Approved for Reprint 3 December 1982.

WESTERN AUSTRALIA

POISONS ACT 1964-1981.

ARRANGEMENT.

Sec.

21915/10/82

Approved for Reprint 3 December 1982.

WESTERN AUSTRALIA.

POISONS.

13° Elizabeth II., No. LXX.

No. 70 of 1964.[1]

28/84

[As amended by Acts:

No. 23 of 1966, assented to 27 October 1966;
No. 28 of 1967, assented to 17 November 1967;
No. 51 of 1967, assented to 5 December 1967;
No. 6 of 1969,[2] assented to 21 April 1969;
No. 87 of 1970,[3] assented to 30 November 1970;
No. 43 of 1978,[4] assented to 29 August 1978;
No. 57 of 1981,[5] assented to 13 October 1981;
No. 63 of 1981, assented to 13 October 1981,

and by Orders in Council published in the *Gazette* on 9/7/81; 18/12/81; 16/7/82; and 13/8/82; and reprinted pursuant to the Amendments Incorporation Act 1938.]

AN ACT to regulate and control the Possession, Sale and Use of Poisons and other Substances; to constitute a Poisons Advisory Committee; and for incidental and other purposes.

[*Assented to 11 December 1964.*]

BE it enacted—

PART I.—INTRODUCTORY PROVISIONS.

1. This Act may be cited as the *Poisons Act 1964-1981.* Short title. Amended by No. 63 of 1981, s. 2.

[1] Proclaimed to come into operation on 1 July 1965. See *Gazette*, 25/6/65, p. 1836.
[2] Proclaimed to come into operation on 13 June 1969. See *Gazette*, 13/6/69, p. 1765.
[3] Proclaimed to come into operation on 2 February 1971. See *Gazette*, 29/1/71, p. 277.
[4] Proclaimed to come into operation on 1 October 1980. *See Gazette*, 29/8/80, p. 3015.
[5] Came into operation 1 September 1982; see section 2.

21915/10/82

Commencement.

2. This Act shall come into operation on a date to be fixed by proclamation.[1]

Arrangement.

3. The arrangement of this Act is as follows:—

Savings.

4. Without limiting the provisions of the Interpretation Act 1918, generally, and in particular the provisions of sections fifteen and sixteen of that Act, and subject to the provisions of this Act, it is hereby declared that the repeal by the Pharmacy Act 1964, of any provision of the Pharmacy and Poisons Act 1910, or by the Police Act Amendment Act 1964, of any provision of Part VIA of the Police Act 1892, so far as that provision relates to poisons or drugs, does not affect any licence or permit granted or issued, or any document made or anything whatsoever done under the provisions so repealed. Any such licence, permit, document or thing so far as it is subsisting or in force at the time of the repeal and could have been granted, issued, made or done under this Act, shall on and after the commencement of this Act continue and have effect for the purposes of this Act (but in the case of a licence or permit only until the date of its expiry), except where this Act expressly or by necessary implication provides otherwise, as if such licence, permit, document or thing had been granted, issued, made or done under a corresponding provision of this Act and that corresponding provision had been in force when the licence, permit, document or thing was granted, issued, made or done, but so that any reference in the

[1] Proclaimed to come into operation 1 July 1965. See *Gazette* 25/6/65, p. 1836.

provision so repealed to the Council of the Pharmaceutical Society of Western Australia shall be read and construed as a reference to the Commissioner of Public Health appointed under the Health Act 1911. 28/84

5. In this Act unless the context requires otherwise— Interpretation. Amended by No. 23 of 1966, s. 2; No. 6 of 1969, s. 3.

"Advisory Committee" means the Poisons Advisory Committee constituted under Part II of this Act;

"automatic machine" means any machine or mechanical device used or capable of being used for the purpose of selling or supplying goods without the personal manipulation or attention of the seller or supplier or his employee or other agent at the time of the sale or supply;

"Commissioner" means the Commissioner of Public Health for the time being appointed under the provisions of the Health Act 1911; 28/84

"dentist" means a dentist registered under the provisions of the Dental Act 1939;

"department" means the department of the Public Service of the State principally assisting the Minister in the administration of this Act; ~~"Director" means the~~ **cutive Director, Public Health and Scientific Support Services in the department;** ". 28/84 28/84

~~"label" includes any tag, brand, mark or statement in writing, that is on or attached to or used in connection with any container or package containing any poison; and "labelled" has a corresponding meaning;~~

"licence" means a licence granted under this Act that is valid and unexpired;

"licensee" means a person who holds or is entitled to exercise a licence under this Act;

"medical practitioner" means a medical practitioner registered under the Medical Act 1894, or any previous corresponding enactment;

"member" means a person occupying any of the offices of the Advisory Committee, including that of chairman;

"pharmaceutical chemist" means a pharmaceutical chemist registered under the provisions of the Pharmacy Act 1964; or any previous corresponding enactment;

"poison" means any substance specified in any of the First, Second, Third, Fourth, Sixth, Seventh and Eighth Schedules or added to any of those Schedules by Order in Council;

"prohibited plant" means any plant from which a drug of addiction may be obtained, derived or manufactured, or such other plant as the Governor declares and is hereby authorized to declare from time to time to be a prohibited plant for the purposes of this Act; and includes any part of such a plant, except in the case of the plant Papaver somniferum, the non-viable seed of that plant;

"public institution" means—

(a) any Government Department, public hospital, University, or technical college or school; or

(b) any other institution or establishment that is not carried on for private gain and that the Governor by Order in Council declares to be a public institution for the purposes of this interpretation;

"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;

"Schedule" means a Schedule in Appendix "A" to this Act;

"specified drug" means any substance that is declared to be a specified drug for the purposes of this Act;

"substance" includes substance, material, compound, preparation, and admixture;

"to cultivate" in relation to a plant includes to sow and to plant;

"veterinary surgeon" means a registered veterinary surgeon under the provisions of the Veterinary Surgeons Act 1960;

"wholesale dealing" means sale or supply by a wholesale dealer in the ordinary course of wholesale business to persons licensed or otherwise expressly authorized by or pursuant to the provisions of this or any other Act, to be in possession of or to sell poisons or other substances specified in any Schedule or added thereto by Order in Council; and includes sale or supply to other persons in wholesale quantities in the ordinary course of wholesale business for use in connection with any prescribed profession, business, trade or industry or any public institution but not for resale.

Construction. Amended by No. 57 of 1981, s. 14.

6. (1) Except as otherwise expressly provided, this Act shall be read and construed as being in aid and not in derogation of the provisions of the Health Act 1911, and of the Misuse of Drugs Act 1981, but those provisions shall be read and construed subject to the express provisions of this Act and where there is any inconsistency between those provisions and the provisions of this Act, the latter provisions shall prevail.

(2) Any reference in any other Act, or in any regulation, rule or by-law made under any other Act, to any narcotic drug to which the Misuse of Drugs Act 1981 applies shall be deemed and be taken to be a reference to any drug of addiction or specified drug within the meaning of this Act.

Administration.

7. (1) Subject to the Minister and the provisions of this Act, the ~~Commissioner~~ Permanent Head shall be responsible for the administration of this Act. 28/84

(2) The cost of the administration of this Act shall be paid out of moneys appropriated by Parliament for the purpose.

PART II.—POISONS ADVISORY COMMITTEE.

Constitution of Poisons Advisory Committee. Amended by No. 63 of 1981. Schedule.

8. (1) For the purposes of this Act an Advisory Committee consisting of twelve members and having the functions prescribed by this Act is constituted under the name of the "Poisons Advisory Committee".

(2) The twelve members of the Advisory Committee shall be comprised of two *ex officio* members and ten nominee members, and of those members—

(a) the *ex officio* members shall be the Com-

Executive Director, or a medical practitioner employed in the department nominated for the purpose by the Permanent Head "; 28/84

the Government Analyst of the State, each by virtue of his office; or while any of those offices is vacant, the person acting in that office; and

(b) the nominee members shall be ten persons appointed by the Governor for terms of tenure of office in accordance with the provisions of section ten of this Act.

(3) Of the ten nominee members referred to in paragraph (b) of subsection (2) of this section—

(a) one shall be a pharmacologist nominated by the Senate of the University of Western Australia;

(b) one shall be a medical practitioner employed in the Department ~~of Public Health~~ specializing in occupational health, nominated by the Minister; 28/84

(c) two shall be medical practitioners, one of whom is a specialist physician, nominated by the body known as The Western Australian Branch of the Australian Medical Association (Incorporated);

(d) one shall be an officer of the Department of Agriculture, nominated by the Minister for Agriculture;

(e) two shall be persons, one of whom shall represent the wholesale dealers within the State engaged in wholesale dealing, nominated by the body known as The West Australian Chamber of Manufactures (Incorporated);

(f) one shall be a veterinary surgeon nominated by the body known as the Veterinary Surgeons' Board constituted under the Veterinary Surgeons Act 1960;

(g) one shall be a person nominated by the body known as The Council of the Pharmaceutical Society of Western Australia; and

(h) one shall be a person nominated by the body known as The Federated Pharmaceutical Service Guild of Australia (W.A. Branch).

officer
(a) of
inated,
ittee.

(4) The Executive Director, or the medical practitioner nominated pursuant to subsection (2) (a) of this section if one be so nominated, shall be the Chairman of the Advisory Committee ".

2184.

Procedure on default of nomination.

equires, by notice in writing to the registrar or secretary of any body referred to in subsection (3) of section eight of this Act, require that body to submit the name of its nominee as provided in that subsection within a period of forty-two days after receipt by the registrar or secretary of such notice, and if upon the expiration of that period, or such extension thereof as the Minister thinks fit and is hereby authorized to grant, he has not received the required name of the nominee, the Minister shall nominate such person to be a nominee member of the Advisory Committee as, having regard to the category in respect of which a person was required to be nominated, he thinks fit.

Term of office of nominee member.

10. (1) Subject to subsection (2) of this section the term of tenure of office of a nominee member expires by effluxion of time on the expiration of a period of three years commencing on the date of his appointment by the Governor.

(2) The respective terms of tenure of office of the persons first appointed to office of nominee member expire by effluxion of time—

(a) in the case of the four nominee members referred to in paragraphs (a), (b) and (c) of subsection (3) of section eight of this Act, at the expiration of one year;

(b) in the case of the three nominee members referred to in paragraphs (d) and (e) of that subsection, at the expiration of two years; and

(c) in the case of the three nominee members referred to in paragraphs (f), (g) and (h) of that subsection, at the expiration of three years,

commencing on the date of his appointment by the Governor to that office.

(3) The term of tenure of an *ex officio* member continues until the member ceases to occupy the office by virtue of which he is an *ex officio* member or until terminated by the Minister.

(4) A person is not rendered ineligible for appointment to the office of member or deputy member because he has previously occupied office as such, unless his appointment has been terminated under the provisions of section twelve of this Act.

(5) A nominee member or the deputy of any member may resign his office of member or deputy member if he sends to the Minister written notice under his hand of his resignation and the Minister accepts such resignation.

Vacation of office.

11. (1) The office of a member becomes vacant if—

(a) he becomes bankrupt, applies to take the benefit of any law for the relief of bankrupt or insolvent debtors, or compounds with his creditors;

(b) he is absent, except on leave granted by the Minister, from three consecutive meetings of the Advisory Committee;

(c) he becomes permanently incapable of performing his duties;

(d) he resigns his office in accordance with the provisions of this Act;

(e) he dies;

(f) the term of his tenure of office expires by effluxion of time;

(g) in the case of an *ex officio* member, the term of tenure is terminated pursuant to subsection (3) of section ten of this Act; **or**

(h) he is convicted of an indictable offence.

(2) On the occurrence of any vacancy in an office of member, a person eligible to be appointed to that office under the provisions of this Part shall in accordance with those provisions be appointed by the Governor to fill the vacancy, and a person so appointed holds office, subject to those provisions, for the remainder of the term of office of the person in whose place he is appointed.

(3) The performance or exercise of the functions, powers, duties or liabilities of the Advisory Committee is not affected by reason only of there being a vacancy in the office of a member.

Dismissal of members.

12. The Governor may terminate the appointment of a member of the Advisory Committee for inability, inefficiency or misbehaviour.

Leave of absence.

13. The Minister may grant leave of absence to a member of the Advisory Committee upon such terms as to remuneration or otherwise as the Governor from time to time determines.

Deputies of members.

14. (1) The Governor may in respect of any member of the Advisory Committee, appoint a person to be the deputy of that member to act in his office during his absence, and the provisions of subsection (3) of section eight and of section nine of this Act apply as well to the nomination and appointment of deputies of nominee members as to the nomination and appointment of the nominee members.

(2) Any person so appointed is entitled, in the absence from a meeting of the Advisory Committee of the member for whom he is the deputy, to attend that meeting, and when so attending shall be deemed to be a member and is authorized to carry out any function that the member of whom he is the deputy could, if present, exercise under this Act.

Acceptance of office.

15. Acceptance of or acting in the office of member or deputy member of the Advisory Committee by any person shall not of itself render the provisions of the Public Service Act 1978, or any other Act applying to persons as officers of the public service of the State, applicable to that member or deputy member, or affect or prejudice the application to him of those provisions if they applied to him at the time of the acceptance of or acting in such office.

Remuneration of members.

16. The members of the Advisory Committee and their deputies, other than those members and deputies who are officers in the public service of the State, are entitled, in respect of their attendances at meetings and carrying out their functions under this Act, to such remuneration and allowances as the Governor determines and is hereby authorized to determine from time to time.

Meetings of Advisory Committee.

17. (1) The Chairman shall convene the first meeting of the Advisory Committee to be held at a time and place appointed by him, and the Advisory Committee shall meet accordingly and shall hold such further meetings as it considers necessary for the conduct of its affairs.

(2) At a meeting of the Advisory Committee—

(a) seven members form a quorum;

(b) the Chairman, or in his absence, the person appointed to be his deputy, shall preside;

(c) if both the Chairman and his deputy are absent, the members present shall elect one of their number present at the meeting to be Chairman thereof;

(d) all questions shall be decided by a majority of votes of the members present and voting;

(e) each member, including the Chairman, shall be entitled to one vote only on the determination of any question;

(f) in the event of an equality of votes, the question shall be determined in the negative.

(3) The Advisory Committee shall cause to be kept minutes of all its proceedings in such manner as the Minister may direct or approve.

Officers of Advisory Committee.

18. (1) The Governor may appoint a secretary to the Advisory Committee and any other officers and servants of the Advisory Committee necessary for carrying out the provisions of this Act.

(2) Any person so appointed may, if required by the terms of his appointment to devote the whole of his time to the service of the Advisory Committee, be appointed under and be subject to the provisions of the Public Service Act 1978.

Functions of Advisory Committee.

19. The functions of the Advisory Committee are to advise the Minister and the ~~Commissioner~~ upon and to make recommendations in relation to—

(a) the necessity to amend any of the Schedules;

(b) the necessity to make, amend or revoke any regulation under this Act;

(c) any matter or thing with regard to the manufacture, distribution, sale, supply, possession, use or labelling of poisons and hazardous substances, or prohibiting the

use of any poison or hazardous substance that the Advisory Committee thinks fit or that the Minister or the ~~Commissioner~~ may refer to it; and

(d) any proposals or questions that may be referred to it with regard to any of the matters mentioned in paragraphs (a), (b) and (c) of this section.

PART III.—POISONS AND OTHER SUBSTANCES.

Division 1.—Classification.

Declaration of poisons or hazardous substances. Schedules. Amended by No. 28 of 1967, s. 2.

20. (1) For the purposes of this Act the substances specified in the First, Second, Third, Fourth, Sixth, Seventh and Eighth Schedules and referred to in subsection (2) of this section are declared to be poisons, and the substances specified in the Fifth Schedule so referred to are declared to be hazardous substances.

(1a) Without limiting the operation of subsection (1) of this section, a substance may be specified in a Schedule, and, pursuant to subsection (2) of this section, declared to be a poison or hazardous substance, as the case requires by reference to—

(a) the manner in which or the purpose for which, it is used or intended for use;

(b) the quantity in which it is supplied;

(c) the nature of the package, including the labelling thereof, in which it is supplied; or

(d) the physical or chemical state or form in which it is supplied.

(2) The substances specified in the Schedules referred to in subsection (1) of this section shall be classified by inclusion in the respective Schedules as follows—

(a) First Schedule: Substances that are of such extreme danger to human life as to warrant distribution thereof being limited to qualified persons;

(b) Second Schedule: Substances that are dangerous to human life if misused or carelessly handled but of necessity are required to be available to the public for medicinal or other purposes without undue restriction;

(c) Third Schedule: Substances that are for therapeutic use, and—

(i) in respect to which personal advice may be required by the purchaser concerning dosage, frequency of administration, and general toxicity;

(ii) with which excessive unsupervised self-medication is unlikely; and

(iii) for which there may exist such urgent need that the supply thereof on prescription only would cause hardship;

(d) Fourth Schedule: Substances the supply of which in the public interest should be restricted to medical, dental or veterinary prescription; and also potentially harmful substances pending evaluation of their toxic or deleterious nature;

(e) Fifth Schedule (Hazardous Substances): Substances of a dangerous nature that are commonly used for domestic purposes and are required to be readily available to the public but in respect of which caution is necessary in their handling, use and storage;

(f) Sixth Schedule: Substances that are required to be readily available to the public for agricultural, pastoral, horticultural or veterinary purposes, or for the control or destruction of pests and vermin, or for industrial purposes;

(g) Seventh Schedule: Substances of exceptional danger that require the taking and exercise of special precautions in their manufacture and use; and

(h) Eighth Schedule (Drugs of Addiction): Substances that are addiction producing drugs or potentially addiction producing drugs, including drugs so classified by the United Nations Organisation or its agencies.

Amendment of Schedules. Amended by No. 28 of 1967, s. 3.

21. The Governor may from time to time by Order in Council, notice of which shall be published in the *Government Gazette*, amend any of the Schedules referred to in section twenty of this Act by—

(a) the addition thereto or the deletion therefrom of any substance;

(aa) the deletion and substitution of all of the items in any Schedule;

(b) the transference of any substance from any Schedule to any other Schedule; or

(c) the alteration of any item in any Schedule,

and every order made under this section shall take effect on and from the day specified for that purpose in the notice, or if no day is so specified, upon the expiration of seven days after the date of publication in the *Government Gazette*, and thereupon the Schedule as so amended shall have the same force and effect as if the amendment effected by the order had been enacted in this Act.

Substances controlled by other laws may be exempted from Act. Inserted by No. 28 of 1967, s. 4.

21A. Where the Minister is of opinion that sufficient provision is made by other laws of the State regulating the supply, sale or use of any substance containing any poison or hazardous substance, he may certify in writing to that effect, and thereupon the Governor may by proclamation exempt that substance from all or any of the provisions of this Act and the regulations.

Sale of any poison may be prohibited.

22. (1) The Governor, on the recommendation of the Advisory Committee, may at any time and from time to time by proclamation prohibit the sale, supply or use of any poison or substance, whether specified in a Schedule or not, either absolutely or except upon and subject to such conditions and for such period or periods as the Governor may think fit.[1]

(2) A proclamation made under this section may be cancelled or from time to time varied, or an error in a proclamation may be rectified, by a subsequent proclamation.

Specified drugs. Inserted by No. 6 of 1969, s. 4. Amended by No. 57 of 1981, s. 15.

22A. (1) The Governor may, by Order in Council, declare any substance to be a specified drug for the purposes of this Act.

(2) Any substance that was, before the coming into operation of the Poisons Act Amendment Act 1969, declared to be a specified drug for the purposes of this Act continues, subject to subsection (3) of this section, to be a specified drug for the purposes of this Act and the Misuse of Drugs Act 1981.

(3) The Governor may, by Order in Council, vary or revoke any Order in Council made under subsection (1) of this section and may in like manner vary or revoke any Order in Council made before the coming into operation of the Poisons Act Amendment Act 1969, declaring any substance to be a specified drug for the purposes of this Act.

[1] See *Gazette* 4 December 1981, p. 4971.

Division 2.—Sale of Poisons.

Persons authorized to sell poisons. Amended by No. 6 of 1969, s. 5; No. 43 of 1978, s. 3.

23. (1) Except as provided by subsection (2) of this section, a person shall not manufacture, distribute, supply, or sell by wholesale or retail any poison unless he is licensed pursuant to the provisions of section twenty-four of this Act to do so.

(1a) Except as provided by subsection (2) of this section, a person shall not write, issue or authorize any prescription or document prescribing the use, sale or supply of a drug of addiction or a specified drug by, to, or in relation to any person.

(2) Subject to this Act—

(a) a pharmaceutical chemist is authorized to manufacture, have in his possession, and to use, supply or sell at his pharmacy in the ordinary course of his retail business any preparation, admixture or extract containing any poison;

(b) a medical practitioner or veterinary surgeon is authorized to have in his possession and to use, supply or sell in the lawful practice of his profession any poison;

(c) any dentist is authorized to have in his possession and to use in the lawful practice of his profession any poison; and

(d) a medical practitioner, veterinary surgeon or dentist is authorized to write, issue or authorize a prescription or document prescribing the use, sale or supply of a drug of addiction or a specified drug in the lawful practice of his profession,

but subject however to such conditions and restrictions as may be prescribed and subject to any notice given by the ~~Commissioner~~ Permanent Head pursuant to the regulations made under paragraph (ha) of subsection (2) of section sixty-four of this Act.

29/84

(3) The provisions of subsection (2) of this section do not authorize any medical practitioner, veterinary surgeon or dentist to sell any poison in an open shop unless he is licensed under this Act to do so.

Licences to sell poisons. Amended by No. 6 of 1969, s. 6.

24. (1) Subject to this Act the ~~Commissioner~~ may grant a licence—

(a) to manufacture any poison;

(b) to manufacture and distribute or sell by wholesale any poison;

(c) to sell by wholesale any poison; or

(d) to sell by retail any poison,

in or at any pharmacy or other premises or place of business specified in the licence, to any person who satisfies the ~~Commissioner~~ that he is a fit and proper person to be the holder of such a licence.

(2) An application for a licence under this section shall be made in the prescribed manner to the ~~Commissioner~~, who may in his discretion grant or refuse the licence.

(3) The ~~Commissioner~~ shall not grant any licence under this section unless and until he is satisfied that the premises of the applicant are suitable for the purpose in respect of which application is made for the licence, and are properly and hygienically equipped for that purpose.

(4) The ~~Commissioner~~ may grant—

(a) to a pharmaceutical chemist, a licence to sell by retail any poison;

(b) to a person who satisfies the ~~Commissioner~~ that he is carrying on a *bona fide* business in such circumstances as may be prescribed, a licence to sell by retail all or any of the poisons specified in the Sixth Schedule;

(c) to a person who satisfies the ~~Commissioner~~ that his place of business is distant at least five miles from the nearest place at which a pharmaceutical chemist conducts a pharmacy, and in such other circumstances as may be prescribed, a licence to sell by retail all or any of the poisons specified in the First, Second and Sixth Schedules;

(d) to such persons and under and subject to such conditions as may be prescribed a licence to sell all or any of the poisons specified in the Seventh Schedule.

(5) The ~~Commissioner~~ may from time to time, by notice, impose such conditions, restrictions and limitations on the sale, supply, use and possession of any poison specified in the Seventh Schedule as he considers necessary for safeguarding the public health.

(6) A notice given by the ~~Commissioner~~ under subsection (5) of this section—

(a) has effect according to its tenor, notwithstanding any other provision of this Act or the terms or conditions of any licence or permit in force thereunder; "this Act" includes regulations. See Act No. 30 of 1918, s. 4.

(b) may be of general application or apply to a particular person or class of persons, in a particular case or class of cases, or to particular circumstances or localities;

(c) has effect, if expressed to apply to any particular person, when served on that person and if not so expressed, when published in the *Government Gazette*; and

(d) may be varied or revoked by the ~~Commissioner~~ by subsequent notice.

(7) Any person who—

(a) having been served with a notice under subsection (5) of this section that is expressed to apply to him, fails to comply with or contravenes any condition, limitation or restriction contained in the notice; or

(b) fails to comply with or contravenes any condition, limitation or restriction contained in a notice published in the *Government Gazette*,

commits an offence and is liable on conviction to a penalty not exceeding two hundred dollars.

Permits to purchase poisons for specified purposes. Amended by No. 23 of 1966, s. 3.

25. (1) The Commissioner may permit fit and proper persons to purchase or otherwise obtain from manufacturers or wholesale dealers poisons for use for industrial, educational, advisory or research purposes, but not for re-sale.

(2) An application for a permit under this section shall be made in the prescribed manner to the Commissioner who may in his discretion grant or refuse the application.

Form of licences and permits and renewal thereof.

26. (1) Every licence or permit issued pursuant to the provisions of this Act shall—

(a) be in the prescribed form;

(b) specify the pharmacy or other premises or place of business in or at which the licence may be exercised, and be limited to one pharmacy or other premises or place of business only;

(c) be subject to such conditions, limitations and restrictions as may be prescribed and as the Commissioner thinks fit;

(d) be issued to the applicant upon payment of the prescribed fee (if any);

(e) remain in force until the thirtieth day of June next following the day of its issue, unless sooner cancelled, suspended or revoked; and

(f) be renewable from year to year.

(2) The holder of a licence or permit under this Act may at least one month prior to the date of the expiration thereof apply to the ~~Commissioner~~ for a renewal of his licence or permit as the case may be, and subject to this Act and payment of the prescribed fee (if any), the ~~Commissioner~~ may renew any licence or permit for the next ensuing year and issue to the applicant a renewed licence or permit as the case may require.

"this Act" includes regulations. See Act No. 30 of 1918, s. 4.

(3) Every renewal of a licence or permit under this section shall take effect from the first day of July in the year to which the renewal relates and shall continue in force until the thirtieth day of June next following that date unless sooner cancelled, suspended or revoked.

27. Every applicant for a licence or permit under this Act or for any renewal thereof shall pay to the ~~Commissioner~~ such fees therefor as are prescribed.

Fees for licences, permits and renewals.

28. The ~~Commissioner~~ may in his discretion cancel, suspend or revoke at any time any licence or permit issued pursuant to the provisions of this Act, and any licence or permit so cancelled, suspended or revoked shall thereupon cease forthwith to have effect and shall be surrendered to the Commissioner on demand.

Commissioner may cancel or suspend licence or permit.

29. (1) Any person aggrieved by the refusal of the ~~Commissioner~~ to grant or renew any licence or permit under this Act, or by an order of the ~~Commissioner~~ cancelling, suspending or revoking any

Appeal against order of Commissioner.

licence or permit, may within six months after notice of such refusal or of such order appeal against the same to a stipendiary magistrate sitting as a court of summary jurisdiction.

(2) The stipendiary magistrate hearing the appeal shall enquire into and decide upon the appeal and may make such order in the matter as he may think just, and his decision shall be final and conclusive.

(3) Every appeal brought pursuant to the provisions of this section shall be brought and conducted in accordance with the regulations.

Licence not to be granted to company or friendly society.

30. (1) A licence under this Part shall not be granted to a company or friendly society although the company or friendly society is lawfully carrying on business as a pharmaceutical chemist; but such a licence may be granted to any pharmaceutical chemist entitled thereto for his own use, who is *bona fide* employed by or engaged with that company or friendly society in the business of a pharmaceutical chemist and may be used by him for the benefit of that company or friendly society.

(2) Where in accordance with the provisions of subsection (1) of this section a licence is used by a pharmaceutical chemist for the benefit of a company or friendly society, that company or friendly society, and the manager or other officers thereof respectively and such pharmaceutical chemist, are jointly and severally liable in respect of any offence under this Act committed by any servant or other agent of that company or friendly society in relation to the possession, sale or use of poisons.

Division 3.—General Provisions.

Sales of poison to be recorded in a book.

31. (1) Every person who sells by retail any poison or class of poison prescribed by regulation for the purposes of this section, shall make a true record of each sale in a book to be kept as prescribed.

(2) A person shall not sell any poison, a record of the sale of which is required to be made in a book pursuant to subsection (1) of this section, on an order by letter, telegram or radiogram unless the purchaser is known to the vendor and the letter, telegram or radiogram is preserved by the vendor and particulars of the date and sender of the order are entered in the book referred to.

Unauthorized sales of poisons.

32. A person shall not—

(a) sell any poison by wholesale unless he is licensed under this Act to do so;

(b) sell any poison by wholesale to any person who is not authorized by or licensed or permitted under this Act to have in his possession or to sell such poison;

(c) except as provided by section one hundred and thirty of the Vermin Act 1918,[1] sell or supply any poison unless he is authorized by or licensed under this Act to do so; or

(d) sell or supply any poison except in accordance with the authority of his licence or permit and the terms and conditions thereof.

Wholesaler not to sell by retail.

33. A wholesale dealer shall not sell any poison by retail unless he is authorized by or licensed under this Act to do so.

Sales to certain persons prohibited. Amended by No. 23 of 1966, s. 4.

34. (1) A person shall not sell any poison or class of poison prescribed by regulation for the purposes of this section to any person—

(a) who is apparently under the age of eighteen years; or

(b) who is unknown to the vendor, unless the sale is made in the presence of an adult witness who is known to the vendor and who knows the purchaser.

[1] Repealed by Agriculture and Related Resources Protection Act 1976.

(2) The witness in whose presence the sale is made pursuant to paragraph (b) of subsection (1) of this section shall, before the delivery of the poison to the purchaser, sign the entry (including the entry of his own name and place of residence) in the book required to be kept under section thirty-one of this Act.

Making false declarations.

35. A person who for the purpose of obtaining for himself or for any other person the grant, issue or renewal of a licence or permit under this Act—

(a) makes any declaration or statement that is false in any material particular; or

(b) knowingly produces or makes use of any such declaration or statement,

commits an offence against this Part.

Drugs not to be used for self administration.

36. A person shall not use or attempt to use, or prescribe, any drug of addiction or specified drug for the purpose of self administration; but a person for whom a medical practitioner has prescribed a drug of addiction or a specified drug in the course of treatment of that person as a patient may take or use that drug to the extent and for the purpose for which it was so prescribed.

New drugs to be classified.

37. (1) Before any new drug is first offered for sale to the public the manufacturer, importer or distributor, as the case may require, of the new drug shall make application to the ~~Commissioner~~ as provided in this section to classify the new drug by determining the Schedule (if any) in which it is to be included and specified and to determine the percentage exemption limit (if any) to be permitted in respect to the new drug.

(2) Every application under this section shall be made in the prescribed manner and on the prescribed form to the ~~Commissioner~~ who shall

submit the application to the Advisory Committee for its consideration and for reference by it to the Poisons Advisory Panel of the body known as the National Health and Medical Research Council.

(3) The Advisory Committee shall forward in writing to the ~~Commissioner~~ its recommendations in relation to the application and the ~~Commissioner~~ upon receipt of and after having regard to those recommendations, shall classify the new drug and determine in which Schedule (if any) it shall be included and specified and may, if he thinks it necessary to do so, determine the percentage exemption limit to be permitted in respect to the new drug.

(4) The ~~Commissioner~~ shall notify in writing the applicant of the classification of the new drug and his determinations in relation thereto pursuant to subsection (3) of this section and thereupon cause the new drug to be added to the Schedule (if any) in which he has determined that it is to be included and specified, in accordance with the provisions of section twenty-one of this Act.

(5) The decision of the ~~Commissioner~~ in respect to any application made to him under the provisions of this section shall be final and conclusive.

(6) In and for the purposes of this section—

"new drug" means a therapeutic substance for use in human therapy that is not included in the latest edition for the time being of any of the respective books called the British Pharmacopoeia, the British Pharmaceutical Codex and the United States Pharmacopoeia, or a substance specified in a Schedule for which the method of manufacture, composition, route of administration or indications for use is changed.

Offence in respect of a new drug.

38. A person who offers for sale or sells, or causes or permits to be offered for sale or sold, to the public any new drug referred to in section thirty-seven of this Act before an order made under section twenty-one of this Act has taken effect to add that new drug to a Schedule, except where the ✳Commissioner has determined that the new drug does not require to be placed in a Schedule, commits an offence against this Act.

Sale of new drug may be prohibited. Amended by No. 23 of 1966, s. 5.

39. (1) Every new drug, whether specified in a Schedule or not, is deemed to be a poison within the meaning of this Act pending notification by the ✳Commissioner of the classification of that new drug and his determinations in relation thereto pursuant to the provisions of section thirty-seven of this Act, but where in respect of any new drug the ✳Commissioner so determines that such new drug does not require to be placed in a Schedule, that new drug shall no longer be deemed to be such a poison.

(2) Notwithstanding the provisions of sections thirty-seven and thirty-eight of this Act, where application is made for classification of a new drug the ✳Commissioner may before the new drug is so classified, if the Advisory Committee so recommends, authorize the sale or supply of that new drug to any person or institution approved by the Advisory Committee, but any such sale or supply shall be made only upon and subject to such conditions as the Advisory Committee thinks fit.

[(3) *Repealed by No. 32 of 1966, s. 5.*]

Offences against this Part. Amended by No. 23 of 1966, s. 6; No. 43 of 1978, s. 4.

40. Except where by this Act it is expressly enacted otherwise, every person who—

(a) contravenes or fails to comply with any of the provisions of this Part;

(b) contravenes or fails to comply with any conditions, limitation or restriction to which any authority, licence or permit issued under this Part is subject;

(ba) contravenes or fails to comply with any conditions, limitation or restriction of any notice given by the Commissioner pursuant to the regulations made under paragraph (ha) of subsection (2) of section sixty-four of this Act;

(c) purchases any poison and gives false information in answer to inquiries that by or under this Act are required to be made by the vendor; or

(d) signs his name as a witness to the sale of any poison to a person unknown to him,

commits an offence against this Part.

Penalty: For a first offence, five hundred dollars; for a second or subsequent offence, three thousand dollars.

PART IV.—DRUGS OF ADDICTION.

Manufacture of heroin permitted in certain cases. Amended by No. 23 of 1966, s. 7; No. 57 of 1981, s. 16.

41. (1) Notwithstanding anything in the Misuse of Drugs Act 1981, it shall not be unlawful for a person to manufacture or prepare heroin for educational, experimental or research purposes—

(a) in any university, college, school or institution that the Governor by Order in Council approves for that purpose; and

(b) under and subject to such conditions as the Governor by Order in Council imposes, and is hereby authorized to impose, in the case of any such approved university, college, school or institution.

(2) In this section "heroin" means diacetylmorphine and includes its salts and any preparation, admixture, extract or other substance containing it.

Licence to cultivate prohibited plants. Inserted by No. 23 of 1966, s. 8. Amended by No. 57 of 1981, s. 17.

41A. (1) Subject to this Act the ~~Commissioner~~ * may grant to any person a licence to cultivate, sell, purchase or have in his possession any prohibited plant.

* Permanent Head. 28/84

(2) A licence granted pursuant to this section shall be subject to such conditions as may be prescribed and as the ~~Commissioner~~ * may in his discretion impose.

[(3) *Repealed by No. 57 of 1981, s. 17.*]

[**42.** *Repealed by No. 57 of 1981, s. 18.*]

[**43.** *Repealed by No. 57 of 1981, s. 19.*]

[**43A.** *Inserted by No. 6 of 1969, s. 7; repealed by No. 43 of 1978, s. 5.*]

Offences generally against this Part. Amended by No. 23 of 1966, s. 9; No. 51 of 1967, s. 2; No. 87 of 1970, s. 4; No. 43 of 1978, s. 6.

44. (1) A person who—

(a) contravenes or fails to comply with any provision of this Part; or

(b) within the State aids and abets, counsels or procures the commission in any place outside the State of any offence punishable under the provisions of any corresponding law in force in that place or does any act preparatory to or in furtherance of any act which if committed within the State would constitute an offence against this Part,

commits an offence against this Part.

(2) A person who commits an offence against this Part, not being an offence for which a penalty is otherwise in this Part expressly provided, is liable upon conviction to a fine of three thousand dollars, or imprisonment for a term of three years, or to both the fine and imprisonment.

(3) A person convicted of an offence against this Part shall forfeit to Her Majesty all articles in respect of which the offence was committed, and the court before which the offender is convicted may order any forfeited articles to be destroyed or otherwise disposed of as the court thinks fit.

(4) A person who—

(a) attempts to commit an offence under this Part; or

(b) solicits or incites another person to commit such an offence,

is, without prejudice to any other liability, liable on summary conviction to the same punishment and forfeiture and to be dealt with as if he had been convicted of the offence which he attempted to commit, or the offence which he solicited or incited another to commit.

Interpretation of "corresponding law".

45. (1) In this Part the expression, "corresponding law" means any law stated in a certificate that purports to have been issued by or on behalf of the Government of—

(a) any British possession (including any territory under Her Majesty's protection, or governed under a trusteeship agreement by the Government of any part of Her Majesty's dominions) outside the State; or

(b) any foreign country (including any protectorate thereof or any territory governed under a trusteeship agreement by the Government thereof),

to be a law providing for the regulation and control in that possession or country of the manufacture, sale, use, export or import of drugs in accordance with the provisions of any of the Conventions referred to in Appendix "B" to this Act.

(2) Any statement in a certificate referred to in subsection (1) of this section as to the effect of the law mentioned in that certificate, or any statement in any such certificate that any facts constitute an offence against that law, shall be conclusive.

PART V.—MISCELLANEOUS PROVISIONS.

Containers of poisons to be marked or labelled.

46. A person shall not sell any poison or hazardous substance unless the container immediately containing it is marked or labelled in such manner and with such particulars as are prescribed.

Medicines for internal use not to be sold in certain containers.

47. (1) A person shall not sell any drug or medicine that is for internal use or any food, drink or condiment in a container—

(a) of like description to that prescribed by the regulations for a container in which any poison intended for external use may be sold; or

(b) of such a description as not to be readily distinguishable by sight and touch, or by either sight or touch, from a container in which a poison intended for external use may be sold.

"This Act" includes regulations. See Act No. 30 of 1918, s. 4.

(2) Nothing in this section affects any other requirement of this Act relating to the containers in which drugs or medicines that are or contain poisons within the meaning of this Act may be sold.

Prohibition against hawking, etc.

Amended by No. 23 of 1966, s. 10.

48. A person shall not, except pursuant to a licence issued by the ~~Commissioner~~ Permanent Head, 28/84—

(a) sell or attempt to sell; or

(b) hawk or peddle, or distribute or cause to be distributed as a sample,

any poison in any street or public place or from house to house.

Penalty: One hundred dollars.

Prohibition against selling by automatic machines. Amended by No. 23 of 1966, s. 11.

49. (1) A person shall not—

(a) install or permit to be installed on or about his premises or elsewhere any automatic machine for the sale or supply of any poison;

(b) sell or supply any poison by means of any automatic machine;

(c) place or permit to be placed, any poison in any automatic machine that is on or about his premises or under his control; or

(d) permit or suffer any person to purchase or be supplied with or otherwise obtain any poison by means of any automatic machine.

(2) A person who contravenes or fails to comply with any provision of subsection (1) of this section commits an offence against this Act and is liable on conviction to a fine of one hundred dollars or to imprisonment for a term not exceeding six months, and in addition to a daily penalty of ten dollars during the time that the offence is continued after conviction.

(3) Any automatic machine in respect of which any person is convicted of an offence under this section may in the discretion of the court before which proceedings for the offence are taken be forfeited to Her Majesty.

Leaving poisons unlabelled an offence. Amended by No. 23 of 1966, s. 12.

50. (1) A person who being in charge or possession of any poison leaves it in any place (whether that place is or is not ordinarily accessible to other persons), unless the bottle or container in which the poison is contained is marked clearly and legibly with the word, "Poison" or with other

prescribed words, and otherwise duly labelled in the manner provided by section forty-six of this Act, commits an offence against this Act.

Penalty: One hundred dollars.

(2) This section does not apply to pharmaceutical chemists in the conduct of their business or to persons granted exemption pursuant to subsection (3) of this section.

28/84 (3) The ~~Commissioner~~ Permanent Head may exempt any person from the provisions of this section where he is of opinion, having regard to the circumstances of the case, that such exemption is warranted.

Calculation of percentages for liquid preparations.

51. For the purposes of this Act percentages in the case of liquid preparations shall (unless other provision in that behalf is made by regulation under this Act) be calculated on the basis that a preparation containing one per centum of any substance means a preparation in which—

(a) one gramme of the substance, if a solid; or

(b) one millilitre of the substance, if a liquid,

is contained in every one hundred millilitres of the preparation, and so in proportion for any greater or less percentage.

PART VI.—SUPPLEMENTARY PROVISIONS.

Orders in Council may be cancelled or amended.

52. An Order in Council made under the provisions of this Act may be cancelled or from time to time varied or amended, or an error in any such Order may be rectified, by a subsequent Order in Council.

Apprehension of offenders. Amended by No. 23 of 1966, s. 13.

53. (1) Any officer or constable of the Police Force and all persons whom he shall call to his assistance, may take into custody with or without a warrant any person found committing any offence—

(a) against section forty-eight of this Act; or

(b) against any provision of Part IV of this Act or any regulation made thereunder prohibiting the sale of any drug of addiction or specified drug, or the cultivation, sale, purchase or possession of any prohibited plant,

whose name and residence are unknown to and cannot readily be ascertained by that officer or constable, or who on demand neglects or refuses to give his name and address or either of them, or gives a false name or address.

(2) The powers conferred by this section upon officers and constables of the Police Force are in addition to and not in diminution of the powers conferred on those officers and constables by the provisions of the Police Act 1892, or of any other Act.

Powers to enter, etc. Amended by No. 23 of 1966, s. 14.

54. (1) Any inspector appointed under the Health Act 1911, or other person authorized in that behalf in writing by the Minister, may at any reasonable time, for the purpose of ascertaining whether the provisions of this Act and the regulations are being complied with,—

(a) enter upon any premises occupied by any person licensed or otherwise authorized under this Act to have in his possession any poison or prohibited plant;

(b) inspect and examine any room or part of the premises entered upon, and any goods or records in those premises;

(c) take an account of any poisons and any prohibited plants in those premises; or

(d) on payment or tender of a reasonable price, demand, take and obtain any sample of any poison or prohibited plant in or upon those premises.

(2) Any person who—

(a) refuses or fails to admit any inspector or authorized person demanding to enter upon premises pursuant to the provisions of this section;

(b) refuses to permit any inspector or authorized person to take or obtain any sample pursuant to the provisions of this section; or

(c) delays or obstructs, or causes or permits to be delayed or obstructed, any inspector or authorized person in the exercise of his powers under this section,

commits an offence against this Act.

Search warrant may be granted. Amended by No. 23 of 1966, s. 15; No. 57 of 1981, s. 20.

55. (1) If it appears to a justice on complaint made on oath before him that there is reasonable ground for suspecting—

(a) that there is in any house or premises any poison or prohibited plant in contravention of this Act or the regulations; or

(b) that any person has in his possession or under his control in any house or premises—

(i) any poison, substance or prohibited plant or any preparation thereof in contravention of this Act or the regulations; or

(ii) any document directly or indirectly relating to or connected with any transaction or dealing that is or would, if carried out, be an offence against any provision of this Act or the regulations, or against the provisions of any corresponding law in force in any place outside the State,

the justice may give to any member of the Police Force a search warrant in the form in Appendix "C" to this Act.

(2) A warrant given under subsection (1) of this section authorizes the member of the Police Force named in the warrant, within one month from the date of the warrant, and with such assistants as may be necessary—

(a) to enter into and upon and search the house or premises specified in the warrant at any time during the day or night, and to open and break open if necessary and search all things found therein or thereon;

(b) to use force if necessary in making entry whether by breaking open doors or otherwise;

(c) to arrest and bring before a stipendiary magistrate or two justices any person found committing any offence in such house or premises against the provisions of Part IV of this Act;

(d) to search all persons found in or upon the house or premises;

(e) to seize, or seize and carry away—

(i) any substance or preparation that may be reasonably suspected of being or containing a poison, or any prohibted plant, found in the house or premises, or in the possession or under the control of any person therein, or that is in that house or premises or under such control in contravention of any provision of this Act or the regulations;

(ii) any articles used or capable of being used for the purpose of preparing, taking or administering any drug of addiction or specified drug for the purposes of addiction; and

(iii) any document referred to in subparagraph (ii) of paragraph (b) of subsection (1) of this section.

(3) All articles seized under subparagraph (ii) of paragraph (e) of subsection (2) of this section shall on conviction of the person in whose possession those articles were found be forfeited to Her Majesty, and the court before which such person was convicted may order all or any of those articles to be destroyed or otherwise disposed of as the court thinks fit.

(4) Subject to subsection (3) of this section, any poison (not being a drug of addiction or a specified drug) seized under the provisions of this section may, at the request of the owner thereof and with the approval in writing of the Minister, be returned to such owner subject to such conditions or limitations as to its use or otherwise as the Minister may in his discretion impose.

(5) The provisions of this section shall be in addition to and not in derogation of the provisons of the misuse of Drugs Act 1981.

Sales by employees, etc.

56. For the purposes of this Act any person on whose behalf a sale is made is deemed to be the person who sells, and every employee, assistant or apprentice of such person is liable to the like penalties as the person on whose behalf he makes any sale.

Persons deemed to have sold poisons.

"This Act" includes regulations.

57. (1) Where any poison or hazardous substance is sold in an unopened package to an inspector or authorized person and in respect of the sale thereof there is a contravention of or failure to comply with any provision of this Act, each of the persons referred to in paragraphs (a) and (b) of this subsection shall, in addition to the person who actually sold the package to the inspector or authorized person, be liable in respect of such contravention or failure, namely—

(a) if the package has a label on or attached to it, any person who appears from that label to have manufactured or prepared

such poison or hazardous substance, or to have imported it into the State, or to have enclosed or caused to be enclosed in that package such poison or hazardous substance, or to have been the wholesale supplier thereof; or

(b) if the package has a label on or attached to it but such label does not disclose any of the particulars referred to in paragraph (a) of this subsection, or if the package has no label on or attached to it, any person who has previously sold the unopened package.

(2) A person to whom the provisions of subsection (1) of this section apply is deemed to have sold the unopened package to the inspector or authorized person as on the day when and at the place where the inspector or authorized person purchased it, and that person is liable to the same penalty as if he had actually sold such package to the inspector or authorized person on that day and at that place.

(3) It shall be a defence to a charge under this section if the person charged shows—

(a) that the contravention or non-compliance is due to the act or default of some subsequent seller;

(b) that the contravention or non-compliance is due to deterioration or other causes beyond his control; or

(c) where the package has a label on or attached to it, that he did not in fact affix or attach the label or cause it to be affixed or attached or enclose or cause to be enclosed the poison or hazardous substance in the package.

(4) Nothing in this section shall affect the liability of any person selling any such unopened package to an inspector or authorized person with respect to any contravention or non-compliance due

to his default or to other causes within his control; and the conviction of any person under the provisions of this section shall not exonerate the person selling such unopened package or any other person from liability with respect to any such contravention or non-compliance.

(5) Without affecting the generality of the application of this or any other provision of this Act to firms or the members of them, where a firm appears from any such label to have imported, manufactured or prepared any poison, or as the case may be, hazardous substance, or to have been the wholesale supplier thereof or to have enclosed the same in a package—

(a) proceedings under this section may be taken (whether in a court of petty sessions or otherwise) and penalties recovered accordingly against any member or members of the firm; and

(b) this section shall be read and construed and have effect as though the name or names of the member or members of the firm had appeared on such label.

(6) In this section—

"authorized person" means a person authorized in writing by the Minister for the purposes of this Act;

"inspector" means an inspector appointed under the Health Act 1911;

"wholesale supplier" means a person who sells or supplies poisons or hazardous substances to any other person for the purpose of re-sale.

Evidence on prosecutions.

58. Whenever in any prosecution for a contravention of or failure to comply with any provision of this Act or any regulations made under this Act it is necessary or proper to prove in respect of any particular article or substance that it is a poison, or as the case may be, hazardous substance, then in every such case—

(a) evidence that any substance commonly sold under the same name or description as that particular article or substance is a poison or hazardous substance shall be *prima facie* proof that such particular article or substance also conforms to the same description accordingly; and

(b) evidence that any particular article or substance or the container thereof is labelled, "Poison" or with other prescribed words, shall be *prima facie* proof that such particular article or substance is a poison or, as the case may be, hazardous substance.

28/K4

Publication of list of licensed persons.

59. The ~~Commissioner~~ Permanent Head shall in the month of August in each year cause to be published in the *Government Gazette* a list of the names and places of business of all persons who hold licences or permits under this Act, and the production of a copy of the *Government Gazette* containing any such list as last published shall be *prima facie* proof in all courts and in all legal proceedings that the persons specified in such list hold such licences or permits.

Proof of certificate of analyst.

60. (1) In any legal proceedings for offences against this Act—

(a) the production of a certificate purporting to be signed by an analyst with respect to any analysis made by him shall, without proof of the signature of the person appearing to have signed the certificate or that he is an analyst, be sufficient evidence—

(i) of the identity of the thing analysed;

(ii) of the result of the analysis; and

(iii) of the matters relevant to such proceedings stated in the certificate,

unless the defendant by not less than three days' notice in writing delivered to the complainant and by a like three days' notice delivered to the analyst (opportunity to deliver which notices shall be afforded the defendant) requires the analyst to attend as a witness; and

(b) the court may, in addition to any other order as to costs, make such order as it thinks just as to the conduct money of the analyst and the expenses and remuneration to be paid for any analysis.

(2) For the purpose of this section, "analyst" means an analyst appointed under the provisions of the Health Act 1911.

Evidence of qualifications.

61. In any legal proceedings under this Act—

(a) the production of a copy of the *Government Gazette* containing the several registers or lists as last published in relation to the time in question of medical practitioners, pharmaceutical chemists, dentists or veterinary surgeons and of persons holding licences or permits under this Act shall, if the name of the defendant does not appear in any of such registers or lists, be *prima facie* proof that he is not a medical practitioner or a registered pharmaceutical chemist, dentist, veterinary surgeon or a person who holds a licence or permit under this Act;

(b) a certificate that any person is or is not, or was or was not, on a certain date or for a certain period a medical practitioner or a registered pharmaceutical chemist, dentist, veterinary surgeon or a person who holds

a licence, permit or authority under this Act shall be *prima facie* proof of the fact therein stated if the certificate purports to be signed—

(i) in the case of a medical practitioner, by the registrar of the Medical Board constituted under the Medical Act 1894;

(ii) in the case of a registered pharmaceutical chemist, by the registrar of the Pharmaceutical Council of Western Australia, constituted under the Pharmacy Act 1964;

(iii) in the case of a registered dentist, by the registrar of The Dental Board of Western Australia, constituted under the Dental Act 1939;

(iv) in the case of a registered veterinary surgeon, by the registrar of the Veterinary Surgeons' Board, constituted under the Veterinary Surgeons Act 1960; and

(v) in the case of a person who holds a licence, permit or authority under this Act, by the ~~Commissioner~~ Permanent Head. 23/84

62. Every person who contravenes or fails to comply with any provision of this Act or any regulation made under this Act commits an offence against this Act and if no penalty is expressly provided with respect to that offence is liable on conviction to a penalty not exceeding one hundred dollars. General penalty. Amended by No. 23 of 1966, s. 16.

63. (1) No act, matter or thing done or omitted to be done in good faith by the Minister or by the ~~Commissioner~~, or by the Advisory Committee or by any member thereof or by the secretary or any other Protection from liability.

officer thereof, or by any inspector or authorized person or by any member of the police force, in the administration or intended administration of this Act, or in the exercise or performance or intended exercise or performance of any of his or its powers, functions or duties under this Act, shall subject the Minister or the ~~Commissioner~~, or the Advisory Committee or any member or the secretary or other officer thereof, or any inspector, authorized person or member of the police force, to any liability in respect thereof.

(2) In this section inspector and authorized persons have the same respective meanings as are given to them in section fifty-seven of this Act.

Regulations. Amended by No. 23 of 1966, s. 17; No. 6 of 1969, s. 8; No. 43 of 1978, s. 7; No. 57 of 1981, s. 21.

64. (1) The Governor may make regulations prescribing all matters that by this Act are required or permitted to be prescribed, or that are necessary or convenient to be prescribed, for carrying out or giving effect to this Act.

(2) Without limiting the generality of the powers conferred by subsection (1) of this section, the Governor may make regulations for or with respect to—

(a) the possession, sale and safe custody of poisons and hazardous substances including the specifications of cupboards and other receptacles and the manner of storage of any poison or hazardous substance;

(b) specifying the containers in which any poison or hazardous substance may be sold, and the shape, size and materials of such containers, and prohibiting the use of such containers for other substances;

(c) marking and labelling, and specifying the particulars (including antidotes) to be included in labels on or attached to, containers of poisons and hazardous substances;

(d) prohibiting or regulating the possession, manufacture, distribution, supply, sale, handling or use of any poisons or hazardous substances either absolutely or except under such circumstances or conditions as may be prescribed;

(da) prohibiting or regulating the cultivation possession, sale or purchase of any prohibited plant either absolutely or subject to such conditions as may be prescribed, and prescribing those conditions;

(e) prescribing precautions to be taken in the manufacture, storage, handling or use of any poisons or hazardous substances;

23/84 A Permanent Head

(f) the application for and the granting, issue, renewal, cancellation and suspension of licences, permits and authorities under this Act;

(g) prescribing the persons to whom and the circumstances and conditions in and under which licences to sell by retail poisons specified in the First, Second, Sixth or Seventh Schedules may be granted under section twenty-four of this Act;

(h) the application for classification under section thirty-seven of this Act of new drugs and the procedure to be followed in relation to such application and to the determination and notice in respect thereof;

(ha) authorizing the ~~Commissioner~~, by notice given to any such person as is referred to in subsection (2) of section twenty-three of this Act, to revoke, in whole or in part, the authority conferred by that subsection on that person in relation to drugs of addiction and specified drugs;

(i) prescribing conditions, limitations and restrictions to which licences and permits under this Act shall be subject;

(j) prescribing the form of, and the particulars to be recorded in, the book required to be kept pursuant to section thirty-one of this Act, and the procedure to be followed in relation to the sale and recording of poisons;

(ja) requiring persons engaged in the cultivation, sale, distribution or supply of any prohibited plant, or the manufacture, sale, distribution or supply of any poison or hazardous substance, to keep such books, records or documents, and furnish such information, relating to such prohibited plant, poison or hazardous substance as the ~~Commissioner~~ may require from time to time, and providing for production of those books, records or documents and the furnishing in writing or otherwise of that information to the ~~Commissioner~~ at such times and in such manner as he may direct;

(k) prescribing the manner in which appeals against decisions of the ~~Commissioner~~ under this Act shall be brought and the procedure to be followed in the conduct of such appeals;

(l) prescribing the precautions to be observed in respect to the sale of poisons or hazardous substances ordered by letter, telegram or radiogram;

(m) the inspection of premises, stocks, books, and documents relating to poisons, hazardous substances and prohibited plants;

(n) prohibiting or regulating the sale of any poison or hazardous substance by methods of self-service other than any such methods prescribed;

(o) providing for the forfeiture of any poison, hazardous substance or prohibited plant unlawfully in the possession of any person and for the disposal of any poison, hazardous substance or prohibited plant so forfeited;

(p) specifying the persons or classes of persons authorized or entitled to purchase, use or be in possession of any poison;

(q) exempting from all or any of the provisions of this Act and the regulations, substances containing any poison that by their nature are not capable of being used in evasion of this Act and the regulations, or that are supplied or sold by a pharmaceutical chemist or in accordance with the prescription of a medical practitioner, dentist or veterinary surgeon for an individual and specific case;

(r) authorizing medical practitioners, and pharmaceutical chemists dispensing medicines and drugs at any public hospital or institution, or persons in charge of laboratories for the purpose of research or instruction, dentists, veterinary surgeons, and such other persons as to the ~~Commissioner~~ may seem proper, to be in possession of any poison or hazardous substance for the purposes of their respective professions or employments, and prescribing the conditions and restrictions upon and subject to which such authority may be given;

(s) regulating the issue by medical practitioners, dentists or veterinary surgeons of prescriptions containing any poison, the dispensing of such prescriptions, and the supply of any such poisons thereunder;

(sa) prohibiting and regulating the issue by medical practitioners, dentists or veterinary surgeons of prescriptions containing any drug of addiction or any specified drug, or

any class of drug of addiction or any class of specified drug, the dispensing of such prescriptions and the supply of drugs of addiction or specified drugs thereunder;

(t) prescribing the colouring of any poison or hazardous substance;

(u) providing for the disposal of automatic machines forfeited pursuant to the provisions of this Act;

(v) prescribing fees to be paid for the issue and renewal of licences and permits under this Act;

(w) prescribing forms to be used for the purposes of this Act;

(x) prescribing a penalty of not more than one hundred dollars for any contravention of or failure to comply with the regulations;

(y) any other matter or thing in any manner relating to poisons, hazardous substances or prohibited plants;

(z) any other purpose that the Governor deems necessary for safeguarding the public and the public health in relation to poisons, hazardous substances and prohibited plants.

(2a) Regulations may be made under this section requiring any person who is licensed or otherwise authorized under this Act to have in his possession, manufacture, supply or sell any poison, drug of addiction or specified drug to—

(a) retain any document, writing, prescription or authorization or record thereof relating to the sale or supply of any drug of addiction or specified drug;

(b) maintain such records relating to the sale or supply of drugs of addiction or specified drugs as may be prescribed;

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

(2b) Regulations made under subsection (2a) of this section may be made so as to apply—

(a) generally, or to a particular drug of addiction or particular specified drug or to particular classes thereof;

(b) generally, or to particular classes of persons,

and may make differing provisions as regards classes of persons and classes of drugs of addiction or specified drugs.

(3) Regulations made under the provisions of this section are in addition to and not in derogation of any regulations made under the Health Act 1911, and under the misuse of Drugs Act 1981, but where and to the extent that inconsistency exists between the regulations made under this section and any regulations made under the Health Act 1911 or the Misuse of Drugs Act 1981, as referred to in this subsection, the regulations made under this section shall prevail.

APPENDIX. "A"[1]

Substituted by *G.G.* 9/7/81, pp. 2725-68. Amended by *G.G.* 18/12/81, pp. 5190-5192; *G.G.* 16/7/82, pp. 2721-23; *G.G.* 13/8/82, pp. 3112-13.

First Schedule.

A substance specified in this Schedule includes any compound and all preparations and admixtures containing any proportion thereof and these are therefore subject to all the restrictions of this Schedule unless specifically exempted or specifically included in any other Schedule.

[1] Appendix "A" substituted see *G.G.* 9/7/81, pp. 2725-68. For previous amendments see Index to Legislation of Western Australia, Table 1.

ACONITE (ROOT OF ACONITUM NAPELLUS).

ANTIMONY and substances containing more than the equivalent of 1 per cent of antimony trioxide, except antimony chlorides in polishes.

ATROPINE and substances containing more than 0.25 per cent of atropine, except atropine methonitrate or when included in the Second Schedule.

BELLADONNA and substances containing more than 0.25 per cent of the alkaloids of belladonna calculated as hyoscyamine.

BROMINE as such.

BRUCINE and substances containing more than 0.2 per cent of brucine.

CONIINE and substances containing more than 0.1 per cent of coniine.

COTARNINE.

CROTON OIL.

HOMATROPINE and substances containing more than 0.25 per cent of homatropine.

HYDROCYANIC ACID, CYANIDES and substances for therapeutic use containing more than the equivalent of 0.15 per cent of hydrocyanic acid.

HYOSCINE and substances containing more than 0.25 per cent of hyoscine, except hyoscine N-butyl-bromide.

HYOSCYAMINE and substances containing more than 0.25 per cent of hyoscyamine.

HYOSCYAMUS and substances containing more than 0.25 per cent of alkaloids of hyoscyamus calculated as hyoscyamine.

LOBELIA and substances containing more than 0.5 per cent of the alkaloids of lobelia except preparations for smoking or burning.

MERCURIC CHLORIDE and substances containing more than 0.5 per cent of mercuric chloride, except when prepared and packed to comply with the requirements of the Sixth or Seventh Schedule.

MERCURIC IODIDE and substances containing more than 2 per cent of mercuric iodide, except when included in the Sixth Schedule.

MERCURIC NITRATE and substances containing mercuric nitrate equivalent to more than 3 per cent mercury (Hg).

MERCURIC-POTASSIUM IODIDE and substances containing mercuric-potassium iodide equivalent to more than 2 per cent of mercuric iodide.

MERCURIC THIOCYANATE except when included in the Sixth Schedule.

MERCURY, organic compounds of, and substances containing more than the equivalent of 0.5 per cent of mercury (Hg) in organic compounds, except for therapeutic use or when included in the Sixth Schedule.

NUX VOMICA.

PHOSPHORUS YELLOW and substances containing more than 0.5 per cent of free phosphorus.

SAVIN, oil of.

STRAMONIUM and substances containing more than 0.25 per cent of alkaloids calculated as hyoscyamine, except preparations for smoking or burning.

STRYCHNINE in preparations made for human therapeutic purposes.

TANSY, oil of.

VERATRUM, except for therapeutic use.

Excluding however, the substances hereinbefore mentioned when contained in any of the following—

Batteries and accumulators.
Ceramics.
Electrical components and electrical lamps.
Explosives.
Fireworks other than fireworks containing arsenic.
Glazed Pottery.
Inorganic Pigments.
Matches.
Motor fuels and lubricants.
Paints other than substances prepared for medicinal or cosmetic purposes.
Paper.
Photographic film.
Photographic paper.
Timber and wallboard.
Vitreous enamels.

Second Schedule.

A substance specified in this Schedule includes any compound and all preparations and admixtures containing any proportion thereof and these are therefore subject to all the restrictions of this Schedule unless specifically exempted or specifically included in any other Schedule.

ACETIC ACID (excluding its salts and its derivatives) in substances for therapeutic use containing more than 80 per cent of acetic acid.

ACETYLDIHYDROCODEINE when compounded with one or more other medicaments, in substances containing 1 per cent or less of acetyldihydrocodeine.

ANAESTHETICS LOCAL the following only—

(i) Benzamine;

(ii) Benzocaine;

(iii) Butylaminobenzoate;

(iv) Butylaminobenzoate picrate;

(v) Lignocaine,

when included in—

(a) lozenges, pastilles, tablets or capsules containing 30 mg or less of such substances;

(b) suppositories or bougies containing 200 mg or less of such substances;

(c) Preparations for external use, other than eye drops, containing 10 per cent or less of such substances.

ANTICHOLINERGIC SUBSTANCES for external use.

ANTIHISTAMINES, all tertiary nitrogenous organic bases which possess pharmacological properties characteristic of antihistamine compounds (except azatadine, chlorcyclizine, cyclizine, meclozine and methapyrilene) in—

(a) oral liquid preparations when compounded with an antitussive, an expectorant or a sympathomimetic amine and when packed, labelled and sold for the relief of coughs and colds and bearing a label warning that the product is not suitable for children under eight years of age;

(b) preparations labelled and packed as eye drops, nasal drops or nasal sprays for topical use;

(c) solid dose preparation containing only buclizine, cinnarizine, diphenhydramine, dimenhydrinate, pheniramine and promethazine as the antihistamine ingredient labelled and packed for the treatment of motion sickness in packs of ten doses or less.

ANTIMONY, in substances containing the equivalent of 1 per cent or less of antimony trioxide, except antimony chlorides in polishes.

ASPIRIN, except—

(a) in tablets or capsules each containing 325 mg or less of aspirin as the only therapeutically active constituent where the tablets or capsules are packed in blister or strip packaging or a container fitted with a child resistant closure, and the tablets or capsules are enclosed in a primary pack containing not more than 25 tablets or capsules;

(b) in individually wrapped powders or granules each containing 650 mg or less of aspirin as the only therapeutically active constituent where the powders or granules are contained in a primary pack containing not more than 12 individually wrapped powders or granules;

(c) when included in the Fourth Schedule.

ATROPINE, except atropine methonitrate, in substances containing 0.25 per cent or less of atropine, and atropine sulphate, 0.6 mg tablets in packs of six, when labelled for the treatment of organophosphorus poisoning.

BELLADONNA, in substances containing 0.25 per cent or less of the alkaloids of belladonna, calculated as hyoscyamine.

BROMHEXINE (N-cyclohexyl-N-methyl—(2-amino-3, 5-dibromobenzyl)—ammonium chloride).

BRUCINE, in substances containing 0.2 per cent or less of brucine, except when used in concentrations of 0.02 per cent or less for the denaturation of alcohol.

BUFEXAMAC in substances containing 5 per cent or less of bufexamac for external human therapeutic use, and in suppositories.

CANTHARIDES (CANTHARIDIN) in substances containing 0.01 per cent or less of cantharidin.

CARBARYL in preparations for external human therapeutic use containing 2 per cent or less of carbaryl.

CARBENOXOLONE in preparations for topical oral use.

CARBETAPENTANE CITRATE, except in preparations containing 0.5 per cent or less of carbetapentane citrate.

CHLOROFORM (excluding its derivatives) except—

(a) in preparations containing 10 per cent or less of chloroform where the chloroform content is declared on the label; or

(b) when included in the Fourth Schedule.

CINNAMEDRINE.

CLEMASTINE in oral dosage units of 1 mg or less in packs of 50 doses or less.

CODEINE when compounded—

(a) in tablets or capsules with either aspirin or paracetamol or salicylamide or either one of their derivatives and containing not more than 10 mg of codeine in each tablet or capsule when—

(i) packed in blister or strip packaging or in containers fitted with a child resistant closure;

and

(ii) in a primary pack containing not more than 25 such tablets or capsules;

(b) when compounded with aspirin or paracetamol or salicylamide in individually wrapped powders containing 10 mg or less of codeine in each individually wrapped powder where the powders are enclosed in a primary pack containing not more than 12 individually wrapped powders;

(c) with one or more other medicaments in divided preparations containing not more than 10 mg of codeine per dosage unit; or

(d) with one or more other medicaments in undivided preparations with a concentration of not more than 0.5 per cent of codeine.

CONIINE in substances containing 0.1 per cent or less of coniine.

DEXTROMETHORPHAN in substances containing 1 per cent or less of dextromethorphan when compounded with one or more other medicaments in such a way that the dextromethorphan contained therein cannot readily be extracted.

DEXTRORPHAN in substances containing 1 per cent or less of dextrorphan.

DIAMINES, phenylene, toluene and all other alkylated benzene diamine derivatives, except when included in the Sixth Schedule.

DICOPHANE (DDT) in substances for human therapeutic use.

DICYCLOMINE in substances for internal use containing 0.1 per cent or less of dicyclomine.

DIMETHISOQUIN in preparations for topical use.

DIPHEMANIL METHYLSULPHATE in substances for topical use.

EPHEDRINE AND PSEUDOEPHEDRINE in substances containing more than a total 0.5 per cent of ephedrine and pseudoephedrine except in substances for external use containing 1 per cent or less of ephedrine and pseudoephedrine.

ERYTHRITYL TETRANITRATE and other nitric esters of polyhydric alcohols.

ETAFEDRINE.

ETHER, except—

(a) in preparations containing 10 per cent or less of ether; or

(b) when included in the Fourth, Fifth or Sixth Schedules.

ETHOHEPTAZINE in substances containing 1 per cent or less of ethoheptazine.

ETHYLMORPHINE when compounded with one or more other medicaments in substances containing 1 per cent or less of ethylmorphine.

FERROUS SULPHATE and other iron preparations for human internal use, except in preparations containing 5 per cent or less of iron.

FLUORIDES, metallic including ammonium fluoride and fluorinated stannous compounds, when intended for therapeutic purposes, except—

(a) in dentifrices containing 0.5 per cent or less of fluoride ion;

(b) in substances containing 15 mg/kg or less of fluoride ion; or

(c) when included in the Fourth or Sixth Schedules.

GLYCERYL TRINITRATE.

GUAIPHENESIN.

(a) in liquid preparations containing 2 per cent or less of guaiphenesin; or

(b) in solid dose preparations containing 120 mg or less of guaiphenesin in each dosage unit.

HEXACHLOROPHANE in preparations for skin cleansing purposes containing 3 per cent or less of hexachlorophane, except—

(a) in preparations for use on infants; or

(b) in preparations for the treatment of animals.

HOMATROPINE in substances containing 0.25 per cent or less of homatropine.

HYDROCYANIC ACID AND CYANIDES in substances containing the equivalent of 0.15 per cent or less of hyrocyanic acid.

8-HYDROXYQUINOLINE and its derivatives for therapeutic use except when included in the Fourth or Sixth Schedules and except non-halogenated derivatives for external use containing 1 per cent or less of non-halogenated derivatives.

HYOSCINE and its derivatives in substances containing 0.25 per cent or less of hyoscine and/or its derivatives, except hyoscine N-butyl-bromide.

HYOSCYAMINE and its derivatives in substances containing 0.25 per cent or less of hyoscyamine and/or its derivatives.

HYOSCYAMUS in substances containing 0.25 per cent or less of alkaloids calculated as hyoscyamine.

IODINE, excluding its salts and derivatives, containing more than 2.5 per cent of available iodine except when included in the Sixth Schedule.

ISOPROPAMIDE in preparations containing 2 per cent or less of isopropamide for cutaneous use.

ISOSORBIDE DINITRATE.

LEAD SALTS and compounds of lead when prepared for medical or cosmetic use, except in substances for hair dressing containing the equivalent of 1 per cent or less of lead (Pb).

LINDANE in preparations for external human therapeutic use containing 2 per cent or less of lindane.

LOBELIA in substances containing 0.5 per cent or less of the alkaloids of lobelia, except substances for smoking or burning.

MALDISON in preparations for external human therapeutic use containing 2 per cent or less of maldison.

MEBENDAZOLE for human therapeutic use.

MERCURIC AMMONIUM CHLORIDE (AMMONIATED MERCURY).

MERCURIC CHLORIDE in substances containing 0.5 per cent or less of mercuric chloride, except when prepared and packed to comply with the requirements of the Sixth and Seventh Schedules.

MERCURIC IODIDE in substances containing 2 per cent or less of mercuric iodide, except when included in the Sixth Schedule.

MERCURIC NITRATE in substances containing the equivalent of 3 per cent or less of mercury (Hg).

MERCURIC OXIDE and all oxides of mercury.

MERCURIC POTASSIUM IODIDE in substances containing the equivalent of 2 per cent or less of mercuric iodide.

MERCURY (METALLIC), as such, except in scientific instruments.

MERCURY, organic compounds of, in substances containing the equivalent of 0.5 per cent or less of mercury (Hg) in organic combination, except—

(a) when included in the Fourth, Sixth or Seventh Schedules; or

(b) as a preservative in substances containing the equivalent of 0.01 per cent or less of mercury (Hg).

METHOXAMINE in substances containing more than 0.5 per cent of methoxamine, except preparations for external use containing 1 per cent or less of methoxamine.

METHOXYPHENAMINE.

N-METHYLEPHEDRINE.

NAPHAZOLINE.

NICOCODINE when compounded with one or more other medicaments in substances containing 1 per cent or less of nicocodine.

NICODICODINE when compounded with one or more other medicaments in preparations containing 1 per cent or less of nicodicodine.

NORCODEINE when compounded with one or more other medicaments in substances containing 1 per cent or less of norcodeine.

OXETHAZAINE.

OXOLAMINE.

OXYMETAZOLINE.

PAPAVERINE.

PARACETAMOL, except—

(a) in tablets or capsules each containing 500 mg or less of paracetamol as the only therapeutically active constituent where the tablets or capsules are packed in blister or strip packaging or in a container fitted with a child resistant closure, and the tablets or capsules are enclosed in a primary pack containing not more than 25 tablets or capsules;

(b) in individually wrapped powders or granules each containing 1 000 mg or less of paracetamol as the only therapeutically active constituent where the powders or granules are enclosed in a primary pack containing not more than 12 individually wrapped powders or granules;

(c) when included in the Fourth Schedule.

PHEDRAZINE.

PHENAMAZOLINE.

PHENAZONE for topical use, and substances containing 150 mg or less of phenazone prepared and labelled for the treatment of migraine and nauseating headaches.

PHENOL and any homologue of phenol boiling below 220°C, creosote, and substances containing more than 3 per cent by weight of such substances or homologues, for therapeutic use.

PHENYLEPHRINE in substances containing more than 0.5 per cent of phenylephrine, except substances for external use containing 1 per cent or less of phenylephrine.

PHENYLPROPANOLAMINE in tablets or capsules containing not more than 30 mg of phenylpropanolamine per tablet or capsule when not in sustained release form, and not more than 50 mg in each tablet or capsule when in sustained release form.

PHOLCODINE when compounded with one or more other medicaments in substances containing 1 per cent or less of pholcodine.

POTASSIUM CHLORATE and substances containing more than 10 per cent of potassium chlorate.

PROCYCLIDINE in preparations containing 5 per cent or less of procyclidine for cutaneous use.

PROPANTHELINE in substances for topical use.

PROPOXUR in preparations for external human therapeutic use containing 0.2 per cent or less of propoxur.

PROPYLHEXEDRINE in appliances for inhalation in which the substance is absorbed upon an inert solid material.

PROPYLPHENAZONE.

PYRANTEL for human therapeutic use.

RINIDOL.

SALICYLAMIDE, except—

(a) in tablets or capsules each containing 500 mg or less of salicylamide as the only therapeutically active constituent where the tablets or capsules are packed in blister or strip packaging or in a container fitted with a child resistant closure, and the tablets or capsules are enclosed in a primary pack containing not more than 25 tablets or capsules;

(b) in individually wrapped powders or granules each containing 1 000 mg or less of salicylamide as the only therapeutically active constituent where the powders or granules are enclosed in a primary pack containing not more than 12 individually wrapped powders or granules;

(c) when included in the Fourth Schedule.

SILVER NITRATE.

SODIUM NITRITE for therapeutic use.

STAVESACRE and substances containing more than 0.2 per cent of stavesacre.

STRAMONIUM in substances containing 0.25 per cent or less of the alkaloids calculated as hyoscyamine, except substances for smoking or burning.

TETRAHYDROZOLINE.

TRAMAZOLINE.

TRICLOFOS in preparations containing 5 per cent or less of triclofos.

TRIMIZOLINE.

TYMAZOLINE.

VIPRYNIUM.

XYLOMETAZOLINE.

ZINC PYRITHIONE, except in preparations containing 2 per cent or less of zinc pyrithione.

Excluding however, the substances hereinbefore mentioned when contained in any of the following—

Batteries and accumulators.

Ceramics.

Electrical components and electric lamps.

Explosives.

Fireworks other than fireworks containing arsenic.

Glazed pottery.

Inorganic pigments.

Matches.

Motor Fuels and lubricants.

Paints other than substances prepared for medicinal or cosmetic purposes.

Paper.

Photographic Film.

Photographic Paper.

Timber and wallboard.

Vitreous enamels.

Third Schedule.

A substance specified in this Schedule includes any compound and all preparations and admixtures containing any proportion thereof and these are therefore subject to all the restrictions of this Schedule unless specifically exempted or specifically included in any other Schedule.

ADRENALINE, natural or synthetic, in substances containing more than 0.01 per cent but not more than 1 per cent of adrenaline.

AMYL NITRITE.

ANTHISTAMINES, all tertiary nitrogenous organic bases which possess pharmacological properties characteristic of anihistamine compounds (except azatadine, chlorcyclizine, cyclizine, meclozine and methapyrilene)—

(a) in oral solid preparations containing 30 dosage units or less except when included in the Second Schedule;

(b) in oral liquid preparations labelled for the treatment of coughs and colds except when included in the Second Schedule.

BENZOYL PEROXIDE in preparations containing 10 per cent or less of benzoyl peroxide for external human therapeutic use.

BROMIDES, inorganic, in extemporaneous preparations for therapeutic use.

BUTYL NITRITE.

CHLORAL HYDRATE in substances containing 5 per cent or less of chloral hydrate, except alpha-chloralose when included in the Sixth Schedule.

CHOLESTYRAMINE for human therapeutic use.

CODEINE when compounded with aspirin or paracetamol or salicylamide in tablets or capsules containing 10 mg or less of codeine in each tablet or capsule or in individually wrapped powders containing 10 mg or less of codeine in each individually wrapped powder, except when included in the Second Schedule.

COLESTIPOL for human therapeutic use.

5,5 DIBROMO-O-CRESOLSULFONPHTHALEIN in solutions for testing for pregnancy.

DIHYDROCODEINE when compounded with one or more therapeutically active medicaments in substances containing 1 per cent or less of dihydrocodeine.

FENOTEROL in metered aerosols delivering 200 micrograms or less of fenoterol per metered dose.

FLAVOXATE.

FOLIC ACID for human use except in preparations containing 500 micrograms or less of folic acid per recommended daily dose.

FOLINIC ACID.

IDOXURIDINE in preparations for cutaneous use containing 0.5 per cent or less of idoxuridine.

INSULIN and substances containing the specific hypoglycaemic principle of the pancreas.

ISOBUTYL NITRITE.

ISOPRENALINE in preparations containing 1 per cent or less of isoprenaline except when contained in metered aerosols delivering more than 100 micrograms per metered dose.

MEFENAMIC ACID in packs of 30 capsules or less when labelled for treatment of spasmodic dysmenorrhea.

NITROFURAZONE for topical use.

NORADRENALINE, in substances containing 1 per cent or less of noradrenaline, except substances containing 0.01 per cent or less of noradrenaline.

NOSCAPINE.

OCTYL NITRITE.

ORCIPRENALINE in metered aerosols delivering 750 micrograms or less of orciprenaline per metered dose.

PREGNANCY TESTING KITS, and preparations and solutions for testing for pregnancy containing Human Chorionic Gonadotrophin bound to red blood cells and Human Chorionic Gonadotrophin antiserum, packed and labelled for use on one occasion only, and sold under the brand names "PREDICTOR", "EPT", and "DISCOVER 2".

QUININE for human therapeutic use.

SALBUTAMOL in metered aerosols delivering 100 micrograms or less of salbutamol per metered dose.

SANTONIN.

SODIUM CROMOGLYCATE in liquid nasal preparations topically applied.

TERBUTALINE in metered aerosols delivering 250 micrograms or less of terbutaline per metered dose.

TRETINOIN for external human therapeutic use.

Fourth Schedule.

A substance specified in this Schedule includes any compound and all preparations and admixtures containing any proportion thereof and these are therefore subject to all the restrictions of this Schedule unless specifically exempted or specifically included in any other Schedule.

ACEDAPSONE.

ACETANILIDE and alkyl acetanilides, for human therapeutic use.

ACETAZOLAMIDE.

ACETOHEXAMIDE.

ACETYLCHOLINE and other choline esters.

ACETYLCYSTEINE.

ACETYLDIHYDROCODEINE when compounded with one or more other medicaments and containing 100 mg or less of acetyldihydrocodeine per dosage unit and with a concentration of more than 1 per cent and not more than 2.5 per cent of acetyldihydrocodeine in undivided preparations.

ACETYL METHYL DIMETHYL OXIMIDO PHENYL HYDRAZINE.

ADIPHENINE.

ADRENALINE, natural or synthetic, in substances containing more than 1 per cent of adrenaline.

ALCURONIUM.

ALLYLISOPROPYLACETYLUREA.

ALPHADOLONE.

ALPHA-RECEPTOR BLOCKING AGENTS including phentolamine and phenoxybenazamine.

ALPHAXOLONE.

ALPRENOLOL.

AMANTADINE.

AMBENONIUM.

AMBUCETAMIDE.

AMBUTONIUM.

AMETHOCAINE.

AMIDOPYRINE, its salts, its derivatives and their salts.

AMIKACIN.

AMILORIDE.

AMINOMETRADINE.

AMINOREX.

AMIPHENAZOLE.

AMISOMETRADINE.

AMITRIPTYLINE and other compounds structurally derived therefrom by substitution in the side chain.

AMODIAQUINE.

AMOXYCILLIN.

AMPHOMYCIN.

AMPHOTERICIN.

AMPICILLIN.

AMYGDALIN (Laetrile).

AMYLOCAINE.

ANABOLIC steroidal agents.

ANAESTHETICS: the following when specifically prepared and packed as therapeutic agents for the induction of inhalation anaethesia.

Chloroform.

Cyclopropane.

Ether.

Ethyl Chloride.

Ethylene.

Fluroxone.

Halothane.

Methoxyflurane.

Nitrous Oxide.

Trichloroethylene.

Vinyl ether.

ANAESTHETICS LOCAL, being synthetic cocaine substitutes, except when included in the Second Schedule.

ANGIOTENSIN AMIDE.

ANTAZOLINE, except when included in the Second or Third Schedules.

ANTIBIOTICS, penicillin, penicillanic acid, streptomycin, chloramphenicol, tetracycline, and any other antibiotic substances however derived and their chemical derivatives, except when included in the Sixth Schedule.

ANTIFOLIC ACID SUBSTANCES including aminopterin, teropterin and orthopterin.

ANTIHISTAMINES, all tertiary nitrogenous organic bases which possess pharmacological properties characteristic of antihistamine compounds not elsewhere specified in these schedules except when included in the Second, Third or Seventh Schedule.

ANTIMALARIAL SUBSTANCES, except Quinine, not elsewhere specified in the Schedules.

ANTIMONY, organic compounds of, for therapeutic use.

ANTITUBERCULOSIS SUBSTANCES including isoniazid and its derivatives, para-aminosalicylic acid and its salts, and thiacetazone.

APOMORPHINE.

APROTININ.

ARSENIC, for human therapeutic use.

ASPIRIN, when compounded with caffeine, or paracetamol or salicylamide.

ATENOLOL.

ATROPINE METHONITRATE.

AZAPERONE.

AZAPETINE.

AZATADINE.

BACITRACIN except—

(a) when included in the Sixth Schedule;

(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle; or

(c) in milk replacers for calves and starter rations for pigs in concentrations of 100 mg/kg or less of the total active antibiotic principle.

BACLOFEN.

BAMIPINE, except when included in the Second or Third Schedules.

BARBITURIC ACID.

BECLAMIDE.

BEMEGRIDE.

BENACTYZINE and other substances structurally derived from diphenyl-methane with ataractic properties when used for therapeutic purposes.

BENSERAZIDE.

BENZAMINE, except when included in the Second Schedule.

BENZHEXOL.

BENZILONIUM.

BENZOCAINE, except when included in the Second Schedule.

BENZOYL PEROXIDE in preparations for external human therapeutic use except when included in the Third Schedule.

BENZPHETAMINE and other substances structurally derived from beta-aminopropylbenzene or beta-aminoisopropylbenzene by substitution in the side chain or by ring closure therein (or both such substitution and such closure), except when included in the Second or Eighth Schedule and except ephedrine, pseudoephedrine and phenylephrine in substances exempted from the Second Schedule.

BENZTROPIN.

BENZYDAMINE.

BENZYLPENICILLIN (including procaine penicillin) except when included in the Sixth Schedule.

BETHAHISTINE.

BETA-RECEPTOR BLOCKING AGENTS not otherwise included in these Schedules.

BETHANIDINE.

BIPERIDEN.

BISMUTH SUBGALLATE for oral use in humans.

BLEOMYCIN.

BORON compounds for human therapeutic or cosmetic use except:—

(a) in preparations for external use containing 1 per cent or less of boron; or

(b) in unit dose preparations for periodontal disease containing 100 mg or less of boron.

BRETYLIUM.

BROMIDES, inorganic, for therapeutic use, except in extemporaneous preparations.

BROMOCRIPTINE.

BROMODIPHENHYDRAMINE, except when included in the Second or Third Schedules.

BROMOFORM for therapeutic use.

BROMPHENIRAMINE, except when included in the Second or Third Schedules.

BROMVALETONE.

BUCLIZINE, except when included in the Second or Third Schedules.

BUFEXAMAC, except when included in the Second Schedule.

BUMETANIDE.

BUSULFAN.

BUTACAINE.

BUTYLAMINOBENZOATE, except when included in the Second Schedule.

BUTYL CHLORAL HYDRATE.

CALCITONIN.

CALCITRIOL.

CALCIUM CARBIMIDE for therapeutic use.

CAMPHORATED OIL.

CAMPHOTAMIDE.

CANDIDICIN.

CANTHARIDES and substances containing more than 0.01 per cent of cantharidin.

CANTRODIFENE.

CAPREOMYCIN.

CAPTODIAME.

CAPURIDE.

CARAMIPHEN.

CARBACHOL.

CARBAMAZEPINE.

CARBARYL, for human therapeutic use except when included in the Second Schedule.

CARBAZOCHROME.

CARBENICILLIN.

CARBENOXOLONE for human therapeutic use, except when included in the Second Schedule.

CARBIDOPA.

CARBIMAZOLE.

CARBINOXAMINE, except when included in the Second or Third Schedules.

CARBOCROMEN.

CARBROMAL.

CARDIAC glycosides not elsewhere specified in these Schedules.

CARINDACILLIN.

CEFOXITIN.

CEPHACETRILE.

CEPHALEXIN.

CEPHALORIDINE.

CEPHALOTHIN.

CEPHAMANDOLE.

CEPHAPIRIN.

CEPHAZOLIN.

CEPHRADINE.

CETOXIME, except when included in the Second or Third Schedules.

CHENODEOXYCHOLIC ACID.

CHINIOFON and other halogenated 8-hydroxyquinoline derivatives for internal human use.

CHLORAL FORMAMIDE.

CHLORAL HYDRATE and substances containing more than 5 per cent of the equivalent of chloral hydrate.

CHLORAMPHENICOL.

CHLORAZANIL.

CHLORBUTOL, in preparations for human oral use except in preparations containing 0.5 per cent or less of chlorbutol as a preservative.

CHLORCYCLIZINE.

CHLORDIAZEPOXIDE and other substances structurally derived from benzodiazepine with ataractic properties when used for therapeutic purposes.

CHLORMERODRIN.

CHLORMETHIAZOLE.

CHLORMEZANONE.

CHLOROBENZYL DISULPHONAMIDE.

1-(4-CHLOROPHENOXY)-1-IMIDAZOL-1-YL-3,3-DIMETHYL-2-BUTANONE for human use.

CHLOROPYRILENE, except when included in the Second or Third Schedules.

CHLOROQUINE.

CHLOROTHIAZIDE and other substances structurally derived from benzothiadazine for therapeutic use.

CHLORPHENIRAMINE, except when included in the Second or Third Schedules.

CHLORPHENOXAMINE, except when included in the Second or Third Schedules.

CHLORPHENTERMINE.

CHLORPROMAZINE and other substances structurally derived from phenothiazine with ataractic properties when used for therapeutic purposes.

CHLORPROPAMIDE.

CHLORPROTHIXENE.

CHLORTERACYCLINE, except when included in the Sixth Schedule.

CHLORTHALIDONE.

CHLORZOXAZONE.

CIMETIDINE.

CINCHOCAINE.

CINNARAZINE, except when included in the Second or Third Schedules.

CISPLATIN.

CLEMASTINE, except when included in the Second or Third Schedules.

CLEMIZOLE, except when included in the Second or Third Schedules.

CLIDINIUM.

CLINDAMYCIN.

CLOBETASONE—17—BUTYRATE EUMOVATE.

CLOFENAMIDE.

CLOFENOXINE.

CLOFIBRATE.

CLOMOCYCLINE.

CLONAZEPAM.

CLONIDINE.

CLOPAMIDE.

CLOPROSTENOL.

CLORAZEPATE.

CLOREXOLONE.

CLOTRIMAZOLE.

CLOXACILLIN.

CLOZOPIN.

CODEINE when compounded with one or more other medicaments—

(a) in divided preparations containing 30 mg or less of codeine per dosage unit; or

(b) in undivided preparations with a concentration of 1 per cent or less of codeine,

except when included in the Second or Third Schedules.

COLASPASE.

COLCHICINE.

COLISTIN.

CORTISONE and steroid suparenal cortical hormones, either natural or synthetic.

COUMARIN derivatives and phenylindanedione derivatives for therapeutic use.

CURARE, TUBOCURARINE, d-TUBOCURARINE, d-TUBOCURARINE-DIMETHYL-ETHER, and all synthetic quatenary ammonium compounds and other compounds having curarising properties.

CYCLANDELATE.

CYCLIRAMINE, except when included in the Second or Third Schedules.

CYCLIZINE.

CYCLOPENTOLATE.

CYCLOSERINE.

CYCRIMINE.

CYPROHEPTADINE, except when included in the Second or Third Schedules.

DACARBAZINE.

DANAZOL.

DAPSONE and all derivatives of 4,4'-diaminodiphenylsulphone.

DEANOL.

DEBRISOQUINE.

DEMACARIUM BROMIDE.

DEMECLOCYCLINE.

DEPTROPINE, except when included in the Second or Third Schedules.

DESIPRAMINE.

DESMOPRESSIN (D.D.A.V.P.).

DEXBROMPHENIRAMINE, except when included in the Second or Third Schedules.

DEXCHLORPHENIRAMINE, except when included in the Second or Third Schedules.

DEXTROMETHORPHAN, except when included in the Second Schedule.

DEXTROPROPOXYPHENE.

DEXTRORPHAN and substances containing more than 1 per cent of dextrorphan.

DIBENZEPIN.

DIBUTAMIDE.

DICHLORPHENAMIDE.

DICHLORALPHENAZONE.

DICLOFENAC.

DICYCLOMINE and substances containing more than 0.1 per cent of dicyclomine.

DIETHAZINE.

DIETHYLCARBAMAZINE for human therapeutic use.

DIETHYLPROPION.

DIFENOXIN in preparations containing 0.5 mg or less of difenoxin per dosage unit and a quantity of atropine sulphate equivalent to at least 5 per cent of the dose of difenoxin.

DIFLUNISAL.

DIGITALIS and its glycosides.

DIHYDRALLAZINE.

DIHYDROCODEINE when compounded with one or more other medicaments in preparations containing 100 mg or less of dihydrocodeine per dosage unit and with a concentration of more than 1 per cent and not more than 2.5 per cent of dihydrocodeine in undivided preparations.

DIHYDROSTREPTOMYCIN, except when included in the Sixth Schedule.

DI-ISOPROPYLAMINE DICHLOROACETATE.

DIMENHYDRINATE, except when included in the Second or Third Schedules.

DIMETHINDENE, except when included in the Second or Third Schedules.

DIMETHISOQUIN, except when included in the Second Schedule.

DIMETHOTHIAZINE, except when included in the Second or Third Schedules.

DIMETHOXANATE.

DIMETHYL SULPHOXIDE for therapeutic use.

DINITROCRESOLS for therapeutic use.

DINITRONAPHTHOLS, for therapeutic use.

DINITROPHENOLS, for therapeutic use.

DINITROTHYMOLS for therapeutic use.

DIPERODON.

DIPHEMANIL METHYLSULPHATE, except substances for topical use.

DIPHENHYDRAMINE, except when included in the Second or Third Schedules.

DIPHENIDOL.

DIPHENOXYLATE in preparations containing 2.5 mg or less of diphenozylate calculated as base per dosage unit, and a quantity of atropine sulphate equivalent to at least 1 per cent of the dose of diphenoxylate.

DIPHENYLPYRALINE, except when included in the Second or Third Schedules.

DIPIVEFRIN.

DIPYRIDAMOLE.

DISOPHENOL.

DISOPYRAMIDE.

DISULFIRAM for therapeutic use.

DITHIAZANINE, except substances containing 2 per cent or less of dithiazanine for veterinary use.

DITOPHAL.

DOBUTAMINE.

DOPAMINE.

DOTHIEPIN.

DOXAPAM.

DOXEPIN.

DOXORUBICIN.

DOXYCYCLINE.

DOXYLAMINE, except when included in the Second or Third Schedules.

DROPERIDOL.

DROSTANOLONE.

DYFLOS.

ECONAZOLE, except when included in the Sixth Schedule.

EMBRAMINE, except when included in the Second or Third Schedules.

EMETINE and substances containing more than 0.2 per cent of emetine.

ENFLURANE.

EPICILLIN.

ERGOT, its alkaloids, their salts, derivatives of such alkaloids, and their salts.

ERYTHROMYCIN except—

(a) when included in the Sixth Schedule;

(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle; or

(c) in milk replacers for calves or starter rations for pigs in concentrations of 100 mg/kg or less of the total active antibiotic principle.

ETHACRYNIC ACID.

ETHAMBUTOL.

ETHAMIVAN.

ETHCHLORVYNOL.

ETHINAMATE.

ETHOGLUCIDE.

ETHOHEPTAZINE and substances containing more than 1 per cent of ethoheptazine.

ETHOPROPAZINE.

ETHOXZOLAMIDE.

ETHYL CHLORIDE for therapeutic use.

ETHYLMORPHINE when compounded with one or more other medicaments in preparations containing not more than 100 mg of ethylmorphine per dosage unit and with a concentration of more than 1 per cent and not more than 2.5 per cent of ethylmorphine in undivided preparations.

ETHYLOESTRENOL.

ETIDOCAINE.

FENCAMFAMIN.

FENFLURAMINE.

FENOPROFEN.

FENOTEROL except when included in the Third Schedule.

FENPIPRAMIDE.

FENPIPRANE.

FLAVOPHOSPHOLIPOL except—

(a) when included in the Sixth Schedule; or

(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle.

FLUCLOXACILLIN.

FLUCYTOSINE.

FLUFENAMIC ACID.

FLUNITRAZEPAM.

FLUORIDES including ammonium fluoride and fluorinated stannous compounds for therapeutic use containing more than the equivalent of 3 per cent of fluorine.

5-FLUOROCYTOSINE.

FLUOROURACIL and other substances structurally derived from uracil with cytotoxic properties, when used for therapeutic purposes.

FLUOXYMESTERONE.

FLURAZEPAM.

FLUSPIRILENE.

FRAMYCETIN.

FRUSEMIDE.

FUSIDIC ACID.

GALANTHAMINE.

GALLAMINE.

GENTAMICIN.

GLIBENCLAMIDE.

GLIBORNURIDE.

GLICLAZIDE.

GLUCAGON.

GLUTETHIMIDE.

GLYCOPYRROLATE.

GLYMIDINE.

GRAMICIDIN.

GRISEOFULVIN.

GUANACLINE.

GUANETHIDINE.

HALCINONIDE.

HALOPERIDOL and other substances structurally derived from butyrophenone with ataratic properties, when used for therapeutic purposes.

HALOPYRAMINE, except when included in the Second or Third Schedules.

HALOTHANE.

HEPARIN.

HETACILLIN.

HEXACHLOROPHANE, and substances containing hexachlorophane for use on infants, and hexachlorophane in all other substances except when included in the Second or Sixth Schedule.

HEXAMETHONIUM.

HEXOCYCLIUM.

HISTAPYRRODINE, except when included in the Second or Third Schedules.

HOMOCHLORCYCLIZINE, except when included in the Second or Third Schedules.

HYDRALLAZINE.

HYDROQUINONE for human therapeutic use in substances containing more than 2 per cent of hydroquinone.

HYDROXYCHLOROQUINE.

1-HYDROXY-PYRIDO (3,2 a)-5-PHENOXAZONE-3-CARBOXYLIC ACID.

HYDROXYUREA.

HYDROXYZINE.

HYGROMYCIN B, except when included in the Sixth Schedule or in preparations containing 5 mg/kg or less of hygromycin B.

HYOSCINE N-BUTYLBROMIDE.

HYPOTHALAMIC RELEASING FACTORS when used for diagnostic purposes.

IBUFENAC.

IBUPROFEN.

IDOXURIDINE, except when included in the Third Schedule.

IMIPRAMINE.

INDOMETHACIN.

INOSITOL NICOTINATE for internal use.

ION EXCHANGE RESINS, anionic and cationic, for internal use in humans, except when used as an excipient in tablets and capsules and except when included in the Third Schedule.

IPRATROPIUM BROMIDE.

IRON COMPOUNDS in injectable preparations for human therapeutic use.

ISOAMINILE.

ISOAMYLAMINE-METHYLHEPTAN.

ISOETHARINE.

ISOMETHEPTENE.

ISOPRENALINE, except when included in the Third Schedule.

ISOPROPAMIDE, except when included in the Second Schedule.

ISOXUPRINE.

KANAMYCIN.

KETAMINE.

KETOPROFEN.

KHELLIN.

KITASAMYCIN except—

(a) when included in the Sixth Schedule; or

(b) in animal feedstuffs for growth promotion in concentrations of 100 mg/kg or less of the total active antibiotic principle.

LABETOLOL.

LAUDEXIUM METHYL SULPHATE.

LEFETAMINE.

LEPTAZOL.

LEVAMISOLE for human therapeutic use.

LEVODOPA.

LIDOFLAZINE.

LIGNOCAINE, except when included in the Second Schedule.

LINCOMYCIN.

LITHIUM salts and substances for therapeutic use containing more than the equivalent of 0.01 per cent of lithium (Li).

LOPERAMIDE.

LORAZEPAM.

LOXAPINE.

LYMECYCLINE.

MAPHENIDE.

MAZINDOL.

MEBEVERINE.

MEBHYDROLIN, except when included in the Second or Third Schedules.

MECAMYLAMINE.

MECLOFENOXATE.

MECLOZINE.

MEDAZEPAM.

MEFENAMIC ACID, except when included in the Third Schedule.

MEFRUSIDE.

MEPACRINE.

MEPENZOLATE.

MEPHENESIN and its derivatives, except guaiphenesin when included in the Second Schedule.

MEPHENTERMINE.

MEPIVACAINE.

MEPROBAMATE.

MEPYRAMINE, except when included in the Second or Third Schedules.

MERCAPTOPURINE and other substances structurally derived therefrom with cytotoxic properties when used for therapeutic purposes.

MERCUROUS CHLORIDE for therapeutic use.

MERCURY, organic compounds of, for therapeutic use, except substances for topical use containing the equivalent of 0.5 per cent or less of mercury (Hg).

METARAMINOL.

METFORMIN.

METHACYCLINE.

METHADIENONE.

METHANDRIOL.

METHANTHELINE.

METHAPHENILINE, except when included in the Second or Third Schedules.

METHAZOLAMIDE.

METHDILAZINE, except when included in the Second or Third Schedules.

METHENOLONE.

METHICILLIN.

METHIMAZOLE.

METHIXENE.

METHOCARBAMOL.

METHOTREXATE.

METHOXSALEN.

METHYLANDROSTANOLONE.

METHYLDOPA.

METHYLPENTYNOL and other substituted alkynes for internal use.

METHYLPERIDOL.

METHYPRYLONE.

METOCLOPRAMIDE.

METOLAZONE.

METOPROLOL.

METRIZAMIDE.

METRONIDAZOLE.

METYRAPONE.

MEXILITINE HYDROCHLORIDE.

MEZLOCILLIN.

MIANSERIN.

MIBOLERONE.

MICONAZOLE.

MINOCYCLINE.

MINOXIDIL.

MITABRONITOL.

MITHRAMYCIN.

MITOMYCIN.

MONENSIN, except in animal feeds containing 120 mg/kg or less of monensin.

MONO-AMINE OXIDASE INHIBITORS, including iproniazid, isocarboxazid, nialamide, phenelzine, pheniprazine and other substances for which mono-amine oxidaze inhibition is claimed, except triparanol.

MONOBENZONE and substances containing more than 2 per cent of monobenzone, for human therapeutic use.

MOPERONE.

MORPHINE ANTAGONISTS including nalorphine, levallorphine and naloxone.

MUSTINE and other substances structurally derived therefrom with cytotoxic properties, when used for therapeutic purposes.

NALIDIXIC ACID.

NANDROLONE.

NAPROXEN.

NEOMYCIN, except when included in the Sixth Schedule.

NEOSTIGMINE.

NICOCODINE when compounded with one or more other medicaments in substances containing 100 mg or less of nicocodine per dosage unit and with a concentration of more than 1 per cent and not more than 2.5 per cent of nicocodine in undivided preparations.

NICODICODINE when compounded with one or more other medicaments in preparations containing 100 mg or less of nicodicodine per dosage unit and with a concentration of more than 1 per cent and not more than 2.5 per cent of nicodicodine in undivided preparations.

NICOTINE in chewing tablets containing 4 mg or less of nicotine per tablet for use as an aid in withdrawal from tobacco smoking.

NICOTINIC ACID, where the recommended daily dose exceeds 250 mg.

NICOTINYL ALCOHOL, for internal use.

NIFENAZONE.

NIKETHAMIDE.

NIRIDAZOLE.

NITRAZEPAM.

NITROFURAN for therapeutic use in humans.

NORADRENALINE and substances containing more than 1 per cent of noradrenaline.

NORCODEINE when compounded with one or more other medicaments and containing 100 mg or less of norcodeine per dosage unit and with a concentration of more than 1 per cent and not more than 2.5 per cent of norcodeine in undivided preparations.

NORETHANDROLONE.

NORTRIPTYLINE

NOVOBIOCIN except—

(a) when included in the Sixth Schedule; or

(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle.

OCTAMYLAMINE.

OCTATROPINE.

OLEANDOMYCIN except—

(a) when included in the Sixth Schedule; or

(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle.

ORCIPRENALINE, except when included in the Third Schedule.

ORGANO-PHOSPHORUS COMPOUNDS with anticholinesterase activity for human therapeutic use except when included in the Second Schedule.

ORNIDAZOLE.

ORNIPRESSIN.

ORPHENADRINE.

ORTHOCAINE.

OXACILLIN.

OXANDROLONE.

OXAZEPAM.

OXPRENOLOL.

OXYBUPROCAINE.

OXYMESTERONE.

OXYMETHOLONE.

OXYPHENBUTAZONE.

OXYPHENCYCLIMINE.

OXYPHENONIUM.

OXYTETRACYCLINE, except when included in the Sixth Schedule.

PAMAQUINE.

PANCURONIUM.

PARACETAMOL when compounded with aspirin or caffeine or salicylamide.

PARALDEHYDE.

PARAMETHADIONE.

PAROMOMYCIN.

PEMOLINE.

PEMPIDINE.

d-PENICILLAMINE.

PENTAMETHONIUM.

PENTHIENATE.

PENTOLINIUM.

PERHEXILINE.

PHENACEMIDE and other substances structurally derived from acetylurea with anticonvulsant properties, when used for therapeutic purposes.

PHENACETIN.

PHENAZONE, except when included in the Second Schedule.

PHENAZOPYRIDINE.

PHENETHICILLIN, except when included in the Sixth Schedule.

PHENFORMIN.

PHENGLUTARIMIDE.

PHENINDAMINE, except when included in the Second or Third Schedules.

PHENIRAMINE, except when included in the Second or Third Schedules.

PHENOXYBENZAMINE.

PHENOXYMETHYL PENICILLIN, except when included in the Sixth Schedule.

PHENSUXIMIDE and other substances structurally derived from succinamide with anticonvulsant properties when used for therapeutic purposes.

PHENTERMINE.

PHENTHIMENTONIUM.

PHENYAPIN.

PHENYLBUTAZONE.

PHENYLPROPANOLAMINE, except when included in the Second Schedule.

PHENYLTOLOXAMINE, except when included in the Second or Third Schedules.

PHENYTOIN and other substances structurally derived from hydantoin with anticonvulsant properties, when used for therapeutic purposes.

PHOLCODINE when compounded with one or more other medicaments and containing 100 mg or less of pholcodine per dosage unit and with a concentration of more than 1 per cent and not more than 2.5 per cent of pholcodine in undivided preparations.

PHYSOSTIGMINE.

PICROTOXIN.

PILOCARPINE and substances containing more than 0.025 per cent of pilocarpine.

PIMOZIDE.

PINDOLOL.

PIPENZOLATE.

PIPERIDOLATE.

PIPOBROMAN.

PIPRADROL.

PITUITARY, its extracts, its active principles and their synthetic substitutes, except when included in the Seventh Schedule.

PIZOTIFEN.

POLYMETHYLENE BISTRIMETHYL AMMONIUM COMPOUNDS.

POLYMYXIN.

POTASSIUM PERCHLORATE for therapeutic use.

PRACTOLOL.

PRAZEPAM.

PREGNANCY TESTING KITS and preparations and solutions for testing for pregnancy unless specifically included in the Third Schedule.

PREGNENOLONE ACETATE, except in substances for topical use.

PRENYLAMINE.

PRILOCAINE.

PRIMAQUINE.

PRIMIDONE.

PROBENECID.

PROCAINAMIDE.

PROCAINE.

PROCARBAZINE.

PROCHLORPERAZINE.

PROCYCLIDINE, except when included in the Second Schedule.

PROGUANIL.

PROLINTANE.

PROMETHAZINE, except when included in the Second or Third Schedules.

PROMIZOLE.

PROPANIDID.

PROPANTHELINE, except in substances for topical use.

PROPRANOLOL.

PROPYLHEXEDRINE, except when included in the Second Schedule.

PROQUAZONE.

PROSTAGLANDINS for veterinary use.

PROTHIONAMIDE.

PROXIMETACAINE.

PYRATHIAZINE, except when included in the Second or Third Schedules.

PYRIDOSTIGMINE.

PYRIMETHAMINE.

PYROXAMINE, except when included in the Second or Third Schedules.

PYRROBUTAMINE, except when included in the Second or Third Schedules.

QUINETHAZONE.

QUINIDINE.

RAUWOLFIA, its alkaloids, their salts, derivatives of such alkaloids, and their salts.

RIFAMPICIN.

RITORDRINE.

ROLITETRACYCLINE.

SALBUTAMOL except when included in the Third Schedule.

SALCATONIN.

SALICYLAMIDE when compounded with aspirin or caffeine or paracetamol.

SELENIUM except—

(a) when included in the Fifth or Sixth Schedules;

(b) in animal feedstuffs in concentrations of 0.1 g/tonne or less of selenium; or

(c) in compressed pellets for the control of selenium —responsive conditions in sheep.

SEX HORMONES, natural or synthetic and their substitutes, in all substances including cosmetics; except their derivatives and their substitutes without sex hormonal activity.

SILVER SULPHADIAZINE.

SISOMYCIN.

SODIUM CROMOGLYCATE except when included in the Third Schedule.

SODIUM FLUORIDE in preparations for human ingestion containing more than 2.2 mg sodium fluoride per dosage unit.

SODIUM NITROPRUSSIDE for human therapeutic use.

SODIUM VALPROATE.

SONTOQUINE.

SPARTEINE.

SPECTINOMYCIN.

SPIRAMYCIN except—

(a) when included in the Sixth Schedule;

(b) in animal feedstuffs for growth promotion in pigs or poultry in concentrations of 50 mg/kg or less of the total antibiotic principle.

SPIRONOLACTONE.

STANOLONE.

STANOZOLOL.

STREPTOMYCIN except when included in the Sixth Schedule.

STRYCHNINE in preparations containing 1.5 per cent or less of Strychnine for the treatment of animals.

STROPHANTHUS and its glycosides and their derivatives.

SULINDAC.

SULPHANILAMIDE, and its derivatives except—

(a) when included in the Sixth Schedule; or

(b) Sulphaquinoxaline when incorporated in baits for destruction of vermin and in animal feedstuffs containing 200 mg/kg or less of Sulphaquinoxaline.

(c) Oryzalin.

SULPHINPYRAZONE.

SULPHOMYXIN.

SULPHONAL and alkyl sulphonals.

SULTHIAME.

SUXAMETHONIUM.

TACRINE.

TAMOXIFEN.

TEMAZEPAM.

TERBUTALINE, except when included in the Third Schedule.

TESTOSTERONE PROPIONATE, TESTOSTERONE DIPROPIONATE AND TESTOSTERONE ENANTHATE except when included in the Sixth Schedule.

TETRABENAZINE.

TETRACYCLINE, except when included in the Sixth Schedule.

THENALDINE, except when included in the Second or Third Schedules.

THENYLDIAMINE, except when included in the Second or Third Schedules.

THIAMBUTOSINE.

THIOTEPA and other substances structurally derived therefrom with cytotoxic properties, when used for therapeutic purposes.

THIOTHIXENE.

THIOURACIL and substances structurally derived therefrom with antithyroid properties, when used for therapeutic purposes.

THIOUREA for therapeutic use.

THONZYLAMINE, except when included in the Second or Third Schedules.

THYROID and its extract, and its active principles.

TIAMULIN.

TICARCILLIN.

TIEMONIUM.

TIGLODINE.

TIMOLOL.

TINIDAZOLE.

TIPEPIDINE.

TOLAZAMIDE.

TOLAZOLINE for internal use.

TOLBUTAMIDE.

TOLPROPAMINE, except when included in the Second or Third Schedules.

TRANEXAMIC ACID.

TRETAMINE.

TRIAMTERENE.

TRIAZIQUONE.

TRIAZOLAM.

TRICLOFOS, except when included in the Second Schedule.

TRICYCLAMOL.

TRIDIHEXETHYL.

TRIFLUPERIDOL.

TRIMEPRAZINE, except when included in the Second or Third Schedules.

TRIMETAPHAN.

TRIMETHOBENZAMIDE, except when included in the Second or Third Schedules.

TRIMETHOPRIM.

TRIMIPRAMINE, and other compounds structurally derived therefrom by substitution in the side chain.

TRIMUSTINE.

TRIOXSALEN.

TRIPELENNAMINE, except when included in the Second or Third Schedules.

TRIPERIDOL.

TRIPROLIDINE, except when included in the Second or Third Schedules.

TROXIDONE, and other substances structurally derived from oxazolidone with anticonvulsant properties, when used for therapeutic purposes.

TYLOSIN except—

(a) when included in the Sixth Schedule; or

(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle; or

(c) in milk replacers for calves or starter rations for pigs in concentrations of 100 mg/kg or less of the total active antibiotic principle.

URETHANE for therapeutic use.

URETHANES and UREIDES having or purporting to have soporific, hypnotic or narcotic properties not specifically included in this or any other Schedule.

VACCINES, SERA, TOXOIDS, antitoxins and antigens for human parenteral use except when specified in the Seventh Schedule.

VACCINES, live virus for veterinary use.

VALNOCTAMIDE.

VERATRUM for therapeutic use.

VIDARABINE.

VINCA ALKALOIDS, including semi-synthetic derivatives.

VIRGINIAMYCIN except—

(a) when included in the Sixth Schedule; or

(b) in animal feedstuffs for growth promotion in concentrations of 50 mg/kg or less of the total active antibiotic principle.

VISNADINE.

VITAMIN A in preparations containing more than 10 000 international units per recommended daily dosage for human use.

VITAMIN D in preparations containing more than 25 micrograms per recommended daily dosage for human use.

XANTHINE OXIDASE INHIBITORS, including allopurinol.

XANTHINOL NICOTINATE.

XYLAZINE.

YOHIMBA, its alkaloids and their salts.

ZENAROL.

Excluding however, the substances hereinbefore mentioned when contained in any of the following—

Batteries and accumulators.

Ceramics.

Electrical components and electric lamps.

Explosives.

Fireworks other than fireworks containing arsenic.

Glazed pottery.

Inorganic pigments.

Matches.

Motor fuels and lubricants.

Paints other than substances prepared for medicinal or cosmetic purposes.

Paper.

Photographic Film.

Photographic Paper.

Timber and wallboard.

Vitreous enamels.

Fifth Schedule.

Hazardous substances.

ACETIC ACID in substances containing 80 per cent or less and more than 30 per cent of acetic acid, except for therapeutic use.

ACETIC ANHYDRIDE in substances containing 80 per cent or less and more than 30 per cent of acetic anhydride.

ACETONE and substances containing more than 25 per cent of acetone when packed in containers of more than 60 ml but not more than 20 litres.

AKLOMIDE and substances containing aklomide.

ALACHLOR and substances containing alachlor.

ALKALINE SALTS, being sodium carbonate, sodium orthosilicate, sodium metasilicate or trisodium phosphate or any combination of any two or more thereof, except—

(a) in preparations containing 10 per cent or less of combined substances;

(b) in solid preparations the pH of which in 1 per cent (w/v) aqueous solution is 11.5 or less; or

(c) in liquid preparations having a pH of 11.5 or less.

ALLOXYDIM SODIUM and substances containing alloxydim sodium.

Alpha-(2-CHLOROPHENYL)-alpha-(4-CHLOROPHENYL)-5-PYRIMIDINEMETHANOL and substances containing alpha-(2-chlorophenyl)-alpha-(4-chlorophenyl)-5-pyrimidinemethanol.

AMITROLE and substances containing amitrole.

AMMONIA in substances containing more than 0.5 per cent but not more than 5 per cent by weight of free ammonia (NH_3) except in medicinal substances for internal use, or when used in appliances for inhalation in which the substance is absorbed upon an inert solid material.

AMMONIUM THIOCYANATE and substances containing ammonium thiocyanate.

ARSENIC, organic compounds of, in substances containing 3 per cent or less of arsenic (As) when prepared for use as herbicides or defoliants.

BARIUM SILICOFLUORIDE when coated on paper in an amount not exceeding 10 mg per sq cm.

BENDIOCARB in preparations containing 2 per cent or less of bendiocarb.

BENTAZONE and substances containing bentazone.

BENTHIOCARB and substances containing benthiocarb.

BENZENE HEXACHLORIDE in substances containing 10 per cent or less of benzene hexachloride.

BENZOYL PEROXIDE except when included in the Third or Fourth Schedules or when used as an approved food additive.

BIOALLETHRIN except in preparations containing 10 per cent or less of bioallethrin.

BUTHIDAZOLE and substances containing buthidazole.

CADMIUM SULPHIDE in substances containing 2.5 per cent or less of cadmium sulphide, for human therapeutic use.

CAPTAFOL and substances containing captafol.

CARBARYL in preparations containing 10 per cent or less of carbaryl except when included in the Second or Fourth Schedules.

CHLORDECONE in substances containing 5 per cent or less of chlordecone.

CHLORETHALIN and substances containing chlorethalin.

CHLORFENAC and substances containing chlorfenac.

CHLORFENSON and substances containing chlorfenson.

CHLORINATING COMPOUNDS AND BLEACHES containing more than 4 per cent of available chlorine, except—

(a) when included in the Seventh Schedule; or

(b) when included elsewhere in this Schedule.

CHLORNIDINE and substances containing chlornidine.

CHLOROCRESOL and substances containing chlorocresol.

(2-CHLOROETHYL) PHOSPHONIC ACID and substances containing (2-chloroethyl) phosphonic acid.

1-(4-CHLOROPHENOXY)-1-IMIDAZOL-1-YL-3,3-DIMETHYL-2-BUTANONE and substances containing that compound in concentrations of more than 2 per cent, except when such compound or substance is included in the Fourth or Sixth Schedule.

CHLOROPROPYLATE and substances containing chloropropylate.

CHLOROTHALONIL and substances containing chlorothalonil.

COPPER SULPHATE.

4-CPA and substances containing 4-CPA.

CUPRIMYXIN and substances containing cuprimyxin, for the treatment of animals.

CYANATRYN and substances containing cyanatryn.

CYANOACRYLATE ACID ESTERS and substances containing cyanoacrylate acid esters.

CYANURIC ACID (excluding its salts and derivatives) and substances containing cyanuric acid (excluding its salts and derivatives).

CYCLOHEXANONE PEROXIDE and substances containing cyclohexanone peroxide.

3-CYCLOHEXYL-6 (DIMETHYLAMINO)-1-METHYL-1,3,5-TRIAZINE 2,4-(1H,3H)-DIONE and substances containing 3-cyclohexyl-6 (dimethylamino)-1-methyl-1,3,5-triazine 2,4-(1H,3H)-dione.

CYPERMETHRIN in substances containing 10 per cent or less of cypermethrin.

2,4-D and substances containing 2,4-D and amines, esters and salts of 2,4-D.

2,4-DB and substances containing 2,4-DB and salts and esters of 2,4-DB.

DDT in preparations containing 10 per cent or less of DDT, except for human therapeutic use.

2,4-DES and substances containing 2,4-DES and its salts.

N,N - DIALLYLDICHLOROACETAMIDE and substances containing N,N - Diallyldichloroacetamide except when in preparations containing 10 per cent or less of N,N - Diallyldichloroacetamide.

DICAMBA and substances containing dicamba.

DICHLONE and substances containing dichlone.

DICHLOROISOCYANURATES and substances containing dichloroisocyanurates in preparations containing more than 4 per cent available chlorine.

1-[2(2,4-DICHLOROPHENYL)-2-(2-PROPENYLOXY) ETHYL]-1H-IMIDAZOLE and substances containing that compound.

3,6-DICHLOROPICOLINIC ACID and substances containing that compound.

DICHLORVOS when—

(a) impregnated in plastic resin strip material containing 20 per cent or less of dichlorvos;

(b) in sustained release resin pellets for veterinary use containing 20 per cent or less of dichlorvos; or

(c) in aerosol packs containing 10 grams or less of dichlorvos.

DICLORAN and substances containing dicloran.

DICOFOL and substances containing dicofol.

DIMETHIRIMOL and substances containing dimethirimol.

DIMETHYLFORMAMIDE in preparations containing 10 per cent or less of dimethylformamide.

1, 1-DIMETHYLPIPERIDINIUM ION and substances containing 1, 1-Dimethylpiperidinium Ion.

DINITRAMINE and substances containing dinitramine.

DIPHENAMID and substances containing diphenamid.

DODINE and substances containing dodine.

EPTC and substances containing EPTC.

ETHEPHON (excluding its salts and derivatives) and substances containing ethephon (excluding its salts and derivatives).

ETHER PREPARATIONS for use in internal combustion engines.

ETHOFUMESATE and substances containing ethofumesate.

ETHOXYQUIN and substances containing more than 10 per cent of ethoxyquin.

ETHYLENE GLYCOL when packed and labelled as a boiling point and/or freezing point modifier and containing 10 mg/kg of denatonium benzoate.

N-(1-ETHYLPROPYL)-3,4-DIMETHYL-2,6-DINITRO-ANILINE and substances containing N-(1-ethylpropyl) -3,4-dimethyl-2,6-dinitroaniline.

EUCALYPTUS OIL.

FENBUTATIN-OXIDE and substances containing fenbutatin-oxide.

FENOPROP and substances containing fenoprop.

FENSON and substances containing fenson.

FENTHION in substances containing 20 per cent or less of fenthion when packed in single use containers having a capacity of 0.3 ml or less.

FLAMPROP-METHYL and substances containing flampropmethyl.

FORMIC ACID (excluding its salts and derivatives) and substances containing formic acid (excluding its salts and derivatives).

FOSPIRATE when impregnated in plastic resin strip material containing 20 per cent or less of fospirate.

GLYPHOSATE and substances containing glyphosate.

HYDROCARBONS, liquid, including Kerosine, Mineral Turpentine, White Petroleum Spirit, Toluene, Xylene and Light Mineral and Parafin Oils but excluding their derivatives, distilling under 300 degrees Celsius, except—

(a) Toluene and Xylene when included in the Sixth Schedule;

(b) in containers having a capacity of more than 20 litres;

(c) in substances containing 25 per cent or less of a total of such liquid hydrocarbons;

(d) in solid or semi-solid cleaning and polishing preparations;

(e) in preparations packed in pressurised aerosol containers;

(f) in adhesives packed in containers each containing 50 grams or less of adhesive.

HYDROCHLORIC ACID in substances containing 10 per cent or less weight-in-weight of hydrochloric acid, except preparations containing 0.5 per cent or less of hydrochloric acid.

HYDROFLUORIC ACID and HYDROSILICOFLUORIC ACID in preparations containing 0.5 per cent or less of hydrofluoric acid or hydrosilicofluoric acid except in substances containing 15 mg/kg or less of fluoride ion.

HYDROGEN PEROXIDE and substances containing more than 6 per cent weight-in-volume (20 vol) of hydrogen peroxide.

IODOFENPHOS and substances containing idofenphos.

2-ISO-BUTYLAMINO-4-ETHYLAMINO-6-METHOXY-1,3,5-TRIAZINE and substances containing 2-iso-butylamino-4-ethylamino-6-methoxy-1,3,5-triazine.

ISOPROPYL-N-(3N-ETHYL-N-PHENYLCARBAMOYLOXY) PHENYLCARBAMATE and substances containing isopropyl-n-(3-n-ethyl-n-phenylcarbamoyloxy) phenylcarbamate.

KEROSINE and substances containing more than 25 per cent of kerosine when packed in containers of 20 litres or less.

LEVAMISOLE in substances containing 15 per cent or less of levamisole for the treatment of animals.

LINDANE in substances containing 10 per cent or less of lindane except when included in the Second Schedule.

LIQUID EPOXY RESINS and all amines and organic anhydrides used as curing agents for epoxy resins.

MALDISON in substances containing 10 per cent or less of maldison, except for human therapeutic use.

MANCOZEB and substances containing mancozeb.

MANEB and substances containing maneb.

MCPA and substances containing MCPA.

MCPB and substances containing MCPB.

MECOPROP and substances containing mecoprop.

METALDEHYDE in preparations containing 2 per cent or less of metaldehyde.

METHABENZTHIAZURON and substances containing methabenzthiazuron.

METHAZOLE and substances containing methazole.

METHIOCARB in pelletted preparations containing 2 per cent or less of methiocarb.

METHOXYCHLOR and substances containing methoxychlor.

METHYLATED SPIRIT, INDUSTRIAL, as defined by the Spirits Act 1906 of the Parliament of the Commonwealth or any Act in substitution for that Act, as amended from time to time, except in containers having a capacity of more than 5 litres, and except in preparations containing 75 per cent or less methylated spirits, industrial.

METHYLENE CHLORIDE and substances containing methylene chloride except in aerosols.

METHYL ETHYL KETONE and substances containing methyl ethyl ketone when packed in containers of 20 litres or less, except in preparations containing 25 per cent or less of ketones included in the Fifth Schedule.

METHYL ETHYL KETONE PEROXIDE and substances containing methyl ethyl ketone peroxide.

METHYL ISO-AMYL KETONE and substances containing methyl iso-amyl ketone when packed in containers of 20 litres or less, except in preparations containing 25 per cent or less of ketones included in the Fifth Schedule.

METHYL ISO-BUTYL KETONE and substances containing methyl iso-butyl ketone when packed in containers of 20 litres or less, except in preparations containing 25 per cent or less of ketones included in the Fifth Schedule.

METHYL N-(FUR-2-YL)-N-(2,6-XYLYL) ALANINATE and substances containing Methyl N-(fur-2-yl)-N-(2,6-xylyl) alaninate.

METHYL N-(METHOXYACETYL)-N-(2,6-XYLYL) ALANINATE and substances containing methyl N-(methoxyacetyl)-N-(2,6-xylyl) alaninate.

3-(METHYLSULPHONYL) BUTANONE O-METHYLCARBAMOYLOXIME in solid preparations containing 10 per cent or less of that compound.

METIRAM and substances containing metiram.

METOLACHLOR and substances containing metolachlor.

METRIBUZIN and substances containing metribuzin.

MEZINEB and substances containing mezineb.

MINERAL TURPENTINE and substances containing more than 25 per cent of mineral turpentine when packed in containers of 20 litres or less.

N-3-PYRIDYLMETHYL-N'-PARA-NITROPHENYLUREA in substances containing 10 per cent or less of N-3-pyridylmethyl-N'-para-nitrophenylurea.

NALED when impregnated in plastic resin strip material containing 20 per cent or less of naled.

NAPHTHALENE and substances containing naphthalene.

NAPHTHALENE ACETIC ACID, and substances containing Naphthalene Acetic Acid except in preparations containing 25 per cent or less of naphthalene acetic acid.

NITRIC ACID in substances containing 10 per cent or less weight-in-weight of nitric acid, except preparations containing 0.5 per cent or less of nitric acid.

NORBORMIDE and substances containing norbormide.

OIL OF TURPENTINE and substances containing more than 25 per cent of oil of turpentine when packed in containers of 20 litres or less.

ORGANO TIN COMPOUNDS in substances containing 1 per cent or less of such compounds.

ortho-PHENYLPHENOL and substances containing that compound.

OXYCARBOXIN and substances containing oxycarboxin.

PARACHLOROMETACRESOL and substances containing parachlorometacresol.

PARADICHLOROBENZENE and substances containing paradichlorobenzene.

PEBULATE and substance containing pebulate.

PETROL and substances containing more than 25 per cent of petrol when packed in containers of 20 litres or less.

PHOSPHORIC ACID (excluding its salts and derivatives) and substances containing phosphoric acid, except—

(a) when packed in containers with a capacity of not less than 10 litres and labelled with the word "CORROSIVE" in bold *sans serif* capital letters of a height of not less than 1 cm;

(b) in preparations containing 3 per cent or less of phosphoric acid;

(c) in solid and semi-solid preparations;

(d) in professional Dental Kits.

PIRIMICARB in substances containing 0.5 per cent or less of pirimicarb.

POLY (HEXAMETHYLENE BIGUANIDE) HYDROCHLORIDE and substances containing poly (hexamethylene biguanide) hydrochloride.

POTASSIUM HYDROXIDE in substances containing more than 0.5 per cent and not more than 5 per cent of potassium hydroxide.

PROMETRYNE and substances containing prometryne.

PROPANIL and substances containing propanil.

PROPIONIC ACID (excluding its salts and derivatives) in preparations containing 80 per cent or less and more than 30 per cent of propionic acid, except for therapeutic use.

PROPOXUR in dust preparations containing 3 per cent or less of propoxur.

PRYNACHLOR and substances containing prynachlor.

PYRETHRINS and substances containing more than 10 per cent of pyrethrins.

QUATERNARY AMMONIUM COMPOUNDS and preparations containing more than 10 per cent quaternary ammonium compounds, except when included in any other Schedule.

QUINOMETHIONATE and substances containing quinomethionate.

QUINTOZENE and substances containing quintozene.

SALICYLANILIDE and substances containing salicylanilide.

S-BENZYL N,N-DL-(SEC-BUTYL)-THIOLOCARBAMATE and substances containing S-benzyl N,N-dl-(sec-butyl)-thiolocarbamate.

2-SEC BUTYLAMINO-4-ETHYLAMINO-6-METHOXY-1,3,5-TRIAZINE and substances containing 2-sec butylamino-4-ethylamino-6-methoxy-1,3,5-triazine.

SELENIUM SULPHIDE in shampoos for the treatment of animals and in preparations containing 2.5 per cent or less of selenium sulphide for topical therapeutic use.

SODIUM ACID SULPHATE.

SODIUM CHLORATE in substances containing 10 per cent or less of sodium chlorate.

SODIUM HYDROXIDE in substances containing more than 0.5 per cent and not more than 5 per cent of sodium hydroxide.

SODIUM NITRATE and substances containing more than 1 per cent of sodium nitrate, except for therapeutic use.

STYRENE (excluding its derivatives) and substances containing styrene when packed in containers of 20 litres or less.

SULPHAMIC ACID except in preparations containing 10 per cent or less of sulphamic acid.

2,3,6-TBA and substances containing 2,3,6-TBA.

TDE in substances containing 10 per cent or less of TDE.

TERBUTHYLAZINE and substances containing terbuthylazine.

TERBUTRYNE and substances containing terbutryne.

2-TERT BUTYLAMINO-4-ETHYLAMINO-6-METHOXY-1,3,5-TRIAZINE and substances containing 2-tert butylamino-4-ethylamino-6-methoxy-1,3,5-triazine.

TETRACHLORVINPHOS and substances containing tetrachlorvinphos.

TRIADIMENOL and substances containing triadimenol.

TRIALLATE and substances containing triallate.

1,1,1-TRICHLOROETHANE when packed in containers of 20 litres or less but more than 50 ml, except when in substances containing 25 per cent or less of 1,1,1-trichloroetane or except when used in aerosols other than for therapeutic use.

TRICHLOROACETIC ACID, alkali salts of.

TRICHLOROISOCYANURATE when compressed in block or tablet form for use in swimming pools.

TRIETAZINE and substances containing trietazine.

VERNOLATE and substances containing vernolate.

ZINC PYRITHIONE in preparations containing 2 per cent or less of zinc pyrithione.

ZINEB and substances containing zineb.

ZIRAM and substances containing ziram.

Excluding, however, the substances hereinbefore mentioned when contained in any of the following—

Batteries and accumulators.

Ceramics.

Electrical components and electric lamps.

Explosives.

Fireworks other than fireworks containing arsenic.

Glazed pottery.

Inorganic pigments.

Matches.

Paints other than substances prepared for medicinal or cosmetic purposes.

Paper.

Photographic film.

Photographic paper.

Timber and wallboard.

Vitreous enamels.

Sixth Schedule.

ACEPHATE and substances containing acephate.

ACETIC ACID and substances containing more than 80 per cent of acetic acid, except for therapeutic use.

ACETIC ANHYDRIDE and substances containing more than 80 per cent of acetic anhydride.

ACROLEIN in substances containing 50 per cent or less of acrolein.

ALLIDOCHLOR and substances containing allidochlor.

ALLYL ALCOHOL in substances containing 50 per cent or less of allyl alcohol.

ALPHA-CHLORALOSE and substances containing alpha-chloralose when prepared for use as a pesticide.

ALPHA-CHLORHYDRIN and substances containing alpha-chlorhydrin.

AMETRYNE and substances containing ametryne.

AMIDITHION and substances containing amidithion.

2-AMINOBUTANE and substances containing 2-amino-butane.

AMINOCARB in substances containing 50 per cent or less of aminocarb.

2-AMINO-5-DIETHYL AMINO TOLUENE and substances containing 2-amino-5-diethyl amino toluene.

2-AMINO-5-N-ETHYL-N-(B HYDROXY ETHYL) AMINO TOLUENE and substances containing 2-amino-5-N-ethyl-N-(B hydroxy ethyl) amino toluene.

2-AMINO-5-N-ETHYL-N-(B METHANE SULPHONAMIDE ETHYL) AMINO TOLUENE and substances containing 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 and substances containing 2-amino-5-N-ethyl-N-B methoxyethyl amino toluene di-p-toluene.

AMITON in substances containing 25 per cent or less of amiton.

AMITRAZ and substances containing amitraz.

AMMONIA and substances containing more than 5 per cent of free ammonia (NH_3) except in substances for internal use or when used in appliances for inhalation in which the substance is absorbed upon an inert solid material.

ANILINE and substances containing more than 1 per cent of aniline.

ARECOLINE and substances containing arecoline.

ARECOLINE-ACETARSOL in substances for the treatment of hydatid infestation in animals.

ARSENIC and substances containing arsenic when used for agricultural, pastoral or horticultural purposes or for the control of termites.

ARSENIC, COMPOUNDS OF, except—

(a) when included in the Fourth or Fifth Schedules; or

(b) in animal feedstuff containing 75 mg/kg or less of arsenic (As).

ARSENIC, ORGANIC COMPOUNDS OF, when prepared for use as herbicides or defoliants, except when included in the Fifth Schedule.

AZOBENZENE and substances containing azobenzene.

AZOCYCLOTIN and substances containing azocyclotin.

BACITRACIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

BARBAN and substances containing barban.

BARIUM SALTS (except barium sulphate) and substances containing barium salts (except barium sulphate) and except barium silicofluoride when included in the Fifth Schedule.

BENDIOCARB—

(a) in wettable powders containing 80 per cent or less of bendiocarb and when packed in containers or primary packs containing not less than 100 g of bendiocarb;

(b) in insoluble granular preparations containing 5 per cent or less of bendiocarb;

(c) except when included in the Fifth Schedule.

BENQUINOX and substances containing benquinox.

BENSULIDE and substances containing bensulide.

BENZENE HEXACHLORIDE and substances containing more than 10 per cent of benzene hexachloride.

BENZYLPENICILLIN including procaine penicillin in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of benzylpenicillin or procaine penicillin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

BERYLLIUM and its salts except as ores.

BINAPACRYL and substances containing binapacryl.

BITHIONOL for the treatment of animals and substances containing bithionol for the treatment of animals.

3-(3-(4'BROMODIPHENYL-4-YL)-3-HYDROXY-1-PHENYLPROPYL)-4-HYDROXYCOUMARIN in preparations containing 0.1 per cent weight in volume or less of 3-(3-(4'bromodiphenyl-4-yl)-3-hydroxy-1-phenylpropyl)-4-hydroxycoumarin.

3-(3-(4'BROMODIPHENYL-4-YL)-1,2,3,4-TETRAHYDRONAPHTHYL)-4-HYDROXYCOUMARIN in preparations containing 0.25 per cent weight in volume or less of 3-(3-(4'bromodiphenyl-4-yl)-1,2,3,4-tetrahydronaphthyl)-4-hydroxycoumarin.

BROMOFORM, except for therapeutic use.

BROMOPHOS and substances containing bromophos.

BROMOPHOS-ETHYL and substances containing bromophos-ethyl.

BROMOXYNIL and substances containing bromoxynil.

BROTIANIDE and substances containing brotianide.

BUNAMIDINE and substances containing bunamidine.

BUTACARB and substances containing butacarb.

2-BUTOXY-2'-THIOCYANO-DIETHYL ETHER and substances containing 2-butoxy-2'-thiocyano-diethyl ether.

BUTYNORATE and substances containing butynorate.

CADMIUM, compounds of, except when included in the Fifth Schedule.

CAMBENDAZOLE and substances containing cambendazole.

CARBADOX except in animal feedstuffs containing 50 mg/kg or less of the total active principle.

CARBARYL and substances containing carbaryl except when included in the Second, Fourth or Fifth Schedules.

CARBOFURAN in substances containing 25 per cent or less of carbofuran.

CARBON BISULPHIDE and substances containing carbon bisulphide.

CHLORDANE and substances containing chlordane.

CHLORDECONE and substances containing more than 5 per cent of chlordecone.

CHLORFENETHOL and substances containing chlorfenethol.

CHLORMEQUAT and substances containing chlormequat.

CHLOROALLYLDIETHYL THIOCARBAMATE (CDED) and substances containing chloroallyldiethyl thiocarbamate (CDED).

CHLOROMETHIURON and substances containing chloromethiuron.

2-CHLORO-N-[(4-METHOXY-6-METHYL-1,3,5-TRIAZIN-2-YL) AMINOCARBONYL] BENZENE SULPHONAMIDE and substances containing that compound.

S-(6-CHLORO-2-OXO-OXAZOLO(4,5-b)-PYRIDIN-3-YLMETHYL)-o-o-DIMETHYL PHOSPHOROTHIOATE and substances containing S-(6-chloro-2-oxo-oxazolo (4,5-b)-pyridin-3-ylmethyl)-o-o-dimethyl phosphorothioate.

CHLOROPHACINONE and substances containing chlorophacinone.

1-(4-CHLOROPHENOXY)-1-IMIDAZOL-1-YL-3,3-DIMETHYL-2-BUTANONE in substances containing more than 40 per cent of that compound, except when included in the Fourth Schedule.

CHLOROPICRIN in preparations containing 5 per cent or less of chloropicrin.

CHLORPYRIFOS and substances containing chlorpyrifos.

CHLORPYRIFOS-METHYL and substances containing chlorpyrifos-methyl.

CHLORTETRACYCLINE in preparations—

(a) for topical application to animals for ocular use only; or

(b) for intramammary infusion in animals, containing not more than 100 000 international units per dose of chlortetracycline when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

CHLORTHIAMID and substances containing chlorthiamid.

5[2-CHLOR-4(TRIFLUOROMETHYL) PHENOXY]-2-NITROBENZOATE and substances containing 5[2-Chlor-4(Trifluoromethyl) phenoxy]-2-nitrobenzoate.

CHROMATES and DICHROMATES and substances containing any of these.

CHROMIC ACID.

CHROMIUM TRIOXIDE (excluding its salts and derivatives) and substances containing chromium trioxide (excluding its salts and derivatives).

COUMAPHOS in preparations containing 5 per cent or less of coumaphos.

COUMARIN DERIVATIVES and phenylindanedione derivatives not specifically included in these Schedules, except for therapeutic use.

COUMATETRALYL and substances containing coumatetralyl.

CROTOXYPHOS and substances containing crotoxyphos.

CRUFOMATE and substances containing crufomate.

CYANAZINE and substances containing cyanazine.

CYCLOSULFYNE and substances containing cyclosulfyne.

CYHEXATIN and substances containing cyhexatin.

CYPERMETHRIN and substances containing cypermethrin except when included in the Fifth Schedule.

CYTHIOATE and substances containing cythioate.

DAZOMET and substances containing dazomet.

DDT and DDT in preparations containing more than 10 per cent of DDT, except for human therapeutic use.

DEMETON-O-METHYL and DEMETON-S-METHYL in substances containing 50 per cent or less of demeton-O-methyl or demeton-S-methyl or both.

DI-ALLATE and substances containing di-allate.

DIAZINON and substances containing diazinon.

DICHLOFENTHION and substances containing dichlofenthion.

DICHLOFLUANID and substances containing dichlofluanid.

DICHLOROEATHLENE and substances containing dichloroethylene.

DICHLOROETHYL ETHER and substances containing dichloroethyl ether.

1-[2-(2,4-DICHLOROPHENYL)-4-ETHYL-1,3-DIOXOLAN-2-YL-METHYL]-1H-1,2,4-TRIAZOLE and substances containing that compound.

0 - (2,4-DICHLOROPHENYL) - 0 - ETHYL - S - PROPYL-PHOSPHORODITHIOATE and substances containing 0-(2,4-dichlorophenyl) - 0 - ethyl - S - propylphosphorodithioate.

N-(3, 4-DICHLOROPHENYL)-N′-(2-(2″ SULFOXY-4′-CHLORPHENOXY) - 5 CHLORPHENYL) UREA (SODIUM SALT) and substances containing N-(3,4-dichlorophenyl)-N′-(2-(2″ sulfoxy-4′-chlorophenoxy)-5 chlorphenyl) urea (Sodium Salt).

DICHLOROPROPANE and substances containing dichloropropane.

DICHLOROPROPENE and substances containing dichloropropene.

DICHLORVOS in substances containing 50 per cent or less of dichlorvos, except when included in the Fifth Schedule.

DICLOFOP-METHYL and substances containing diclofop-methyl.

DIETHYLENE DIOXIDE and substances containing diethylene dioxide.

N,N-DIETHYL-p-PHENYLENE DIAMINE and substances containing N,N-diethyl-p-phenylene diamine.

DIFENZOQUAT and substances containing difenzoquat.

DIHYDROSTREPTOMYCIN in preparations for intramammary infusion in animals containing not more than 100 000 international units per dose of dihydrostreptomycin when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

DIMETHANONAPHTHALENE and all substitution and/or addition products of dimethanonaphthalene including aldrin and dieldrin and substances containing any of these.

DIMETHOATE and substances containing dimethoate.

1,3-DI(METHOXYCARBONYL)-1-PROPEN-2-YL DIMETHYL PHOSPHATE in substances containing 50 per cent or less of 1,3-di(methoxycarbonyl)-1-propen-2-yl dimethyl phosphate.

DIMETHYLFORMAMIDE and substances containing dimethylformamide except when included in the Fifth Schedule.

2-(2',4'-DIMETHYL-PHENYLIMINO)-3-METHYL-4-THIAZOLINE and substances containing 2-(2',4'-dimethylphenylimino)-3-methyl-4-thiazoline.

DIMETHYL SULPHOXIDE and substances containing dimethyl sulphoxide, except for therapeutic use.

DIMETILAN in substances containing 50 per cent or less of dimetilan.

DIMETRIDAZOLE and substances containing dimetridazole.

DINITROCRESOLS, DINITROPHENOLS and their homologues in substances containing 5 per cent or less of such compounds, except for therapeutic use.

DINOCAP and substances containing dinocap.

DIOXACARB and substances containing dioxacarb.

DIPHACINONE and substances containing diphacinone.

DIQUAT and substances containing diquat.

DISODIUM METHYL ARSONATE in substances prepared for use as a herbicide.

DISULFIRAM and substances containing disulfiram except for therapeutic use.

DISULFOTON in substances containing 25 per cent or less of disulfoton.

DITHIANON and substances containing dithianon.

DITHIAZANINE in substances containing 2 per cent or less of dithiazanine for veterinary use.

DITHIOCARBAMATES and derivatives of dithiocarbamates and substances containing these when prepared for use for agricultural, pastoral or horticultural purposes, except when included in the Fifth Schedule.

3,3'-DI-(TRIFLUOROMETHYL)-4,4'-DICHLORO-N, N'DIPHENYLUREA and substances containing 3,3'-di-(trifluoromethyl)-4,4'-dichloro-N,N'diphenylurea.

DIUREDOSAN and substances containing diuredosan.

ECONAZOLE in preparations containing econazole for external animal use.

ENDOSULFAN in substances containing 50 per cent or less of endosulfan.

ENDOTHAL and substances containing endothal.

EPICHLOROHYDRIN and substances containing more than 1 per cent of epichlorohydrin.

ERYTHROMYCIN in preparations—

(a) for intramammary infusion in animals, containing not more than 100 000 international units per dose of erythromycin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose; or

(b) in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

ETHER SOLVENT and substances containing ether solvent except substances included in the Fifth Schedule.

ETHIOFENCARB and substances containing ethiofencarb.

ETHOATE-METHYL and substances containing ethoate-methyl.

ETHOPROPHOS in substances containing 50 per cent or less of ethoprophos.

5-ETHOXY-3-TRICHLOROMETHYL 1,2,4-THIADAZOLE and substances containing 5-ethoxy-3-trichloromethyl 1,2,4-thiadazole.

ETHYL BROMIDE and substances containing ethyl bromide.

ETHYLENE CHLOROHYDRIN and substances containing ethylene chlorohydrin.

ETHYLENE DIBROMIDE and substances containing ethylene dibromide.

ETHYLENE DICHLORIDE and substances containing ethylene dichloride.

ETHYLENE GLYCOL when packed and labelled as an anti-freeze, except when included in Fifth Schedule.

ETHYLENE OXIDE and substances containing ethylene oxide.

FAMPHUR in substances containing 50 per cent or less of famphur.

FENAMINOSULF in substances containing 50 per cent or less of fenaminosulf.

FENAMIPHOS in granular preparations containing 5 per cent or less of fenamiphos.

FENAZAFLOR and substances containing fenazaflor.

FENCHLORPHOS and substances containing fenchlorphos.

FENITROTHION and substances containing fenitrothion.

FENTHION and substances containing fenthion, except when included in the Fifth Schedule.

FENTHION-ETHYL in substances containing 50 per cent or less of fenthion-ethyl.

FENVALERATE and substances containing fenvalerate.

FERBAM and substances containing ferbam.

FERROCYANIDES and FERRICYANIDES and substances containing more than 1 per cent of ferrocyanides and/or ferricyanides.

FLAVOPHOSPHOLIPOL in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

FORMALDEHYDE and substances containing more than 5 per cent of formaldehyde.

FORMETANATE in substances containing 50 per cent or less of formetanate.

FORMOTHION in substances containing formothion.

FOSPIRATE in substances containing fospirate, except when included in the Fifth Schedule.

FUMAGILLIN and substances containing fumagillin.

GUAZATINE and substances containing guazatine.

HEPTACHLOR and substances containing heptachlor.

HEXACHLOROBENZENE and substances containing hexachlorobenzene.

HEXACHLOROPHANE in substances for the treatment of animals.

HYDRAZINE and substances containing hydrazine.

HYDROCHLORIC ACID and substances containing more than 10 per cent by weight of hydrochloric acid (HCL).

HYDROFLUORIC ACID and HYDROSILICOFLUORIC ACID, their salts and all substances containing any of those compounds or their salts, except—

(a) when used for human therapeutic purposes;

(b) in dentifrices containing 0.5 per cent or less of fluoride ion;

(c) in preparations containing 3 per cent or less of sodium fluoride or sodium silicofluoride when used as preservatives;

(d) when included in the Second, Fourth, Fifth or Seventh Schedules;

(e) in substances containing 15 mg per kg or less of fluoride ion;

(f) ammonium fluosilicate in preparations containing 3.2 per cent or less of ammonium fluosilicate for pesticide purposes.

8-HYDROXYQUINOLINE and its derivatives in substances for topical use on animals.

HYGROMYCIN B in animal feedstuff premixes for use as a anthelmintic containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of Hygromycin B.

IMIDOCARB and salts of imidocarb, and substances containing imidocarb or salts of imidocarb.

IODINE and substances containing iodine—

(a) in Iodophors, except those containing 1.5 per cent or less of available iodine;

(b) in other liquid preparations containing 2.5 per cent or less of available iodine; or

(c) in preparations for animal treatment only, except in solid or semi-solid preparations containing 2.5 per cent or less of available iodine.

IOXYNIL and substances containing ioxynil.

IRON COMPOUNDS in preparations for the treatment of animals.

ISOCARBOPHOS in substances containing 50 per cent or less of isocarbophos.

ISOCYANATES, free organic, except in paints containing 0.1 per cent or less of free organic isocyanates.

KITASAMYCIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 100 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

LASALOCID and substances containing lasalocid.

LAURYLISOQUINOLINIUM BROMIDE and substances containing laurylisoquinolinium bromide.

LEAD COMPOUNDS except—

(a) in preparations for therapeutic or cosmetic use;

(b) in pencil cores, finger colours, showcard colours, pastels, crayons, poster paint/colours or coloured chalks containing 0.01 per cent or less of lead.

LEPTOPHOS in substances containing 50 per cent or less of leptophos.

LINDANE and substances containing lindane except when included in the Second or Fifth Schedule.

MALDISON and substances containing maldison, except when included in the Second or Fifth Schedule.

MEBENDAZOLE and substances containing mebendazole for veterinary use.

MECARBAM in substances containing 50 per cent or less of mecarbam.

MECLOFENAMIC ACID in substances for veterinary use.

MENAZON and substances containing menazon.

MERCURIC CHLORIDE and substances containing mercuric chloride when labelled and packed for photographic use only.

MERCURIC IODIDE and substances containing mercuric iodide, for agricultural, industrial, pastoral or horticultural use.

MERCURIC THIOCYANATE and substances containing mercuric thiocyanate for photographic purposes.

MERCUROUS CHLORIDE and substances containing mercurous chloride, except when included in the Fourth Schedule.

MERCURY, ORGANIC compounds of, and substances containing these, for use in agricultural, industrial, pastoral or horticultural use, except when included in the Seventh Schedule.

METACRESOLSULPHONIC ACID AND FORMALDEHYDE CONDENSATION PRODUCT in preparations for animal use.

METALDEHYDE and substances containing more than 2 per cent of metaldehyde.

METAXANINE and substances containing metaxanine.

METHAM-SODIUM and substances containing metham-sodium.

METHIOCARB and substances containing methiocarb, except when included in the Fifth Schedule.

METHOMYL in substances containing 50 per cent or less of methomyl.

0-2-METHOXYCARBONYLPROP-1-ENYL-0,0-DIMETHYL-PHOSPHOROTHIOATE and substances containing that compound.

METHYL ALCOHOL and substances containing methyl alcohol excluding its derivatives, except methylated spirits.

METHYL BISTHIOCYANATE and substances containing that compound except in substances containing 1 per cent or less of that compound.

METHYL CHLORIDE and substances containing methyl chloride.

METHYL ISOTHIOCYANATE and substances containing methyl isothiocyanate.

1-(B-METHYL SULPHONAMIDE ETHYL)-2-AMINO-3-N,N-DIETHYLAMINO BENZENE and substances containing 1-(B-methyl sulphonamide ethyl)-2-amino-3-N,N-diethylamino benzene.

3-(METHYLSULPHONYL) BUTANONE O-METHYLCARBAMOYLOXIME and substances containing that compound, except when included in the Fifth Schedule.

MOLINATE and substances containing molinate.

MONOCROTOPHOS in substances containing 50 per cent or less of monocrotophos.

NALED and substances containing naled, except when included in the Fifth Schedule.

NAPHTHALOPHOS in substances containing naphthalophos, except when included in the Seventh Schedule.

N-[5-CHLORO-4-[(4-CHLOROPHENYL)-CYANO-METHYL]-2-METHYLPHENYL]-2-HYDROXY-3,5-DIIODOBENZAMIDE and substances containing that compound.

NEOMYCIN in preparations for topical application to animals for ocular use only.

NICOTINE in preparations containing 3 per cent or less of nicotine when labelled and packed for animal use, except tobacco in any form.

NIMIDANE and substances containing nimidane.

NITHIAMIDE and substances containing more than 20 per cent of nithiamide.

N-METHYL CARBAMATES and derivatives thereof and substances containing these for use as pesticides, except when specifically included in any other schedule.

N-3-PYRIDYLMETHYL N'-PARA-NITROPHENYLUREA and substances containing N-3-pyridylmethyl N'-para-nitrophenylurea, except when included in the Fifth Schedule.

NITRIC ACID and substances containing more than 10 per cent by weight of nitric acid.

NITROBENZENE and substances containing more than 0.1 per cent of nitrobenzene, except in soaps containing 1 per cent or less of nitrobenzene or in solid or semi-solid polishes.

NITROFURAN and its derivatives, in preparations packed and labelled for the treatment of ornamental caged birds or ornamental fish only.

NITROPHENOLS, ORTHO, META and PARA and substances containing these.

NITROSCANATE and substances containing nitroscanate.

NITROXYNIL and substances containing nitroxynil.

NOVOBIOCIN in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of novobiocin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

2-n-OCTYL-4-ISOTHIAZOLIN-3-ONE and substances containing 2-n-octyl-4-isothiazolin-3-one.

OLAQUINDOX and substances containing olaquindox, for growth promotion in pigs, except in animal feedstuffs in concentrations of 100 mg/kg or less of the total active principle.

OLEANDOMYCIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

OMETHOATE in substances containing 50 per cent or less of omethoate.

ORGANOPHOSPHORUS COMPOUNDS, including organic fluorophosphates, organic pyrophosphates and organic thiophosphates and substances containing these, except—

(a) when specifically included in any other schedule;

(b) for human therapeutic use.

ORGANO-TIN COMPOUNDS, being di-alkyl, tri-alkyl and tri-phenyl tin compounds where the alkyl group is methyl, ethyl, propyl or butyl not included elsewhere in these Schedules, except—

(a) in plastics;

(b) in paints containing 3 per cent or less of such compounds, calculated as a proportion of the non-volatile content of the paint; or

(c) in other preparations containing 1 per cent or less of such compounds.

ORTHO-DICHLOROBENZENE and substances containing ortho-dichlorobenzene.

ORTHO-TOLIDINE when packed and labelled in concentrations of 0.1 per cent or less of ortho-tolidine for the testing of water.

OXALIC ACID, water soluble oxalates and substances containing these, except laundry blue.

OXANTEL EMBONATE and substances containing oxantel embonate prepared for the treatment of animals.

OXFENDAZOLE and substances containing oxfendazole.

OXYCLOZANIDE and substances containing oxyclozanide.

OXYTETRACYCLINE in preparations—

(a) for topical application to animals for ocular use only;

(b) for intramammary infusion in animals, containing not more than 100 000 international units per dose of oxytetracycline, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

OXYTHIOQUINOX and substances containing oxythioquinox.

PARAQUAT in granular preparations containing 3 per cent or less of paraquat and in surface sprays prepacked in pressurised containers containing not more than 350 grams of pressurised spray containing paraquat at a concentration of not more than 6.3 g/kg.

PARBENDAZOLE and substances containing parbendazole.

PENTACHLOROPHENOL and substances containing pentachlorophenol, except in substances containing 0.5 per cent or less of pentachlorophenol.

PERFLUIDONE and substances containing perfluidone.

PERMANGANATES and substances containing permanganates.

PHENETHICILLIN in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of phenethicillin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

PHENKAPTON and substances containing phenkapton.

PHENOL and any homologue of phenol boiling below 220° C, creosote, and substances containing more than 3 per cent by weight of such substances or homologues, except for therapeutic use.

PHENOXYMETHYLPENICILLIN in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of phenoxymethylpenicillin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

PHENYLENE DIAMINES and alkylated phenylene diamines, not elsewhere specified in this schedule:

(a) when used in hair dyes;

(b) in preparations packed and labelled for photographic purposes;

(c) in preparations packed and labelled for testing water except diethyl-or dimethyl-para-phenylenediamine in tablets containing 10 mg or less in opaque strip packaging labelled for water testing.

PHOSALONE and substances containing phosalone.

PHOSMET and substances containing phosmet.

PHOSPHIDES METALLIC and substances containing metallic phosphides.

PHOSPHORUS YELLOW in substances containing 0.5 per cent or less of free phosphorus.

PHOXIM and substances containing phoxim.

PICRIC ACID and substances containing more than 5 per cent of picric acid.

PINDONE and substances containing pindone.

PIPEROPHOS and substances containing piperophos.

PIRIMICARB and substances containing pirimicarb, except when included in the Fifth Schedule.

PIRIMIPHOS-ETHYL and substances containing pirimiphos-ethyl.

PIRIMIPHOS-METHYL and substances containing pirimiphos-methyl.

POTASSIUM BROMATE and substances containing more than 0.5 per cent of potassium bromate.

POTASSIUM CYANATE and substances containing potassium cyanate.

POTASSIUM HYDROXIDE and substances containing more than 5 per cent of potassium hydroxide.

PROFENOPHOS and substances containing profenophos.

PROGESTERONE in a silicon rubber elastomer when used as a controlled-release implant for synchronization of oestrus in cattle.

PROMACYL and substances containing promacyl.

PROMECARB in preparations containing 50 per cent or less of promecarb.

PROPACHLOR and substances containing propachlor.

PROPIONIC ACID (excluding its salts and derivatives), except—

(a) in preparations containing 80 per cent or less of propionic acid; and

(b) for therapeutic use.

PROPOXUR except when included in the Second or Fifth Schedules.

PYRAZOPHOS and substances containing pyrazophos.

N-3-PYRIDYLMETHYL N^1-p-NITROPHENYLUREA and substances containing N-3-pyridylmethyl N^1-p-nitrophenylurea, except when included in the Fifth Schedule.

RAFOXANIDE and substances containing rafoxanide.

SELENIUM, compounds of—

(a) in preparations containing a 2.5 per cent or less of selenium—

(i) when packed and labelled for the blueing of gun barrels;

(ii) when packed and labelled for photographic purposes;

(b) in preparations containing 0.1 per cent or less of selenium when packed and labelled as vaccines, drenches or pastes for treatment of animals;

(c) in preparations containing 0.5 per cent or less of selenium when packed and labelled as other injections for treatment of animals;

(d) in premixes containing 2 per cent or less of selenium when packed and labelled for incorporation into animal feeds to provide 0.1 g/tonne or less of selenium.

SODIUM BROMATE and substances containing more than 0.5 per cent of sodium bromate.

SODIUM CHLORATE and substances containing more than 10 per cent of sodium chlorate.

SODIUM HYDROXIDE and substances containing more than 5 per cent of sodium hydroxide.

SPIRAMYCIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

STREPTOMYCIN in preparations for intramammary infusion in animals, containing not more than 100 000 international units per dose of streptomycin, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose.

STRYCHNINE in grain baits containing 0.5 per cent or less of strychnine and registered as a pesticide.

SULFALLATE and substances containing sulfallate.

SULPHANILAMIDE and its derivatives, in preparations packed and labelled for the treatment of ornamental caged birds or ornamental fish only.

SULPHAQUINOXALINE when packed and labelled for use as a coccidiostat in poultry, except preparations containing 200 mg/kg or less of sulphaquinoxaline.

SULPHURIC ACID and substances containing sulphuric acid, except—

(a) in fire extinguishers; and

(b) substances containing 0.5 per cent or less by weight of sulphuric acid (H_2SO_4).

SULPROPHOS and substances containing sulprophos.

2,4,5-T and substances containing 2,4,5-T or salts, amines, esters or ethers of 2,4,5,-T.

TCA and substances containing TCA.

TCMTB (2-(THIOCYANOMETHYL THIO) BENZOTHIAZOLE) and substances containing TCMTB (2-(thiocyanomethyl thio) benzothiazole).

TDE and substances containing TDE, except when included in the Fifth Schedule.

TEMEPHOS and substances containing temephos.

TERPENES, CHLORINATED, and substances containing chlorinated terpenes.

TESTOSTERONE PROPIONATE, DI-PROPIONATE AND ENANTHATE in preparations labelled solely for treatment and prevention of pizzle and sheath rot in wethers.

TETRACHLOROETHYLENE, except when packed in containers of 50 ml or less and except when prepared for the treatment of humans or for veterinary purposes.

TETRACYCLINE and salts of tetracycline in preparations—

(a) for topical application to animals for ocular use only;

(b) for intramammary infusion in animals, containing not more than 100 000 international units per dose of tetracycline, when suitably coloured with Brilliant Blue FCF or other approved colour as a marker and when packed in applicator devices specially designed for the purpose; or

(c) when packed and labelled for the treatment of ornamental caged birds or ornamental fish only.

TETRADIFON and substances containing tetradifon.

TETRAMISOLE and substances containing tetramisole, including levamisole, for veterinary use, except when included in the Fifth Schedule.

THIAZAFLURON and substances containing thiazafluron.

THIOMETON and substances containing thiometon.

THIOUREA and substances containing thiourea, except for therapeutic use.

THIRAM and substances containing thiram.

TOLUENE excluding its derivatives when packed in containers of 20 litres or less, except—

(a) in preparations containing 50 per cent or less of Toluene or both Toluene and Xylene; or

(b) when packed in containers of 50 ml or less.

TRIADIMEFON and substances containing triadimefon.

S,S,S-TRIBUTYLPHOSPHOROTHIOLATE and substances containing S,S,S-tributylphosphorothiolate.

TRICHLORFON and substances containing trichlorfon.

TRICHLOROETHYLENE and substances containing trichloroethylene, except for therapeutic use and except when packed in containers of 50 ml or less.

3,5,6-TRICHLOROPYRID-2-YLOXYACETIC ACID and substances containing 3,5,6-Trichloropyrid-2-yloxyacetic acid.

TRICHLOROPHENOL and substances containing trichlorophenol.

TRIDEMORPH and substances containing tridemorph.

TRIETHYL PHOSPHATE and substances containing triethyl phosphate.

TYLOSIN and its salts in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

VAMIDOTHION in substances containing 50 per cent or less of vamidothion.

VIRGINIAMYCIN in animal feedstuff premixes for growth promotion purposes containing concentrations greater than 50 mg/kg but not more than 20 000 mg/kg of the total antibiotic principle.

WARFARIN and substances containing warfarin, except for therapeutic use.

XYLENE excluding its derivatives, when packed in containers of 20 litres or less except—

(a) in preparations containing 50 per cent or less of xylene or of both Xylene and Toluene; or

(b) when packed in containers of 50 ml or less.

ZINC CHLORIDE and substances containing more than 5 per cent of zinc chloride.

ZINC PHENOLSULPHONATE and substances containing more than 5 per cent of zinc phenolsulphonate.

ZINC SULPHATE, except for human therapeutic use and except in preparations containing 5 per cent or less of zinc sulphate.

Excluding, however, the substances hereinbefore mentioned when contained in any of the following—

Batteries and accumulators.

Blankets mothproofed with dieldrin in the mill during finishing as directed by C.S.I.R.O.

Ceramics.

Electrical components and electric lamps.

Explosives.

Fireworks other than fireworks containing arsenic.

Glazed Pottery.

Lubricants, unless specified in the Schedule.

Matches.

Motor fuels, other than those containing methyl alcohol.

Paints other than prepared for medicinal or cosmetic purposes.

Paper.

Photographic film.

Photographic paper.

Selenium contained in animal feeds containing 100 mg per tonne or less of selenium in total feed.

Timber and wallboard.

Vitreous enamels.

Seventh Schedule.

Special Poisons.

Substances or preparations of exceptional danger which require special precautions and restrictions in manufacture, use and sale.

ACROLEIN and substances containing more than 50 per cent of acrolein.

ALDICARB and substances containing aldicarb.

ALLYL ALCOHOL and substances containing more than 50 per cent of allyl alcohol.

AMINOCARB and substances containing more than 50 per cent of aminocarb.

4-AMINO-PYRIDINE and substances containing 4-amino-pyridine.

ANTU and substances containing antu.

ARPRINOCID and substances containing arprinocid.

BENDIOCARB and substances containing bendiocarb, except when included in the Fifth or Sixth Schedule.

BENZENE and substances containing benzene except—

(a) preparations containing 1.5 per cent v/v or less of benzene.

(b) petrol and fuels for internal combustion engines containing 5 per cent or less of benzene.

(c) motor fuels containing more than 5 per cent but not more than 20 per cent of benzene when packed in containers of 20 litres or less.

BETA HYDROXYETHYL HYDRAZINE and substances containing beta hydroxyethyl hydrazine.

3-(3-(4'BROMODIPHENYL-4-YL)-3-HYDROXY-1-PHENYLPROPYL)-4-HYDROXYCOUMARIN and substances containing 3-(3-(4'bromodiphenyl-4-yl)-3-hydroxy-1-phenylpropyl)-4-hydroxycoumarin except when included in the Sixth Schedule.

3(3-(4'BROMODIPHENYL-4-YL)-1,2,3,4-TETRAHYDRONAPHTHYL)-4-HYDROXYCOUMARIN and substances containing 3(3-(4'bromodiphenyl-4-yl)-1,2,3,4-tetrahydronaphthyl)-4-hydroxycoumarin except when included in the Sixth Schedule.

CAMPHECLOR and substances containing campheclor.

CARBOFURAN and substances containing more than 25 per cent of carbofuran.

CARBON TETRACHLORIDE and substances containing carbon tetrachloride.

CARCINOGENIC SUBSTANCES—

2-Acetyl Aminofluorene.
Alphanapthylamine.
4-Aminobiphenyl.
Benzidine.
Betanaphthylamine.
Beta Propriolactone.
Bis-Chloromethyl Ether.
3,3'-Dichlorobenzidine.
4-Dimethylamino Azobenzene.
Methyl Chloromethyl Ether.
4,4-Methylene Bis-(2-Chloroaniline).
4-Nitrobiphenyl.
N-Nitrosodimethylamine

and substances containing any of those compounds.

CHLORDIMEFORM and substances containing chlordimeform.

CHLORINE as such.

5-CHLORO-3-METHYL-4-NITRO PYRAZOLE and substances containing 5-chloro-3-methyl-4-nitro pyrazole.

CHLOROPICRIN and substances containing more than 5 per cent of chloropicrin.

CLOMIPHENE and other products specifically prepared to stimulate ovulation and substances containing those products.

COUMAPHOS and substances containing coumaphos except when included in the Sixth Schedule.

CYCLOFENIL and substances containing cyclofenil.

DICROTOPHOS and substances containing dicrotophos.

0,0-DIETHYL-0-(2,5-DICHLORO-4-(METHYLTHIO) PHENYL) THIONOPHOSPHATE and substances containing 0,0-diethyl-0-(2,5-dichloro-4-(methylthio) phenyl) thionophosphate.

DIMETILAN and substances containing more than 50 per cent of dimetilan.

DINITROCRESOLS, DINITROPHENOLS and their homologues and substances containing more than 5 per cent of such compounds either separately or together, except for therapeutic use.

DULCIN and substances containing dulcin.

ENDOSULFAN and substances containing more than 50 per cent of endosulfan.

ETHOXYETHYL MERCURY CHLORIDE and substances containing ethoxyethyl mercury chloride.

ETHYL MERCURY CHLORIDE and substances containing ethyl mercury chloride.

FENAMINOSULF and substances containing more than 50 per cent of fenaminosulf.

FENAMIPHOS and substances containing fenamiphos, except when included in the Sixth Schedule.

FENSULFOTHION and substances containing fensulfothion.

FLUOROACETAMIDE and substances containing fluoroacetamide.

FLUORACETIC ACID, its salts and substances containing fluoracetic acid or its salts.

FORMETANATE and substances containing more than 50 per cent of formetanate.

HALOFUGINONE and substances containing halofuginone except in prepared stockfeed containing 3 g/tonne or less of halofuginone.

HYDROCYANIC ACID and CYANIDES, and substances containing more than the equivalent of 0.15 per cent of hydrocyanic acid, except for therapeutic use.

ISOCARBOPHOS and substances containing more than 50 per cent of isocarbophos.

LEPTOPHOS and substances containing more than 50 per cent of leptophos.

LIVE VIRUS VACCINE of the strain known as A2/ENGLAND 42/72.

MERCURIC CHLORIDE when prepared for use for agricultural, industrial, pastoral or horticultural purposes.

METHAPYRILENE and substances containing methapyrilene.

METHOMYL and substances containing more than 50 per cent of methomyl.

METHYL BROMIDE and substances containing methyl bromide.

MIREX and substances containing mirex.

NAPHTHALOPHOS and substances containing more than 50 per cent of naphthalophos, except when specially prepared and packed as a sheep drench.

NICOTINE and its salts and substances containing nicotine and its salts except when included in the Fourth or Sixth Schedule or tobacco in any form.

ORGANO-PHOSPHORUS COMPOUNDS.

Substances containing more than 25 per cent of—

Amiton oxalate.
Azinphos-methyl.
Carbophenothion.
Demeton.
Dimefox.
Disulfoton.
Mazidox.
Mevinphos.
Parathion.
Phorate.
Phosfolan.
Schradan.
Sulfotep.
TEPP.

Substances containing more than 50 per cent of—

Azinphos-ethyl.
Chlorfenvinphos.
Coumithioate.
Demeton-methyl.
Demeton-S-methyl.
Dichlorvos.
Diethyl methylcoumarinyl phosphorothioate.

1, 3-di (Methoxycarbonyl)-1-propen-2-yl-dimethyl phosphate.

Dioxathion.

Endothion.

EPN.

Ethion.

Ethoprophos.

Famphur.

Fenthion-ethyl.

Mecarbam.

Methamidophos.

Methidathion.

Methyl-carbophenothion.

Mipafox.

Monocrotophos.

Omethoate.

Oxydemeton-methyl.

Parathion-methyl.

Phosphamidon.

Prothoate.

Thionazin.

Triamiphos.

Vamidothion.

ORTHO-TOLIDINE, except when included in the Sixth Schedule and except in solid state therapeutic diagnostic reagents.

OXAMYL, and substances containing oxamyl.

PARAQUAT, and substances containing paraquat except when included in the Sixth Schedule.

POLYCHLORINATED BIPHENYLS and substances containing polychlorinated biphenyls.

PROMECARB and substances containing more than 50 per cent of promecarb.

PROSTAGLANDINS and substances containing prostaglandins except when included in the Fourth Schedule.

S-ALPHA-CYANO-m-PHENOXY-BENZYL(1R,3R)-3-(2,2-DIBROMOVINYL)-2-DIMETHYLCYCLOPROPANE and substances containing S-alpha-cyano-m-phenoxy-benzyl (1R,3R)-3-(2,2-dibromovinyl)-2-dimethylcyclopropane.

S-(2-CHLORO-1-PHTHALIMIDOETHYL)-0,0-DIETHYL-PHOSPHORODITHIOATE and substances containing S-(2-chloro-1-phthalimidoethyl) -0,0-diethylphosphorodithioate.

STRYCHNINE and its salts and substances containing these, except when included in the First, Fourth or Sixth Schedules.

TERBUFOS and substances containing terbufos.

TETRACHLOROETHANE and substances containing tetrachloroethane.

THALLIUM and its salts and substances containing thallium or its salts.

THIOFANOX and substances containing thiofanox.

TRIAZBUTIL and substances containing triazbutil.

TRICHLOROISOCYANURATE, except when included in the Fifth Schedule and except in preparations containing 4 per cent or less of available chlorine.

VINYL CHLORIDE.

Eighth Schedule.

Includes any active principle, alkaloid, derivative, natural or synthetic, salt, compound and all preparations and admixtures containing any proportion thereof and these are therefore subject to all the restrictions of this Schedule unless specifically exempted.

A substance specified in this Schedule includes every ester and ether of the substance and every salt of such ester or ether and except in the case of Levomethorphan and Levorphanol every stereoisomer of the substance and every salt of such stereoisomer.

ACETORPHINE (0^3-acetyl-7, 8 dihydro-7a (1 (R)-hydroxy-1-methylbutyl)-0^6-methyl-6, 14-endoetheno-morphine).

ACETYLDIHYDROCODEINE, except when included in the Second or Fourth Schedule.

ACETYLMETHADOL (3-acetoxy-6-dimethylamino-4, 4-diphenylheptane).

ALLYLPRODINE (3-allyl-1-methyl-4-phenyl-4-propionoxypiperidine).

ALPHACETYLMETHADOL (alpha-3-acetoxy-6-dimethylamino-4, 4-diphenyl-heptane).

ALPHAMEPRODINE (alpha-3-ethyl-1-methyl-4-phenyl-4-propionoxypiperidine).

ALPHAMETHADOL (alpha-6-dimethylamino-4, 4-diphenyl-3-heptanol).

ALPHAPRODINE (alpha-1, 3-dimethyl-4-phenyl-4-propionoxypiperidine).

AMPHETAMINE.

ANILERIDINE (1-para-aminophenethyl-4-phenylpiperidine-4-carboxylic acid ethyl ester).

BENZETHIDINE (1-(2-benzyloxyethyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

BENZYLMORPHINE (3-benzylmorphine).

BETACETYLMETHADOL (beta-3-acetoxy-6-dimethylamino-4, 4 diphenyl-heptane).

BETAMEPRODINE (beta-3-ethyl-1-methyl-4-phenyl-4-propionoxypiperidine).

BETAMETHADOL (beta-6-dimethylamino-4, 4-diphenyl-3-heptanol).

BETAPRODINE (beta-1, 3-dimethyl-4-phenyl-4-propionoxypiperidine).

BEZITRAMIDE (1-(3-syano-3, 3-diphenylpropyl)-4-(2-oxo-3-propionyl-1-benzimidazolinyl)-piperidine.

BUFOTENINE.

CANNABIS AND CANNABIS RESIN AND EXTRACTS AND TINCTURES OF CANNABIS.

CLONITAZENE (2-para-chlorobenzyl-1-diethylaminoethyl-5-nitro-benzimidazole).

COCAINE (methyl ester of benzoylecgonine).

COCA LEAF.

CODEINE (3-methyl morphine), except when included in the Second, Third or Fourth Schedules.

CODEINE-N-OXIDE.

CODOXIME (dihydrocodeinone-6-carboxymethyloxime).

CONCENTRATE OF POPPY STRAW (the material arising when poppy straw has entered into a process of concentration of its alkaloids).

DESOMORPHINE (dihydrodesoxymorphine).

DEXAMPHETAMINE.

DEXTROMORAMIDE ((+)-4-(2-methyl-4-oxo-3, 3-diphenyl-4-(1-pyrrolidinyl) butyl) morpholine).

DIACETYLMORPHINE (heroin).

DIAMPROMIDE (N-(2-(methylphenethylamino) propyl) propionanilide).

DIETHYLTHIAMBUTENE (3-diethylamino-1, 1-di-(2'-thienyl)-1-butene).

DIFENOXIN (1-(3-cyano-3, 3 diphenylpropyl)-4-phenylisonipecotic acid) excluding preparations containing, per dosage unit, not more than 0.5 mg of difenoxin and a quantity of atropine sulphate equivalent to at least 5 per cent of the dose of difenoxin.

DIHYDROCODEINE, except when included in the Third or Fourth Schedule.

DIHYDROMORPHINE.

DIMENOXADOL (2-dimethylaminoethyl-1-ethoxy-1, 1-diphenylacetate).

DIMEPHEPTANOL (6-dimethylamino-4, 4-diphenyl-3-heptanol).

2, 5-DIMETHOXY-4-BROMOAMPHETAMINE.

2, 5-DIMETHOXY-4-METHYLAMPHETAMINE.

DIMETHYLTHIAMBUTENE (3-dimethylamino-1, 1-di(2' thienyl)-1-butene).

DIMETHYLTRYPTAMINE.

DIOXAPHETYL BUTYRATE (ethyl 4-morpholino-2, 2-diphenylbutyrate).

DIPHENOXYLATE (1-(3-cyano-3, 3-diphenylpropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester) excluding preparations containing, per dosage unit, not more than 2.5 mg of diphenoxylate calculated as base, and a quantity of atropine sulphate equivalent to at least 1 per cent of the dose of diphenoxylate.

DIPIPANONE (4, 4-diphenyl-6-piperidine-3-heptanone).

DROTEBANOL (3, 4-dimethoxy-17-methylmorphinan-6B, 14-diol).

ECGONINE, ITS ESTERS AND DERIVATIVES WHICH ARE CONVERTIBLE TO ECGONINE AND COCAINE.

ETHYLMETHYLTHIAMBUTENE (3-ethylmethlamino-1, 1-di-(2'-thienyl)-1-butene).

ETHYLMORPHINE (3-ethylmorphine) and substances containing more than 2.5 per cent of ethylmorphine.

ETONITAZENE (1-diethylaminoethyl-2-para-ethoxybenzyl-5-nitro-benzimidazole).

ETORPHINE (7, 8-dihydro-7a (1 (R)-hydroxy-1-methylbutyl)-O^6,methyl-6, 14-endoethenomorphine).

ETOXERIDINE (1-(2-(2-hydroxyethoxy)ethyl)-4-phenyl piperidine-4-carboxylic acid ethyl ester).

FENTANYL (1-phenethyl 4-N-propionyl-anilino piperidine).

FURETHIDINE (1-(2-tetrahydrofurfuryloxyethyl)-4- phenylipiperidine-4-carboxylic acid ethyl ester).

HALLUCINOGENIC SUBSTANCES structurally derived from methoxyphenethylamine.

HEPTANE DERIVATIVES having addiction properties, not specifically included elsewhere in this Schedule.

HEROIN.

HYDROCODONE (dihydrocodinone).

HYDROMORPHINOL (14-hydroxydihydromorphine).

HYDROMORPHONE (dihydromorphinone).

HYDROXYPETHIDINE (4-meta-hydroxyphenyl-1-methylpiperidine-4-carboxylic acid ethyl ester).

ISOMETHADONE (6-dimethylamino-5-methyl-4, 4-diphenyl -3-hexanone).

KETOBEMIDONE (4-meta-hydroxyphenyl-1-methyl-4-proprionylpiperidine).

LEVOMETHORPHAN ((—)-3-methoxy-N-methylmorphinan).

LEVOMORAMIDE ((—)4-(2-methyl-4-oxo-3, 3-diphenyl-4-(1-pyrrolidinyl) butyl) morpholine).

LEVOPHENACYLMORPHAN ((—)-3-hydroxy-N-phenacylmorphinan).

LEVORPHANOL ((—)-3-hydroxy-N-methylmorphinan).

LYSERGIC ACID DIETHYLAMIDE (LSD).

MECLOQUALONE.

MESCALINE.

METAZOCINE (2'-hydroxy-2, 5, 9-trimethyl-6, 7-benzomorphan).

METHADONE (6-dimethylamino-4, 4-diphenyl-3-heptanone).

METHADONE-INTERMEDIATE (4-cyano-2-dimethylamino-4, 4-diphenyl-butane).

METHAQUALONE.

METHYLAMPHETAMINE.

METHYLDESORPHINE (6-methyl-delta-6-desoxymorphine).

METHYLDIHYDROMORPHINE (6-methyldihydromorphine).

METHYLPHENIDATE.

1-METHYL-4-PHENYLPIPERIDINE-4-CARBOXYLIC ACID ESTERS.

METOPON (5-methyldihydromorphinone).

MORAMIDE-INTERMEDIATE (2-methyl-3-morpholino-1, 1-diphenylpropane carboxylic acid).

MORPHERIDINE (1-(2-morpholinoethyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

MORPHINE.

MORPHINE DERIVATIVES not specifically included elsewhere in this or any other Schedule.

MORPHINE METHOBROMIDE AND OTHER PENTAVALENT NITROGEN MORPHINE DERIVATIVES.

MORPHINE-N-OXIDE.

MORPHINE SUBSTITUTES not specifically included elsewhere in this Schedule.

MYROPHINE (myristylbenzylmorphine).

NICOCODINE, except when included in the Second or Fourth Schedules.

NICODICODINE, except when included in the Second or Fourth Schedules.

NICOMORPHINE (3, 6-dinicotinylmorphine).

NORACYMETHADOL ((±)-alpha-3-acetoxy-6-methylamino-4, 4-diphenyl-heptane).

NORCODEINE, except when included in the Second or Fourth Schedule.

NORLEVORPHANOL ((—) -3-hydroxymorphinan).

NORMETHADONE (6-dimethylamino-4, 4-diphenyl-3-hexanone).

NORMORPHINE (n-demethylated morphine).

NORPIPANONE (4,4-diphenyl-6-piperidine-3-hexanone).

OPIUM in any form, except the alkaloids papaverine and noscapine.

OXYCODONE (14-hydroxydihydrocodeinone).

OXYMORPHONE (14-hydroxydihydromorphinone), except Naloxone.

PCE (N-ethyl-1-phenylcyclohexylamine).

PENTAZOCINE.

PETHIDINE (1-methyl-4-phenylpiperidine-4-carboxylic acid ethyl ester).

PETHIDINE-INTERMEDIATE A (4-cyano-1-methyl-4-phenylpiperidine).

PETHIDINE INTERMEDIATE B (4-phenylpiperidine-4-carboxylic acid ethyl ester).

PETHIDINE INTERMEDIATE C (1-methyl-4-phenylpiperidine-4-carboxylic acid).

PHENADOXONE (6-morpholino-4, 4-diphenyl-3-heptanone).

PHENAMPROMIDE (N-(1-methyl-2-piperidinoethyl) propionanilide).

PHENAZOCINE (2-hydroxy-5, 9-dimethyl-2-phenethyl-6, 7-benzomorphan).

PHENCYCLIDINE.

PHENMETRAZINE.

PHENOMORPHAN (3-hydroxy-N-phenethylmorphinan).

PHENOPERIDINE (1-(3-hydroxy-3-phenylpropyl)-4-phenylpiperidine-4-carboxylic acid ethyl ester).

PHOLCODINE except when included in the Second or Fourth Schedules.

PHP or PCPY (1-(-phenylcyclohexl) pyrrolidine).

PIMINODINE (4-phenyl-1-(3-phenylaminopropyl) piperidine-4-carboxylic acid ethyl ester).

PIPERIDINE DERIVATIVES having addiction properties, not specifically included elsewhere in this Schedule.

PIRITRAMIDE (1-(3-cyano-3, 3-diphenylpropyl)-4-(1-piperidino) piperidine-4-carboxylic acid amide).

PROHEPTAZINE (1, 3-dimethyl-4-propionoxyazacycloheptane).

PROPERIDINE (1-methyl-4-phenylpiperidine-4-carboxylic acid isopropyl ester).

PROPIRAM.

PSILOCIN.

PSILOCYBIN.

RACEMETHORPHAN ((±)-methoxy-N-methylmorphinan).

RACEMORAMIDE ((±)-4-(2-methyl-4-oxo-3, 3-diphenyl-4-(1-pyrrolidinyl) butyl) morpholine).

RACEMORPHAN ((±)-3-hydroxy-N-methylmorphinan).

SUFENTANIL.

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

TETRAHYDROCANNABINOLS.

THEBACON (acetyl dihydrocodeinone).

THEBAINE.

TILIDENE.

TRIMPERIDINE (1, 2, 5-trimethyl-4-phenyl-4-propionoxy-piperidine).

APPENDIX "B"

CONVENTIONS.

S. 45. The International Opium Convention signed at the Hague on the 23rd day of January, 1912.

The Convention that is referred to as the Geneva Convention in the preamble to the Dangerous Drugs Act 1925, of the Parliament of the United Kingdom, and as having been signed on behalf of His Majesty on the 19th day of February, 1925.

The Single Convention on Narcotic Drugs, 1961, signed at New York on the 30th day of March, 1961.

APPENDIX "C"

S. 55. FORM OF WARRANT.

To wit } To

WHEREAS it appears to me.....a Justice of the Peace, by the complaint on oath of (A.B.) of (address) in the State (occupation), pursuant to the provisions of section 55 of the Poisons Act 1964, that there is reasonable ground for suspecting that in the house or premises situated at (situation) in the State (here state the subject matter of the suspicion).

This is therefore to authorize and require you with such assistants as may be necessary to enter into and upon and search such house or premises at any time during the day or night and there to open or break open if necessary and search all things found therein or thereon and to search all persons found therein or thereon and if necessary to use force in making such entry into or upon such house or premises, whether by breaking open doors or otherwise, and to arrest and bring before a stipendiary magistrate or two Justices of the Peace all persons found therein or thereon and seize all substances and preparations found in or on such house or premises, or in the possession or under the control

of any person therein as may reasonably be suspected of being or containing a poison or are in contravention of any provision of the Poisons Act 1964, or the regulations made thereunder, and all articles used or capable of being used for the purpose of preparing, taking or administering any drug of addiction or specified drug for the purposes of addiction, and all documents relating to any transaction or dealing that would if carried out be an offence against the said Act or regulations, or any corresponding law in force outside the State, to be dealt with according to law:

And for so doing this shall be your Warrant.

Given under my hand at .....

in Western Australia this .....

day of ....., 19.....

By Authority: WILLIAM C. BROWN, Government Printer