.....19.....

Valid until 30 June 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 6A

Poisons Act 1964

APPLICATION FOR LICENCE TO SELL BY RETAIL POISONS SPECIFIED IN THE 7TH SCHEDULE

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
* Perth.

I, Mr. / Mrs. / Miss ..... (Full Name)

hereby apply for a licence to sell, by retail, on behalf of.....
.....poisons specified in the 7th Schedule:—

.....
.....
.....

I declare that—

(a) I have attained the age of 21 years.

(b) The poisons will be sold only at premises situated at.....
.....

(c) The poisons will be sold only by myself or by an assistant who is not less than 16 years of age.

(d) The poisons will not be sold to anyone who is apparently under 16 years of age.

(e) .....

Date ....
Signature of Applicant.

Fee $.....with application

---

* The present address is 60 Beaufort Street Perth.

Form 6B

Poisons Act 1964

POISONS PERMIT (DISTRIBUTION OF SAMPLES)

This permit is granted to .....of ....., representative of....., licensed manufacturers of, or wholesale dealers in, drugs containing poisons specified in the First, Second, Third or Fourth Schedules to the *Poisons Act 1964;* and authorizes him/her to procure samples of the aforesaid drugs from .....

(Name of manufacturers or wholesalers)

and supply them to persons authorized by regulation 8A of the *Poisons Regulations 1965* (as amended) to receive them.

Dated at Perth....., 19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 6C

Poisons Act 1964

APPLICATION FOR POISONS PERMIT (DISTRIBUTION OF SAMPLES)

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
Perth.

I, ...of .....being a representative of ....., licensed manufacturer of, or wholesale dealer in, drugs containing poisons specified in the First, Second, Third or Fourth Schedules to the *Poisons Act 1964*, hereby apply for a permit to procure from .....

(Name of manufacturers or wholesalers)

and to supply to persons authorized to receive them, samples containing drugs specified in the First, Second, Third or Fourth Schedules to the *Poisons Act 1964.*

Dated at Perth ....., 19.....

Valid until 30 June, 19.....

.....
Signature of applicant

Fee $.....with application

Form 7

Poisons Act 1964

POISONS PERMIT (INDUSTRIAL)

This permit is granted to .....and authorizes him to purchase on behalf of..... from a manufacturer or wholesale dealer—

(a) the poisons specified in the..... Schedules to the *Poisons Act 1964*;

(b) the following poisons:—

.....
.....
.....

This permit is issued subject to the following conditions:—

(1) the poisons will be stored only at premises situated at .....
.....;

(2) the poisons will not be resold;

(3) the poisons will be used only for the following purposes:—

.....
.....
.....

(4) .....
.....
.....

Dated at Perth....., 19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 7A

Poisons Act 1964

APPLICATION FOR POISONS PERMIT (INDUSTRIAL)

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
‡ Perth.

I, Mr. / Mrs. / Miss .....
(Full Name)

hereby apply on behalf of.....for a permit to purchase from a manufacturer or wholesale dealer—

(a) the poisons specified in the *1st, 2nd, 3rd, 4th, 6th, 7th, 8th Schedules to the *Poisons Act 1964*; or

(b) the following poisons:—

.....
.....
.....

In support of this application I declare that—

(1) the poisons will be stored only at premises situated at .....

(2) the poisons will no be resold;

(3) the poisons will be used only for the following purposes:—

.....
.....
.....

(4) .....
.....
.....

Date.....

.....
Signature of Applicant.

Fee $.....with application

---

* Strike out whichever does not apply.
‡ The present address is 60 Beaufort Street, Perth.

Form 8

Poisons Act 1964

POISONS PERMIT (EDUCATIONAL, ADVISORY OR RESEARCH)

This permit is granted to .....and authorizes him to purchase on behalf of..... from a manufacturer or wholesale dealer—

(a) the poisons specified in the..... Schedules to the *Poisons Act 1964*;

(b) the following poisons:—

.....
.....
.....

This permit is issued subject to the following conditions:—

(1) The poisons will be stored only at premises situated at..... .....,

(2) the poisons will not be resold;

(3) the poisons will be used only for the following purposes:—

.....
.....
.....

(4) .....
.....
.....

Dated at Perth.....,19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 8A

Poisons Act 1964

APPLICATION FOR POISONS PERMIT (EDUCATIONAL, ADVISORY OR RESEARCH)

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
‡ Perth.

I, Mr. / Mrs. / Miss .....
(Full Name)

hereby apply on behalf of.....for a permit to purchase from a manufacturer or wholesale dealer—

(a) the poisons specified in the *1st, 2nd, 3rd, 4th, 6th, 7th, 8th Schedules to the *Poisons Act 1964*; or

(b) the following poisons:—

.....
.....
.....

In support of this application I declare that—

(1) The poisons will be stored only at premises situated at.....
.....

(2) the poisons will not be resold;

(3) the poisons will be used only for the following purposes:—

.....
.....
.....

(4) .....
.....
.....

Date..... .....
Signature of Applicant.

---

* Strike out whichever does not apply.
‡ The present address is 60 Beaufort Street, Perth.

Form 9

Poisons Act 1964

LICENCE TO HAWK, PEDDLE OR DISTRIBUTE POISONS

This licence is granted to.....and authorizes him to sell by hawking or peddling, or to distribute as a sample, the following poisons:—

.....
.....
.....

This licence is issued under section 48 of the *Poisons Act 1964*, and is subject to the following conditions, limitations and restrictions:—

.....
.....
.....

Dated at Perth.....19.....

Valid until.....19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 9A

Poisons Act 1964

APPLICATION FOR LICENCE TO HAWK, PEDDLE OR DISTRIBUTE POISONS

To The Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
* Perth.

I, Mr. / Mrs. / Miss .....
(Full Name)

of .....hereby apply for a licence to hawk,
(Address)

peddle, or distribute as a sample, the following poisons:—

.....
.....
.....

In support of this application I declare that—

(a) I have attained the age of 21 years;

(b) the poisons will not be sold by an assistant under 18 years of age;

(c) the poisons will not be sold to anyone who is apparently under 16 years of age;

(d) the poisons will be sold or distributed only in the areas specified in the licence;

(e) .....
.....

Date.....19.....

.....
Signature of Applicant.

---

* The present address is 60 Beaufort Street, Perth.

Form 10

Poisons Act 1964

APPLICATION FOR CLASSIFICATION OF A NEW DRUG

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
*Perth.

I (or we).....of
....., herewith make
application for classification of the new drug.....
.....

I (or we) request that this drug be—

(a) included in Schedule .....;

(b) exempted from inclusion in any Schedule;

(c) preparations containing not more than .....per cent of the drug be.....

In support of this application I (we) submit the following information:—

1. The (a) approved name of the drug.....

(b) generic name of the drug .....

2. The trade name (or names) .....

3. The proprietary name (or names).....

4. The chemical name .....

5. The chemical nature.....

6. The chemical structure and formula .....

7. Its description in precise chemical terms, together with its physical details .....

8. The nature and limits of any impurities present.....

9. Particulars of the tests and standards applied to control its potency, purity and safety during manufacture and storage .....

10. Full details of investigations made with respect to the safety and efficacy of the drug, including tests carried out by universities and/or research institutions, and clinical trials.

Note.—Full reports are required of adequate tests which will show whether or not the substance will be safe. The reports shall include detailed data derived from appropriate animal and other biological experiments in which the methods used and the results obtained are clearly set forth. Details of any reports which could bias an evaluation of the safety of the substance shall NOT be omitted. Special attention shall be given to—

(a) acute, sub-acute and chronic toxicity;

(b) uniformity of response within a species and among different species;

(c) occurrence of unusual or alarming reactions, such as carcinogenesis;

(d) known side effects;

(e) occurrence of sensitivity tolerance or idiosyncracy in response to the substance;

(f) metabolism, rate, extent and mode of elimination of the substance;

(g) any tendency towards accumulation in the body;

(h) any special incompatibility;

(i) method of assay.

11. A statement of the amounts of all ingredients, route of administration, proposed dosage, the claims to be made for such drug and a description of the pharmaceutical forms in which it is proposed to be sold.

12. Full details of proposed labelling and packaging.

13. Evidence of approval or rejection by any other statutory body or authority.

14. Complete bibliography of any publications relating to pharmacological and therapeutic actions, including clinical trials.

Date.....

.....
Signature of Applicant.

---

* The present address is 60 Beaufort Street, Perth.

Form 11

Poisons Act 1964

PERMIT TO SUPPLY FOR VETERINARY USE THE PREPARATIONS REFERRED TO IN REGULATION 39 (2)

This permit is granted to .....and
authorizes him to sell on behalf of.....
at premises situated at.....
the preparations referred to in regulation 39 (2).

Dated at Perth.....19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 11A

Poisons Act 1964

APPLICATION FOR PERMIT TO SUPPLY FOR VETERINARY USE THE PREPARATIONS REFERRED TO IN REGULATION 39 (2)

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
* Perth.

I, Mr. / Mrs. / Miss ..... (Full Name)

hereby apply for a permit to sell on behalf of .....
at premises situated at .....the preparations referred to in regulation 39 (2) of the *Poisons Regulations 1965*.

Date.....19.....

.....
Signature of Applicant.

Fee $.....with application

---

* The present address is 60 Beaufort Street, Perth.

Form 11AA

Poisons Act 1964

STOCKFEED MANUFACTURER'S PERMIT

This permit is granted to .....and authorizes him to sell by retail on behalf of.....
.....to any person producing the written order of a veterinary surgeon such mixture containing the following Fourth Schedule drugs as may be specified in the order, and within the limits as to quantity and composition set out in the order:

Fourth Schedule drugs to which this permit applies—

.....
.....

This permit is issued subject to the following conditions:—

(1) the mixture will be stored at and sold from premises situated at.....
.....;

(2) .....
.....
.....

Dated at Perth....., 19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 11AB

Poisons Act 1964

APPLICATION FOR STOCKFEED MANUFACTURER'S PERMIT

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
*Perth, WA 6000.

I, Mr. / Mrs. / Miss .....
(Full Name)

hereby apply on behalf of.....for a permit to sell by retail upon the written order of a veterinary surgeon mixtures of stockfeed containing the following Fourth Schedule drugs:

.....
.....
.....

In support of this application I declare that—

(1) the mixtures will be stored at and sold from premises situated at.....
.....;

(2) .....
.....
.....

Date.....

.....
Signature of Applicant.

Fee $.....with application

---

*The present address is 60 Beaufort Street, Perth.

Form 12

Poisons Act 1964

NOTICE OF APPEAL UNDER SECTION

IN the Court of Petty Sessions

at .....

BETWEEN

.....Appellant

and

.....Respondent

TAKE NOTICE that pursuant to the provisions of section 29 of the *Poisons Act 1964*, I intend to appeal to the Magistrate of the abovenamed Court against your (a).....on the.....day of.....19..... (b) .....
.....
.....

Dated this.....day of.....19.....

.....
Appellant.

To the Executive Director, Public Health and Scientific Support Services
And to.....

(a) State whether refusal, cancellation, order, etc..
(b) Set out particulars of the decision from which you are appealing.

Form 13

Poisons Act 1964

POISONS PERMIT (DEPARTMENTAL AND HOSPITAL)

THIS permit is granted to .....and authorizes him to purchase on behalf of.....from a manufacturer or wholesale dealer—

(a) the poisons specified in the.....Schedules to the *Poisons Act 1964.*
(b) the following poisons—

.....
.....
.....

This permit is issued subject to the following conditions:—

(1) the poisons will be stored only at premises situated at .....
.....

(2) the poisons will not be resold unless the poisons referred to above have been purchased on behalf of a public hospital;

(3) the poisons will be used only for the following purposes:—

.....
.....
.....

(4) .....
.....
.....

Dated at Perth....., 19.....

Valid until 30 June, 19.....

.....
Executive Director, Public Health and Scientific Support Services.

Form 13A

Poisons Act 1964

APPLICATION FOR POISONS PERMIT (DEPARTMENTAL AND HOSPITAL)

To the Executive Director, Public Health and Scientific Support Services,
Health Department of Western Australia,
*Perth.

I, Mr. / Mrs. / Miss ..... (Full Name)

hereby apply on behalf of.....for a permit to purchase from a manufacturer or wholesale dealer—

(a) the poisons specified in the **1st, 2nd, 3rd, 4th, 6th, 7th, 8th Schedules to the *Poisons Act 1964*; or

(b) the following poisons:—

.....
.....
.....

In support of this application I declare that—

(1) the poisons will be stored only at premises situated at .....
.....

(2) ‡ the poisons will not be resold;

(3) the poisons will be used only for the following purposes:—

.....
.....
.....

(4) .....
.....
.....

Date.....

.....
Signature of Applicant.

---

*The present address is 60 Beaufort Street, Perth.
**Strike out whichever does not apply.
‡Strike out if permit is sought on behalf of a public hospital.

[*Appendix A amended in Gazettes 14 June 1967 pp. 1582-83; 22 September 1969 p. 2876; 3 May 1974 p. 1435; 5 October 1979 pp. 3085-86; 7 June 1985 p. 1941; 23 May 1986 p. 1716; 15 May 1987 p. 2121.*]

APPENDIX B

DRUG REGISTER OF DRUGS OF ADDICTION USED AND RECEIVED

Date on which Transaction was Effected	Person, Body or Firm to Whom Sold or Supplied, or from Whom Received		Amount Used Sold or Supplied	Balance	Prescription Number	Prescriber	Dispenser
	Name and Address	Amount Received					

APPENDIX C

POISONS AND HAZARDOUS SUBSTANCES REQUIRED TO BE LABELLED WITH FIRST AID MEASURES

Except where otherwise stated, a poison or hazardous substance in this Appendix includes any derivative, compound, preparation or admixture included in relation to that item in a Schedule in Appendix A to the Poisons Act.

ACEPHATE
ACETIC ACID
ACETIC ANHYDRIDE
ACETONE
ACIFLUORFEN
ACROLEIN
AKLOMIDE
ALACHLOR
ALDICARB
ALDRIN
ALKALINE SALTS
ALLIDOCHLOR
ALLOXYDIM
ALLYL ALCOHOL
AMETRYN
AMIDITHION
AMINES AND ORGANIC ANHYDRIDES used as curing agents for epoxy resins
2-AMINOBUTANE
AMINOCARB
4-AMINOPYRIDINE
AMITON
AMITRAZ
AMITROLE
AMMONIA
AMMONIUM BIFLUORIDE
AMMONIUM THIOCYANATE
ANILINE
ANTICOAGULANTS
ANTIMONY COMPOUNDS
ANTU
ARECOLINE
ARPRINOCID
ARSENIC COMPOUNDS
AVERMECTIN B1
AZAMETHIPHOS
AZINPHOS-ETHYL
AZINPHOS-METHYL
AZOBENZENE
AZOCYCLOTIN
BARBAN
BARIUM COMPOUNDS except for SULPHATE
BENDIOCARB
BENOMYL
BENQUINOX
BENSULIDE
BENTAZONE
BENTHIOCARB
BENZENE
BENZOYL PEROXIDE
5-BENZYLFUR-3-YLMETHYL (1^1R,3^1S.E) -2^1,2^1-DIMETHYL-3^1 (2-OXO-2,3,4,5-TETRAHYDRO-3-THIENYLIDINEMETHYL) -CYCLOPROPANE CARBOXYLATE
BHC
BINAPACRYL
BIOALLETHRIN
BLEACHES containing more than 4 per cent available CHLORINE
BORAX
BORIC ACID
BORON COMPOUNDS
BORON TRIFLUORIDE
BRODIFACOUM

BROMADIOLONE
BROMOFORM
BROMOPHOS
BROMOPHOS-ETHYL
BROMOXYNIL
BROTIANIDE
BRUCINE
BUNAMIDINE
BUTACARB
BUTHIDAZOLE
BUTOXYCARBOXIM
BUTOXY-2'-THIOCYANO-DIETHYL ETHER
BUTYNORATE
CACODYLIC ACID
CADMIUM COMPOUNDS
CALCIUM HYPOCHLORITE
CAMPHECHLOR
CAMPHOR
CAMPHORATED OIL
CAPTAFOL
CARBARYL
CARBENDAZIM
CARBOFURAN
CARBON BISULPHIDE
CARBON TETRACHLORIDE
CARBOPHENOTHION
ALPHA-CHLORALOSE
CHLORAMPHENICOL for animal use
CHLORDANE
CHLORDECONE
CHLORDIMEFORM
CHLORFENAC
CHLORFENETHOL
CHLORFENSON
CHLORFENVINPHOS
CHLORINE
CHLORMEQUAT
CHLORNIDINE
N-[5-CHLORO-4-[(4-CHLOROPHENYL)-CYANOMETHYL]-2-METHYLPHENYL]-2-HYDROXY-3,5-DIIODOBENZAMIDE in concentrations less than 5 per cent
5-CHLORO-3-METHYL-4-NITROPYRAZOLE
2-CHLORO-N-[(4-METHOXY-6-METHYL-1-3,5,-TRIAZIN-2-YL) AMINOCARBONYL] BENZENE SULPHONAMIDE (Chlorsulfuron)
CHLOROCRESOL
CHLOROFORM
ALPHA-CHLOROHYDRIN
CHLOROMETHIURON
CHLOROPHACINONE
(BETA-[4-CHLOROPHENOXY]-ALPHA-[1,1-DIMETHYLETHYL]1H-1,2,4-TRIAZOLE-1-ETHANOL]) (Triadimenol)
1-(4-CHLOROPHENOXY)-1-IMIDAZOL-1-YL-3,3-DIMETHYL-2-BUTANONE (Climbazole)
CHLOROPICRIN
CHLOROPROPYLATE
CHLOROTHALONIL
CHLORPYRIFOS
CHLORPYRIFOS-METHYL
CHLORTETRACYCLINE for animal use
CHLORTHIAMID
CHLORTHIOPHOS
CHROMATES
CHROMIUM TRIOXIDE
CLANOBUTIN
CLOFENTEZINE
CLOPYRALID
COPPER SULPHATE
COUMAPHOS
COUMARIN DERIVATIVES
COUMATETRALYL
4-CPA

CREOSOTE
CRESOLS
CROTON OIL
CROTOXYPHOS
CRUFOMATE
CYANATRYN
CYANAZINE
CYANIDES
CYANOACRYLIC ACID ESTERS
(ALPHA-CYANO-4-FLUORO-3-PHENOXY)-BENZYL-3-[2-(4-CHLORPHENYL)-2-CHLOROVINYL] -2,2-DIMETHYL CYCLOPROPANE-CARBOXYLATE (Flumethrin)
CYCLOHEXANONE PEROXIDE
CYFLUTHRIN
CYHALOTHRIN
CYHEXATIN
CYOMETRINIL
CYPERMETHRIN
CYTHIOATE
2,4-D
DAZOMET
2,4-DB
DDT
DELTAMETHRIN
DEMETON
DEMETON-S-METHYL
2,4-DES
DI-(METHOXYCARBONYL)-1-PROPEN-2-YL-DIMETHYL PHOSPHATE
DI-ALLATE
DIALIFOS
N,N-DIALLYLDICHLOROACETAMIDE
DIAZINON
1,2-DIBROMO-3-CHLOROPROPANE
DICAMBA
DICHLOFENTHION
DICHLOFLUANID
DICHLONE
O-DICHLOROBENZENE
P-DICHLOROBENZENE
DICHLOROETHYLENE
DICHLOROETHYL ETHER
1-[2-(2,4-DICHLOROPHENYL)-2-(2-PROPENLOXY) ETHYL-1H-IMIDAZOLE
1-[2-(2,4-DICHLOROPHENYL)-4-PROPYL-1,3,DIOXALAN-2-YLMETHYL]-1H1,2,4-TRIAZOLE (Propiconazole)
1-[[2-(2,4-DICHLOROPHENYL)-4-ETHYL-1,3-DIOXOLAN-2-YL] METHYL]-1H1,2,4,TRIAZOLE (Etaconazole)
3,6-DICHLOROPICOLINIC ACID
1,2-DICHLOROPROPANE
1,3-DICHLOROPROPENE
N-(3,4-DICHLOROPHENYL)-N^1-[2-(2^{11}SULFOXY-4^1-CHLORPHENOXY)-5CHLORPHENYL] UREA (SODIUM SALT)
DICHLORVOS
DICHROMATES
DICLOBUTRAZOL
DICLOFOP-METHYL
DICLORAN
DICOFOL
DICROTOPHOS
DIELDRIN
DIENOCHLOR
DIETHYLENE DIOXIDE (Dioxane)
DIFENACOUM
DIFENZOQUAT
2,3-DIHYDRO-5,6-DIMETHYL-1,4-DITHIIN-1,1,4,4 TETRAOXIDE (Dimethipin)
DIMEFOX
DIMETHIRIMOL
DIMETHOATE
DIMETHYL FORMAMIDE
DIMETHYL SULPHOXIDE

2-(2^1,4^1-DIMETHYL-PHENYLIMINO)-3-METHYL-4-THIAZOLINE (Cymiazole)
DIMETILAN
DIMETRIDAZOLE
DINITRAMINE
DINITROCRESOLS
DINITROPHENOLS
DINOCAP
DINOSEB
DIOXACARB
DIOXATHION
DIPHACINONE
DIPHENAMID
DIQUAT
DISTILLATE
DISULFIRAM
DISULFOTON
DITHIANON
DITHIOCARBAMATES
$3,3^1$-DI-(TRIFLUROMETHYL)-$4,4^1$-DICHLORO-N,N^1-DIPHENYL UREA
DIUREDOSAN
DSMA
DNOC
DODINE
ENDOSULFAN
ENDOTHAL
ENDRIN
EPICHLOROHYDRIN
EPOXY RESINS LIQUID
EPTC
ETACONAZOLE
ETHEPHON
ETHER
ETHIOFENCARB
ETHION
ETHOATE-METHYL
ETHOFUMESATE
ETHOPROPHOS
ETHOXYQUIN
ETHYL BROMIDE
ETHYLENE CHLOROHYDRIN
ETHYLENE DIBROMIDE
ETHYLENE DICHLORIDE
ETHYLENE GLYCOL
ETHYLENE OXIDE
ETHYL FORMATE
ETRIDAZOLE
EUCALYPTUS OIL
FAMPHUR
FENAMINOSULF
FENAMIPHOS
FENARIMOL
FENAZAFLOR
FENBUTATIN OXIDE
FENCHLORPHOS
FENITROTHION
FENOPROP
FENSON
FENSULPHOTHION
FENTHION
FENTHION-ETHYL
FENVALERATE
FERBAM
FERRICYANIDES
FERROCYANIDES
FLAMPROP-METHYL
FLUAZIFOP-BUTYL
FLUCHLORALIN
FLUCYTHRINATE
FLUORACETAMIDE

FLUORIDES
FLUOROACETIC ACID
FORMALDEHYDE
FORMETANATE
FORMIC ACID
FORMOTHION
FOSPIRATE
FURALAXYL
GLUTARALDEHYDE
GLYPHOSATE
GUAZATINE
HALOFUGINONE
HALOXON
HCB
HEPTACHLOR
HEXACHLOROPHANE when included in the sixth schedule
HEXAZINONE
HYDRAZINE
HYDROCARBONS, LIQUID
HYDROCHLORIC ACID
HYDROFLUORIC ACID (excluding its salts)
HYDROGEN PEROXIDE
HYDROQUINONE
HYDROSILICOFLUORIC ACID
IMIDOCARB DIPROPIONATE
IODINE
IODOFENPHOS
IODOPHORS
IOXYNIL
IRON COMPOUNDS in solid and liquid preparations for animal treatment
ISOCARBOPHOS
ISOCYANATES, FREE ORGANIC
ISOFENPHOS
ISOPROPYL-N-(3-N-ETHYL-N-PHENYLCARB-AMOYLOXY) PHENYL-CARBAMATE (Phenisopham)
KEROSENE
LAURYL ISOQUINOLINIUM BROMIDE
LEAD ARSENATE
LEAD COMPOUNDS
LEPTOPHOS
LEVAMISOLE
LINDANE
MALDISON
MANCOZEB
MANEB
MAZIDOX
MCPA
MCPB
MEBENDAZOLE
MECARBAM
MECLOFENAMIC ACID
MECOPROP
MENAZON
MEPIQUAT
MERCURIC CHLORIDE
MERCURIC IODIDE
MERCURIC NITRATE
MERCURIC OXIDE
MERCURIC POTASSIUM OXIDE
MERCURIC THIOCYANATE
MERCUROUS CHLORIDE
MERCURY, METALLIC
MERCURY, ORGANIC COMPOUNDS
METACRESOLSULPHONIC ACID and FORMALDEHYDE CONDENSATION PRODUCT
METALAXYL
METALDEHYDE
METAXANINE
METHABENZTHIAZURON
METHACRIFOS

METHAM
METHAMIDOPHOS
METHAZOLE
METHIDATHION
METHIOCARB
METHOMYL
O-2-METHYOXYCARBONYL, PROP-1-ENYL-O, O-DIMETHYLPHOSPHOROTHIOATE (Methacrifos)
METHOXYCHLOR
METHYL ALCOHOL
METHYL BROMIDE
N-METHYL CARBAMATES
METHYL CHLORIDE
METHYL ETHYL KETONE
METHYL ETHYL KETONE PEROXIDE
METHYL ISOAMYL KETONE
METHYL ISOBUTYL KETONE
METHYL ISOTHIOCYANATE
METHYL SALICYLATE (Liquid)
METHYLATED SPIRITS
METHYLENE BISTHIOCYANATE
METHYLENE CHLORIDE
3-(METHYLSULPHONYL) BUTANON-O-METHYLCARBAMOXLOXIM (Butoxycarboxim)
METIRAM
METOLACHLOR
METRIBUZIN
MEVINPHOS
MEZINEB
MINERAL TURPENTINE
MIPAFOX
MIREX
MOLINATE
MONOCROTOPHOS
MSMA
NAA
NABAM
NALED
NAPHTHALENE
NAPHTHALOPHOS
NAPTALAM
NARASIN
NICOTINE
NIMIDANE
NITHIAMIDE
NITRIC ACID
NITROBENZENE
NITROOXYNIL
NITROPHENOL
NORBORMIDE
2-N-OCTYL-4-ISOTHIAZOLIN-3-ONE (Octhilinone)
OESTRADIOL 17-BETA
OFURACE
OIL OF TURPENTINE
OLAQUINDOX
OMETHOATE
ORGANO-PHOSPHORUS COMPOUNDS
OXADIAZON
OXALIC ACID
OXAMYL
OXFENDAZOLE
OXYCARBOXIN
OXYFLUORFEN
OXYTETRACYCLINE for animal use
OXYTHIOQUINOX
PARAQUAT
PARATHION
PARATHION-METHYL
PARBENDAZOLE
PEBULATE

PENCONAZOLE
PENDIMETHALIN
PENTACHLORONITROBENZENE
PENTACHLOROPHENOL
PERACETIC ACID
PERFLUIDONE
PERMANGANATES
PETROL
PHENKAPTON
PHENOLS
PHENYLENE DIAMINES
O-PHENYLPHENOL
PHORATE
PHOSALONE
PHOSFOLAN
PHOSMET
PHOSPHAMIDON
PHOSPHIDES METALLIC
PHOSPHONIC ACID (over 10 per cent content)
PHOSPHORIC ACID
PHOSPHORUS, YELLOW
PHOXIM
PICRIC ACID
PINDONE
PIPEROPHOS
PIRIMICARB
PIRIMIPHOS-ETHYL
PIRIMIPHOS-METHYL
POLY (HEXAMETHYLENE BIGUANIDE) HYDROCHLORIDE
POLYCHLORINATED BIPHENYLS
POTASSIUM BROMATE
POTASSIUM CYANATE
POTASSIUM HYDROXIDE
POTASSIUM SULPHIDE
PROCHLORAZ
PROFENOFOS
PROGESTERONE
PROMACYL
PROMECARB
PROMETRYN
PROPACHLOR
PROPANIL
PROPARGITE
PROPETAMPHOS
PROPICONAZOLE
PROPIONIC ACID
PROPOXUR
PROTHIOPHOS
PRYNACHLOR
PYRAZOPHOS
PYRETHRINS
N-3-PYRIDYLMETHYL-N^1-P-NITROPHENYLUREA (Pyrinuron)
PYRITHIONE ZINC
QUATERNARY AMMONIUM COMPOUNDS
QUINTOZENE
SALICYLANILIDE
SALINOMYCIN
SCHRADAN
SECBUMETON
SELENIUM COMPOUNDS
SETHOXYDIM
SODIUM BIFLUORIDE
SODIUM BROMATE
SODIUM CHLORATE
SODIUM DICHLOROISOCYANURATE
SODIUM HYDROGEN SULPHATE
SODIUM HYDROXIDE
SODIUM HYPOCHLORITE
SODIUM NITRITE

SODIUM SULPHIDE
SODIUM TRICHLOROISOCYANURATE
STRYCHNINE
STYRENE
SULFALLATE
SULFOTEP
SULPHAMIC ACID
SULPHURIC ACID
SULPROPHOS
2, 4, 5-T
2, 3, 6-TBA
TCMTB
TDE
TEMEPHOS
TEPP
TERBUMETON
TERBUTHYLAZINE
TERBUTRYN
TERPENES, CHLORINATED
TESTOSTERONE for animal use
TETRACHLORETHANE
TETRACHLOROETHYLENE
TETRACHLORVINPHOS
TETRACYCLINE for animal use
TETRADIFON
TETRAMISOLE
THALLIUM
THIAZAFLURON
THIOBENCARB
THIODICARB
THIOFANOX
THIOMETON
THIOUREA
THIRAM
TIN, ORGANIC COMPOUNDS
O-TOLIDINE
TOLUENE
TRI-ALLATE
TRIADIMEFON
TRIADIMENOL
TRIAZBUTIL
S,S,S-TRIBUTYLPHOSPHOROTHIOATE
TRICHLOROACETIC ACID and its alkali salts
1,1,1-TRICHLOROETHANE
TRICHLOROETHYLENE
TRICHLOROISOCYANURIC ACID
TRICLOROPHENOL
TRICHLORPHON
TRICLOPYR
TRIDEMORPH
TRIETAZINE
TRIETHYL PHOSPHATE
TRIFLUOROMETHANE SULPHONIC ACID
TURPENTINE OIL
VAMIDOTHION
VERNOLATE
WARFARIN
WHITE SPIRIT
XYLENE
XYLENOLS
ZINC CHLORIDE
ZINC P-PHENOLSULPHONATE
ZINC SULPHATE
ZINEB
ZIRAM

[*Appendix C substituted in Gazette 7 December 1979 pp. 3799-3802; amended in Gazettes 16 July 1982 p. 2727; 28 January 1983 pp. 341-42; 15 March 1985 pp. 941-42; 23 May 1986 pp. 1716-19; erratum 20 June 1986 pp. 2050-52.*]

APPENDIX D

POISONS AND HAZARDOUS SUBSTANCES REQUIRED TO BE LABELLED WITH A WARNING STATEMENT

WS 1 "Avoid contact with the skin and eyes"

Acifluorfen
Avermectin B1
Benzoyl Peroxide except when included in the Second, Third or Fourth Schedule.
Alpha—(2-chlorophenyl)-alpha-(4-chlorophenyl)-5-pyrimidine-methanol
Alpha-cyano-3-phenoxybenzyl 2,2-dimethyl-3-3 (2,2 dichlorovinyl) cyclopropane
5-Chloro-3-methyl-4-nitropyrazole
Chlorthiophos
Clopyralid
Cyclohexanone Peroxide
Cypermethrin
1-[2 (2,4-Dichlorophenyl)-2-(2-propenyloxy) ethyl]-1 H imidazole
3,6-Dichloropicolinic acid
0,0-diethyl-0-(2,4,5-dichloro methyltio phenyl) thionophosphate
Ethephon
Formaldehyde
Guazatine
Hydrochloric acid
Hydrofluoric acid and hydrosilicolfluoric acid, their salts and all substances containing any of these compounds when included in the Fifth Schedule.
Hydroquinone
Isofenphos
Mepiquat
Metacresolsulphonic acid and formaldehyde condensation product.
Methylene bisthiocyanate
N-(3,4-dichlorphenyl)-N^3- 2-(sulfoxy-4^1-chlorophenoxy)-5-chlorphenyl urea (sodium salt)
Nitric Acid
Ofurace
ortho-Phenylphenol
Oxalic acid and metallic oxalates
Oxfendazole
Oxythioquinox
Phenol and any homologue of phenal boiling below 220°.
Phosphonic acid
Phosphoric acid
Prothiophos
Sethoxydim
Sodium chlorate
Sulphuric acid
Thiobencarb
Zinc chloride

WS 2 "Avoid contact with skin and eyes and avoid breathing its dust, vapour or spray mist.

Acrolein
Amidothion
Aminocarb
2-Amino-5-diethylamino toluene
2-Amino-5-N-ethyl-N-(B hydroxy ethyl) amino toluene
2-Amino-5-N-ethyl-N-(B methane sulphonamide ethyl) amino toluene
2-Amino-5-N-ethyl-N B methoxyethyl amino-toluene di-p-toluene
Aniline
Arsenic, organic compounds when prepared for use as herbicides and defoliants.
Azobenzene
Azocyclotin
Benzene
Beryllium
BHC
Biothionol for the treatment of animals.
Bromophos
Bromophos-ethyl
2-Butoxy-2^1-thiocyano-diethyl ether
4-n-Butyl-4H-1,2,4-triazole
Camphechlor

Carbaryl except when included in the Second Schedule.
Carbon disulphide
Chlordane
Chlordimeform
Chlorfenethol
Chlorinating Compounds and Bleaches
Chloropicrin
Chlorpyrifos
Chromate and dichromates of alkali metals and ammonium.
Chromic acid
Crotoxyphos
Crufomate
Cyanoacrylic Acid Esters
(S)-alpha-cyano-m-phenoxybenzyl (1R,3R)-3-(2,2-dibromovinyl)-2-dimethylcyclopropane carboxylate
Cyhalothrin
DDT except for human therapeutic use.
Deltamethrin
Demeton-O-methyl
Demeton-S-methyl
Dialifos
Diazinon
Dibromochloropropane
Dichlofenthion
Dichloroethyl ether
Dichloroethylene
Dichloroisocyanurates
Dichlorvos except when included in Schedule 5.
Diethylene dioxide
N,N-Diethyl-p-phenylene diamine
2,3,-Dihydro-5,6-dimethyl-1,4-dithiin-1,1,4,4-tetraoxide (Dimethipin)
Dimethanonaphthalene and all substitution and/or additional products thereof.
Dimethoate
1,3-Di (methoxycarbonyl)-1-propen-2 yl-dimethyl phosphate
Dimethyl sulphoxide
Dimetilan
Dinitrocresols and their homologues except for therapeutic use.
Dinitrophenols and their homologues except for therapeutic use.
DSMA
Endosulfan
Endothal
Epichlorohydrin
Epoxy resins liquid, and all amines and organic anhydrides used as curing agents for epoxy resins.
Ether solvent
Ethoate-methyl
Ethofumesate
Ethyl bromide
Ethylene dibromide
Ethylene oxide
Famphur
Fenchlorphos
Fenitrothion
Fenthion
Formic acid
Formothion
Glyphosate
Halofuginone
Heptachlor
Hydrazine
Isocyanates, free organic
Lindane except when included in the Second Schedule.
Maldison except when included in the Second Schedule.
Menazon
Metham-sodium
Methiocarb
Methyl alcohol except in methylated spirit.
Methyl bromide
Methyl chloride

Methylene chloride
Methyl isothiocyanate
1-(B-Methyl sulphonamido ethyl)-2-amino-3-N,N-diethylamino benzene.
Naled
Naphthalophos
Nicotine and its salts except in tobacco or chewing tablets.
Nimidane
Nitrobenzene
2-n-Octyl-4-Isothiazolin-3-one
Omethoate
Pentachlorophenol
Peracetic acid
Phenkapton
Phosalone
Phosmet
Phosphides, metallic
Poly (hexamethylene biguanide) hydrochloride
Promecarb
Propachlor
Propetamphos
Propoxur except when included in the Second Schedule.
Selenium, compounds of, in preparations other than for human therapeutic use.
Styrene
TDE
Temephos
Terbuthylazine
Terpenes, chlorinated
Tetrachloroethylene except for therapeutic use.
Tetradifon
Thiometon
Toluene
Triazbutyl
S,S,S-Tributylphosphorothiolate
Trichloroethylene except when specially prepared for medical purposes.
Trichloroisocyanuric acid
Trichlorophenol
Trichlorphon
Tridemorph
Triethyl phosphate
Trifluoromethane sulphonic acid (10 per cent or less)
Vamidothion
Xylene

WS 3 "Warning—this substance is caustic—avoid contact with the skin and eyes".
Acetic acid 80 per cent and over
Potassium hydroxide
Potassium sulphide
Sodium hydroxide
Sodium sulphide

WS 4 "Flammable".
Acrolein
Benzene
Carbon disulphide
dichloroethylene
Diethylene dioxide
Ether solvent
Ethylene oxide
Hydrocarbons, liquid, distilling under 300°C when tested according to method D86-61 of the American Society for Testing Materials.
Kerosene
Methyl alcohol
Methylated spirit
Mineral turpentine
Oil of turpentine
Petrol
Toluene
White spirit
Xylene

WS 5 "Avoid contact with food".
Arsenic, organic compounds, when prepared for use as herbicides or defoliants.
DSMA
Endothal
Insecticide preparations

WS 6 "Wear protective gloves when mixing or using".
Cyhalothrin
Deltamethrin
Dimethyl sulphoxide
Halofuginone
Liquid epoxy resins and all amines and organic anhydrides used as curing agents for epoxy resins.
Phenols
2-n-Octyl-4-isothiazolin-3-one
Terbuthylazine
Triazbutyl

WS 7 "Do not use with other asthma sprays or remedies and avoid frequent and prolonged use except on medical advice".
Asthma sprays containing adrenaline, natural or synthetic, its salts, noradrenaline and substances structurally derived therefrom by substitution in the amine group, their salts.

WS 8 "Should not be taken for periods longer than four weeks except on medical advice".
8-Hydroxyquinoline, its derivatives and their salts when prepared for internal use.

WS 9 "Warning—this product contains ingredients which may cause skin irritation of certain individuals and a preliminary test according to accompanying direction should first be made. This product must not be used for dyeing the eyelashes or eyebrows; to do so may be injurious to the eye".
Amines, aromatic, including phenylene diamine, toluene diamine and other aromatic amines when used in hair dyes.

WS 10 "Warning—milk from animals treated with this preparation is unfit for human consumption and must be discarded for (to be stated) hours following the cessation of treatment to ensure that the milk is free from residues".
Antibiotic preparations for intramammary treatment of animals.

WS 11 "Warning—should not be used for human beings. For animal treatment only".
CHLORTETRACYCLINE in preparations for topical application to animals for ocular use only.
NEOMCYIN in preparations for topical application to animals for ocular use only.
OXYTETRACYCLINE in prepartions for topical application to animals for ocular use only.
SULPHAQUINOXALINE when packed and labelled for use as a coccidiostat in poultry except preparations containing 200 mg/kg or less of sulphaquinoxaline.
TETRACYCLINE in preparations for topical application to animals for ocular use only.
TESTOSTERONE CYPIONATE, DIPROPIONATE, ENANTHATE and PROPIONATE in preparations for the treatment of animals.

WS 12 "In the directions for use on the label it must be clearly stated that the concentration of antibiotics in the feed as given to stock should not exceed 100 mg/kg of the active antibiotic principle".
ANTIBIOTIC PREMIXES for growth promotion purposes.

WS 13 "(1) Do not use in food cupboards;
(2) Do not use in nurseries and sick rooms where people may be continuously exposed".
DICHLORVOS when impregnated in plastic resin strip material containing 20 per cent or less dichlorvos.

WS 14 "For external washing only. Rinse skin thoroughly after use".
S-(2-chloro-1-phthalimidoethyl)-0,0-diethylphosphorodithioate
Hexachlorophane in preparations for skin cleansing purposes containing 3 per cent or less of hexachlorophane.

WS 15 "Vapour is harmful to health on prolonged exposure: use only in a well ventilated area".
Benzene
Carbon tetrachloride
Methylene chloride in paint or lacquer removers
Tetrachlorethane

WS 16 "Unless adequately fired utensils glazed with this preparation must not be used as containers for food or beverages; to do so may cause lead poisoning".
Glazing preparations containing lead compounds.

[*WS 17 deleted.*]

WS 18 "Highly reactive oxidizing chlorine compound may cause fire or explosion or produce severe burns:
Do not allow to get damp.
Store under cover in a dry, clean, well-ventilated place,
Do not allow to come in contact with acids, reducing agents, ammonium compounds, wood shavings, saw dust, papers, fabric, petrol, kerosene or other combustible material."
Dry chlorinating compounds and bleaches containing 10 per cent or more chlorine.
Trichloroioscyanuric acid

WS 19 "Warning—this product contains ingredients which may cause skin irritation in certain individuals. Avoid contact with skin and eyes and avoid breathing its dust".
Captafol
Metolachlor

WS 20 "Whenever liquid concentrate is handled, always wear an approved respirator, polyethylene gloves, rubber boots and goggles.
During application always wear a respirator if exposed to vapour particularly when working in enclosed areas such as a glasshouse".
1,3-Dichloropropene

WS 21 "An anticholinesterase compound" (to appear immediately below the approved name or the list of declared contents on the label)".
Carbaryl except when included in the Second Schedule.
Dialifos
Dichlorvos in aerosol packs containing 10 g or less dichlorvos.
Isofenphos
Maldison except when included in the Second Schedule.
Methacrifos
Organophosphorus and carbamate compounds for pesticidal use except—
(a) di-allate, tri-allate, dazomet, mancozeb, maneb, metiram, propineb, thiram, zineb and ziram;
(b) impregnanted plastic resins, strips or granules and aerosol packs for household use.
Propetamphos
Propoxur
Thiobencarb
Thiodicarb

WS 22 "This substance is strongly alkaline; Avoid contact with skin and eyes".
Alkaline salts

WS 23 "May be fatal if inhaled or swallowed".
Oxamyl

WS 24 "When mixing or spraying, wear PVC neoprene gloves, hat, waterproof coat and trousers (worn outside rubber boots) and a respirator. Wash protective clothing daily after use."
Dialifos
Isopropyl-N-(3-N-ethyl-N-phenyl-carbamoyloxy) phenylcarbamate
Oxamyl

WS 25 "Wear goggles when using as a fine spray".
Acifluorfen
Metolachlor
Thiobencarb

WS 26 "Warning—this medication may be dangerous when used in large amounts or for a long period: or
Caution—this preparation is for the relief of minor and temporary ailments and should be used strictly as directed. Prolonged use without medical supervision could be harmful."
Aspirin
Paracetamol
Salicylamide

WS 27 "For use under Medical Supervision Only".
Aspirin in sustained release preparations containing 650 mg or more of aspirin.
Salsalate

WS 28 "Attacks eyes—protect eyes when using and avoid contact with skin".
Methylethyl ketone peroxide
Narasin

WS 29 "Forms dangerous gas near radiators or naked flames—no smoking".
Methylene Chloride in paint or lacquer removers.

WS 30 "Do not use on broken skin, wash hands thoroughly after use".
Lead compounds in hair cosmetics.

WS 31 "Use of this product is not necessary in areas supplied with fluoridated water".
Sodium fluoride in preparations for human ingestion containing 2.2 mg or less of sodium fluoride per dosage unit.

WS 32 "This substance is highly corrosive. Avoid contact with the skin and avoid breathing its vapour. Contact with the eyes even for short periods can cause blindness".
Hydrofluoric acid except when included in the Fifth Schedule.
Hydrosilicofluoric acid except when included in the Fifth Schedule.
Trifluoromethane sulphonic acid (over 10 per cent)

WS 33 Harmful if inhaled, wear a cloth dust mask or disposable paper dust mask during handling or mixing.
Benomyl.

WS 34 Not for Therapeutic Use.
Dimethyl sulphoxide when not packed and labelled for therapeutic use.

WS 35 Warning—should not be mixed with other medication except on veterinarian's advice.
Dimethyl sulphoxide when packed and labelled for use on animals.

WS 36 Warning—use of this preparation during pregnancy should be avoided.

WS 37 "Not for use in food producing animals on a flock or herd basis".
Chloramphenicol

WS 38 "WARNING—causes birth defects".
Etretinate
Isotretinoin
Thalidomide

[*Appendix D substituted in Gazette 7 December 1979 pp. 3799-3805; amended in Gazettes 16 July 1982 pp. 2727-28; 28 January 1983 p. 342; 15 March 1985 p. 942; 23 May 1986 pp. 1719-20; erratum 20 June 1986 pp. 2053-54; amended in Gazette 11 July 1986 p. 2339.*]

APPENDIX E

POISONS REQUIRED TO BE LABELLED WITH A WARNING STATEMENT AND FIRST AID MEASURES

Except where otherwise stated a substance in this Appendix includes any derivative, compound, preparation or admixture included in relation to that substance in the Seventh Schedule to the Act.

The label for any poison specified in this Appendix shall contain the following warning statement and first aid statement appropriate to the particular item—

	WARNING STATEMENT	FIRST AID STATEMENT
ACROLEIN	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. FLAMMABLE.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. Remove from contaminated area. Apply artificial respiration if not breathing. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
ALDICARB		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
ALLYL ALCOHOL		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.

	WARNING STATEMENT	FIRST AID STATEMENT
AMINOCARB	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
4-AMINOPYRIDINE		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. If skin contact occurs, remove contaminated clothing and wash skin thoroughly.
AMITON	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
ANTU		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
APRINOCID		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
ARSENIC		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
AZINPHOS ETHYL	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
AZINPHOS METHYL	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
BENDIOCARB	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
BENZENE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. FLAMMABLE. VAPOUR IS HARMFUL TO HEALTH ON PROLONGED EXPOSURE. USE ONLY IN A WELL VENTILATED AREA.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, do NOT induce vomiting. Give a glass of water. If skin contact occurs, remove contaminated clothing and wash skin throughly. Remove from contaminated area. Apply artificial respiration if not breathing.
BRODIFACOUM		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
BROMADIOLONE		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
CAMPHECHLOR	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Avoid giving milk or oils. If skin contact occurs, remove contaminated clothing and wash skin thoroughly.
CAPTAFOL	WARNING—THIS PRODUCT CONTAINS INGREDIENTS WHICH MAY CAUSE SKIN IRRITATION IN CERTAIN INDIVIDUALS. AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING ITS DUST. WEAR GOGGLES WHEN USING AS A FINE SPRAY.	If poisoning occurs, contact a doctor or Poisons Information Centre.

	WARNING STATEMENT	FIRST AID STATEMENT
CARBOFURAN	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
CARBON-TETRA-CHLORIDE	VAPOUR IS HARMFUL TO HEALTH ON PROLONGED EXPOSURE. USE ONLY IN A WELL VENTILATED AREA.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Avoid giving alcohol. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. Remove from contaminated area. Apply artificial respiration if not breathing. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
CARBOPHENO-THION	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
CHLORDIME-FORM	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
CHLORFEN-VINPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
CHLORINE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. Remove from contaminated area. Apply artifical respiration if not breathing. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
5-CHLORO-3-METHYL-4-NITROPYRA-ZOLE	AVOID CONTACT WITH THE SKIN AND EYES.	If poisoning occurs, contact a doctor or Poisons Information Centre.

	WARNING STATEMENT	FIRST AID STATEMENT
CHLORO-PICRIN	AVOID CONTACT WITH THE SKIN AND EYES.	If poisoning occurs, contact a doctor or Poisons Information Centre. If skin contact occurs remove contaminated clothing and wash skin thoroughly. Remove from contaminated area. Apply artificial respiration if not breathing. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
CHLOR-THIOPHOS	AVOID CONTACT WITH THE SKIN AND EYES. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
COUMAPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
CYANIDES	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If breathing, crush one amyl nitrite ampoule in handkerchief and hold under patient's nose from 1 or 2 seconds. Repeat up to 5 times at intervals of one minute. If not breathing, wipe patient's lips and apply artificial respiration.
CYHALOTHRIN	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. WEAR PROTECTIVE GLOVES WHEN MIXING OR USING.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
DELTA-METHRIN	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. WEAR PROTECTIVE GLOVES WHEN MIXING OR USING.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
DEMETON	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABLE).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin throughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
DEMETON-O-METHYL DEMETON-S-METHYL	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
DIALIFOS	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL). WHEN MIXING OR SPRAYING, WEAR P.V.C. OR NEOPRENE GLOVES, HAT, WATERPROOF COAT AND TROUSERS (WORN OUTSIDE RUBBER BOOTS) AND A FACE SHIELD. WASH PROTECTIVE CLOTHING DAILY AFTER USE.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
1,2-DIBROMO-3-CHLORO-PROPANE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
DICHLORVOS	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
DICROTOPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If the poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
DIMEFOX	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
1,3-DI-(METHOXY CARBONYL)-1-PROPEN-2-YL-DIMETHYL PHOSPHATE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
DIMETILAN	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
DINITRO-CRESOLS	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
DINITRO-PHENOLS	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
DIOXATHION	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
DISULFOTON	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atrophine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
ENDOTHION	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
ENDRIN		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (AFP) if available. Avoid giving milk or oils. If skin contact occurs, remove contaminated clothing and wash skin thoroughly.
ETHION	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
ETHOPROPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
ETHYLENE DIBROMIDE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. Remove from contaminated area. Apply artificial respiration if not breathing. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.

	WARNING STATEMENT	FIRST AID STATEMENT
FAMPHUR	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
FENAMINOSULF		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
FENAMIPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
FENSULFOTHION	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
FENTHIONETHYL	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
FLUCYTHRINATE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. WHENEVER LIQUID CONCENTRATE IS HANDLED, WEAR AN APPROVED RESPIRATOR, POLYETHYLENE GLOVES, RUBBER BOOTS AND GOGGLES. DURING APPLICATION ALWAYS WEAR A RESPIRATOR IF EXPOSED TO VAPOUR PARTICULARLY	

	WARNING STATEMENT	FIRST AID STATEMENT
FLUCYTHRINATE *(cont.)*	WHEN WORKING IN ENCLOSED AREAS SUCH AS A GLASSHOUSE. WHEN MIXING OR SPRAYING WEAR P.V.C. OR NEOPRENE GLOVES, HAT, WATERPROOF COAT AND TROUSERS (WORN OUTSIDE RUBBER BOOTS) AND A FACE SHIELD. WASH PROTECTIVE CLOTHING DAILY AFTER USE.	
FLUORACETAMIDE		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
FLUOROACETIC ACID	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
FORMETENATE	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter of an hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
HALOFUGINONE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. WEAR PROTECTIVE GLOVES WHEN MIXING OR USING.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
ISOCARBOPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
ISOFENPHOS	AVOID CONTACT WITH THE SKIN AND EYES. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
LEPTOPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
MAZIDOX	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
MECARBAM	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
MERCURIC CHLORIDE		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. If skin contact occurs, remove contaminated clothing and wash skin thoroughly.
METHAMIDOPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
METHIDA-THION	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
METHOMYL	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
METHYL BROMIDE	AVOID CONACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. Remove from contaminated area. Apply artificial respiration if not breathing.
MEVINPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine table every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
MIPAFOX	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
MIREX		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Avoid giving milk or oils. If skin contact occurs, remove contaminated clothing and wash skin thoroughly.

	WARNING STATEMENT	FIRST AID STATEMENT
MONOCROTO-PHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
NAPHTHALO-PHOS	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
NICOTINE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
NIMIDANE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre.
OMETHOATE	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
ORTHO-TOLI-DINE		If poisoning occurs, contact a doctor or Poisons Informaton Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available.
OXAMYL	MAY BE FATAL IF INHALED OR SWALLOWED. WHEN MIXING OR SPRAYING, WEAR P.V.C. OR NEOPRENE GLOVES, HAT, WATERPROOF COAT AND TROUSERS (WORN OUTSIDE RUBBER BOOTS) AND A FACE SHIELD. WASH PROTECTIVE CLOTHING DAILY AFTER USE.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
OXYFLUOROFEN		If poisoning occurs, contact a doctor or Poisons Information Centre. If in eyes, hold eyes open, flood with water for a least 15 minutes and see a doctor.
PARAQUAT		If poisoning occurs get to a doctor or hospital quickly. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
PARATHION	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above. If in eyes, hold eyes open, flood with water for at least 15 minutes and see a doctor.
PARATHION METHYL	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
PHENKAPTON	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST, VAPOUR OR SPRAY MIST.	If poisoning occurs, contact a doctor or Poisons Information Centre. Remove from contaminated area. Apply artificial respiration if not breathing.
PHORATE	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
PHOSFOLAN	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATEDLY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
PHOSPHA-MIDON	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs, If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
PROMECARB	AVOID CONTACT WITH SKIN AND EYES AND AVOID BREATHING DUST VAPOUR OR SPRAY MIST. AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by the skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
PROTHOATE	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
SCHRADAN	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor of Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
STRYCHNINE		If poisoning occurs, contact a doctor or Poisons Information Centre. Give activated charcoal and keep patient quiet, in a dark place if possible.
SULFALLATE		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed induce vomiting. Use Ipecac Syrup (APF) if available.
SULFOTEP	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.

	WARNING STATEMENT	FIRST AID STATEMENT
TEPP	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
TERBUPHOS	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OF THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
TETRACHLOROETHANE	VAPOUR IS HARMFUL TO HEALTH ON PROLONGED EXPOSURE. USE ONLY IN A WELL VENTILATED AREA.	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Avoid giving milk or oils. Avoid giving alcohol. If skin contact occurs, remove contaminated clothing and wash skin thoroughly. Remove from contaminated area. Apply artificial respiration if not breathing.
THALLIUM		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available.
THIONAZIN	AN ANTICHOLINESTERASE COMPOUND (TO APPEAR IMMEDIATELY BELOW THE APPROVED NAME OR THE LIST OF DECLARED CONTENTS ON THE LABEL).	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available. Give one Atropine tablet every quarter hour until dryness of the mouth occurs. If poisoned by skin absorption, remove contaminated clothing and wash skin thoroughly. Give Atropine tablets as above.
TRIAZBUTIL		If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, induce vomiting. Use Ipecac Syrup (APF) if available.
TRICHLOROISOCYANURIC ACID	HIGHLY REACTIVE OXIDIZING CHLORINE COMPOUND MAY CAUSE FIRE OR EXPLOSION OR PRODUCE SEVERE BURNS. DO NOT ALLOW TO GET DAMP. STORE UNDER COVER IN A DRY, CLEAN,	If poisoning occurs, contact a doctor or Poisons Information Centre. If swallowed, do NOT induce vomiting. Give a glass of water. If skin contact occurs, remove contaminated clothing and wash skin thoroughly.

	WARNING STATEMENT	FIRST AID STATEMENT
TRICHLORO-ISOCYANURIC ACID *(cont.)*	WELL VENTILATED PLACE. DO NOT ALLOW TO COME IN CONTACT WITH ACIDS, REDUCING AGENTS, AMMONIUM COMPOUNDS, WOOD SHAVINGS, SAW DUST, PAPERS, FABRIC, PETROL, KEROSENE OR OTHER COMBUSTIBLE MATERIAL.	

[*Appendix E substituted in Gazette 15 March 1985 pp. 943-54; amended in Gazettes 28 February 1986 p. 617; 23 May 1986 p. 1720.*]

APPENDIX F

[*Appendix F repealed in Gazette 1 August 1986 p. 2739.*]

APPENDIX G

Form No.		Annual Fee $
1.	Licence to Procure, Manufacture and Supply Poisons (other than drugs of addiction) by Wholesale Dealing	65
2.	Licence to Procure, Manufacture and Supply by Wholesale Dealing Drugs of Addiction	65
3.	Pharmaceutical Chemist's Licence to Sell Poisons	25
4.	Licence to Sell by Retail Poisons Specified in the Sixth Schedule	15
5.	Licence to Sell by Retail, Poisons Specified in the First, Second or Sixth Schedule	25
6.	Licence to Sell by Retail, Poisons Specified in the Seventh Schedule	15
6B.	Poisons Permit (Distribution of Samples)	25
7.	Poisons Permit (Industrial)	15
8.	Poisons Permit (Educational, Advisory or Research)	No fee
9.	Licence to Hawk, Peddle or Distribute Poisons	15
10.	Classification of a New Drug	No fee
11.	Permit to Supply for Veterinary Use the preparations referred to in Regulation 39 (2)	15
11AA.	Stockfeed Manufacturer's Permit	25
13.	Poisons Permit (Departmental and Hospitals) The fee for renewal is the same as for the original.	No fee

[*Appendix G substituted in Gazette 12 April 1985 pp. 1285-86.*]

APPENDIX H

FOURTH SCHEDULE DRUGS REFERRED TO IN REGULATION 39 (1)

Except where otherwise stated, a substance in this Appendix does not include any derivative but does include any compound, preparation or admixture included in relation to that substance in the Fourth Schedule to the Act. Where a method of application or of administration is stated in relation to a substance, the entry in this Appendix in relation to that substance applies only to compounds, preparation or admixtures of that substance which have been prepared for that method of application or administration—

Acepromazine Maleate.
Antihistamines.
Apomorphine.
Benzyl penicillin for parenteral injection.
Chlorpromazine (but not to be supplied for use on horses).
Procaine penicillin for parenteral injection.
Streptomycin.

[*Appendix H substituted in Gazette 8 February 1985 p. 520; erratum 19 April 1985 p. 1409.*]

APPENDIX I

[*Appendix I repealed in Gazette 23 May 1986 p. 1720.*]

APPENDIX J

(reg. 35A)

THIRD SCHEDULE POISON SALES TO BE RECORDED

AMYL NITRITE;

BUTYL NITRITE;

CHLORAL HYDRATE, when included in the Third Schedule;

CLOTRIMAZOLE, when included in the Third Schedule;

ECONAZOLE, when included in the Third Schedule;

ISOCONAZOLE, when included in the Third Schedule;

MICONAZOLE, when included in the Third Schedule.

[*Appendix J inserted in Gazette 20 September 1985 p. 3743; amended in Gazettes 23 May 1986 p. 1721; 23 January 1987 p. 187.*]

APPENDIX K

(reg. 21A)

POISONS REQUIRED TO BE LABELLED WITH A WARNING STATEMENT RELATING TO DRIVING A MOTOR VEHICLE AND OPERATING MACHINERY

AMITRIPTYLINE.

AZATADINE.

BACLOFEN.

BARBITURIC ACID and its derivatives.

BENZTROPINE.

BROMPHENIRAMINE.

BUCLIZINE.

BUPRENORPHINE.

CHLORAL HYDRATE when included in the 3rd or 4th Schedule.

CHLORDIAZEPOXIDE and other substances structurally derived from benzodiazepine with ataractic properties when used for therapeutic purposes, including—

- BROMAZEPAM.
- CLONAZEPAM.
- DIAZEPAM.
- FLUNITRAZEPAM.
- FLURAZEPAM.
- LORAZEPAM.
- MEDAZEPAM.
- OXAZEPAM.
- PRAZEPAM.
- TEMAZEPAM.

CHLORPHENIRAMINE.

CHLORPROMAZINE and other substances structurally derived from phenothiazine with ataractic properties when used for therapeutic purposes, including—

- FLUPHENAZINE.
- PERICYAZINE.
- PERPHENAZINE.
- PROMAZINE.
- THIETHYLPERAZINE.
- THIOPROPAZATE.
- THIORIDAZINE.
- TRIFLUOPERAZINE.

CLEMASTINE.

CLOMIPRAMINE.

CLONIDINE.

CLORAZEPATE.

CODEINE except when included in the 2nd or 3rd Schedules.

CYCLOSERINE.

CYPROHEPTADINE.

DANTROLENE.

DESIPRAMINE.

DEXCHLORPHENIRAMINE.

DEXTROMORAMIDE.

DIFENOXIN.

DIHYDROCODEINE when included in the 3rd Schedule.
DIMENHYDRINATE.
DIMETHINDENE.
DIPHENHYDRAMINE.
DIPHENOXYLATE.
DIPHENYLPYRALINE.
DOTHIEPIN.
DOXEPIN.
DOXYLAMINE.
DROPERIDOL.
ECGONINE, its esters and derivatives which are convertible to ecgonine and cocaine.
ETHYLMORPHINE when included in the 4th Schedule.
FENFLURAMINE.
GLUTETHIMIDE.
HALOPERIDOL.
HYDROCODONE.
HYDROMORPHONE.
HYDROXYZINE.
IMIPRAMINE.
MAZINDOL.
MEBHYDROLIN.
MECLOZINE.
MEPROBAMATE.
MEPYRAMINE.
METHADONE.
METHAQUALONE.
METHDILAZINE.
MORPHINE its salts and derivatives.
NALBUPHINE.
NORMETHADONE.
NORTRIPTYLINE.
OPIUM in any form except the alkaloids noscapine and papaverine.
OXYCODONE.
PENTAZOCINE.
PETHIDINE its salts and derivatives.
PHENIRAMINE.
PHENOPERIDINE.
PHENYLTOLOXAMINE.
PHOLCODINE.
PROCHLORPERAZINE.

PROMETHAZINE.

PROTRIPTYLINE.

THENYLDIAMINE.

THIOTHIXENE.

TRIMEPRAZINE.

TRIMIPRAMINE.

TRIPROLIDINE.

[*Appendix K added in Gazette 11 July 1986 pp. 2339-40.*]

NOTES

1. This reprint is a compilation as at 22 July 1987 of the *Poisons Regulations 1965* and includes all amendments in the reprint published in the *Gazette* on 15 September 1981 and all amendments effected by the other regulations referred to in the following Table.

Table of Regulations

Regulations	Gazettal	Commencement	Miscellaneous
Poisons Act Regulations 1965	29 June 1965 pp. 1883-1914	1 July 1965	
(Regulations effecting amendments included in the previous reprint are not referred to in this Table)			
Poisons Act Amendment Regulations 1981	6 November 1981 p. 4527	6 November 1981	
Poisons Amendment Regulations 1982	16 July 1982 pp. 2727-28	16 July 1982	
Poisons Amendment Regulations (No. 2) 1982	24 December 1982 p. 4904	24 December 1982	
Poisons Amendment Regulations 1983	28 January 1983 pp. 341-42	28 January 1983	
Poisons Amendment Regulations (No. 2) 1983	23 September 1983 pp. 3803-07	23 September 1983	
Poisons Amendment Regulations 1984 (Erratum 13 April 1984 p.1020)	6 April 1984 p. 928	6 April 1984	
Health Legislation Amendment Regulations 1984	29 June 1984 pp. 1780-84	1 July 1984	
Poisons Amendment Regulations (No.2) 1984	12 October 1984 p. 3267	12 October 1984	
Poisons Amendment Regulations 1985 (Erratum 19 April 1985 p. 1409)	8 February 1985 pp. 519-20	8 February 1985	
Poisons Amendment Regulations (No. 2) 1985	8 Febuary 1985 pp. 520-21	8 February 1985	
Poisons Amendment Regulations (No. 3) 1985 (Erratum 29 March 1985 p. 1110)	15 March 1985 pp. 941-54	15 March 1985	
Poisons Amendment Regulations (No. 5) 1985	12 April 1985 pp. 1285-86	1 July 1985	

Regulations	Gazettal	Commencement	Miscellaneous
Poisons Amendment Regulations (No. 6) 1985	31 May 1985 p. 1882	31 May 1985	
Poisons Amendment Regulations (No. 4) 1985	7 June 1985 p. 1941	7 June 1985	
Poisons Amendment Regulations (No. 6) 1985	5 July 1985 p. 2392	5 July 1985	
Poisons Amendment Regulations (No. 8) 1985	20 September 1985 p. 3743	20 September 1985	
Poisons Amendment Regulations 1986	31 January 1986 pp. 332-33	31 January 1986	
Poisons Amendment Regulations (No. 2) 1986	28 February 1986 pp. 616-17	28 February 1986	
Poisons Amendment Regulations (No. 3) 1986	28 February 1986 p. 618	28 February 1986	
Poisons Amendment Regulations (No. 4) 1986 (Erratum 20 June 1986 pp. 2049-54)	23 May 1986 pp. 1716-20	23 May 1986	
Poisons Amendment Regulations (No. 5) 1986 (Erratum 30 May 1986 p. 1769)	23 May 1986 p. 1721	23 May 1986	
Poisons Amendment Regulations (No. 7) 1986	11 July 1986 pp. 2339-40	15 July 1986	
Poisons Amendment Regulations (No. 6) 1986	1 August 1986 p. 2739	1 August 1986	
Poisons Amemdment Regulations (No. 8) 1986	21 November 1986 p. 4269	21 November 1986	
Poisons Amendment Regulations (No. 9) 1986	21 November 1986 p. 4270	21 November 1986	
Poisons Amendment Regulations (No. 10) 1986	5 December 1986 pp. 4466-67	5 December 1986	
Poisons Amendment Regulations (No. 12) 1986	19 December 1986 pp. 4874-75	19 December 1986	
Poisons Amendment Regulations 1987	23 January 1987 p. 187	23 January 1987	
Poisons Amendment Regulations (No. 2) 1987	20 March 1987 p. 954	20 March 1987	
Poisons Amendment Regulations (No. 3) 1987	15 May 1987 p. 2121	15 May 1987	

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