CHAPTER 105 PHARMACY AND POISONSCHAPTER 105

PHARMACY AND POISONS

ARRANGEMENT OF SECTIONS

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CHAPTER 105 PHARMACY AND POISONSCHAPTER 105

AN ACT TO CONTROL THE PRACTICE OF PHARMACY AND THE SALE AND DISTRIBUTION OF POISONS

[28th July 1941]

PART I

1. This Act may be cited as the Pharmacy and Poisons Act.

2. In this Act, unless the context otherwise requires—

"Board" means the Pharmacy and Poisons Board appointed under this Act;

5 of 1941 6 of 1953 7 of 1958 6 of 1967
LN 78 of 1973 LN 46A of 1978 LN 88 of 1978 LN 60 of 1981 LN 5 of 1988

Short title

Interpretation

7 of 1958, s. 4
6 of 1967, sched LN 46A of 1978

"Chairman" means the Chairman of the Board appointed under this Act;

"Court" means the High Court;

"member" means a member of the Board constituted under this Act;

"poison" includes the several substances mentioned in the poisons list in Schedule B;

"qualified medical practitioner", "qualified dentist" and "qualified veterinary surgeon" means a medical practitioner, a dentist and a veterinary surgeon respectively holding a diploma or certificate entitling him to practise his profession in the United Kingdom or in any other country approved by the Minister;

"register" means the register of pharmacists registered under this Act;

"registered pharmacist" means a person registered under this Act;

"Under Secretary (Health), Ministry of Health and Medical Services" means the officer for the time being holding the office of Under Secretary (Health), Ministry of Health and Medical Services.

PART II ADMINISTRATIONPART II

ADMINISTRATION

3.—(1) For the purposes of this Act there is hereby constituted an authority to be called the "Pharmacy and Poisons Board".

(2) The Board shall be a body corporate with perpetual succession and a common seal and shall be capable of suing and being sued.

(3) All courts, judges and persons acting judicially shall take judicial notice of the seal of the Board affixed to any document and shall deem that it was duly affixed.

4.—(1) The Board shall consist of the Under Secretary (Health), Ministry of Health and Medical Services and of two members who shall be appointed from time to time by the Minister.

(2) The Under Secretary (Health), Ministry of Health and Medical Services shall be ex officio Chairman of the Board.

(3) The Chairman and one member shall form a quorum.

(4) The Chairman shall have an original vote and, in the event of equality of voting, a second or casting vote.

5. All meetings of the Board shall be convened by the Chairman by notice in writing to the other members of the Board, specifying the time and place of meeting.

6.—(1) For the purposes of this Act the Board may, by writing under the hand of the Chairman, summon any person to attend the meeting of the

Meetings of the Board

Board may summon person to attend and give evidence

(2) The Board may in its discretion, on the application of any party to any proceedings before the Board, by writing under the hand of the Chairman, summon any person to appear as a witness before the Board.

7. The Chairman of the Board may administer an oath to any person appearing before the Board, whether the witness has been summoned or appears without being summoned before the Board, and may examine the witness upon oath.

8. If any person served with a summons to attend the Board fails without reasonable cause to attend the Board or to produce any documents, books or writings in his custody or control, which he was required by the summons to produce, he shall be guilty of an offence and shall be liable to a penalty of one hundred dollars.

9. If any person appearing as a witness before the Board refuses to be sworn, or to answer any question relevant to the proceedings before the Board put to him by any member thereof, he shall be guilty of an offence and shall be liable to a penalty of one hundred dollars:

Chairman may administer oath

Person failing to appear when summoned

Person refusing to make oath

Provided that nothing contained in this section shall render any person compellable to answer any question in respect of any matter which would have been protected from disclosure on the ground of privilege if the proceedings had been held in any court.

10. Any witness before the Board who knowingly gives false testimony touching any matter material to any inquiry shall be guilty of an offence, and shall be liable to a penalty of two hundred dollars or to imprisonment for twelve months.

11. The members of the Board shall not be personally liable for any act or default of the Board done or omitted to be done, in good faith, in administering this Act.

12.—(1) The Board may demand and collect, in advance, such fees as are prescribed.

False testimony Member not liable for acts of Board

Members not liable for acts of Board

Fees

(2) Such fees and all penalties and other moneys received or realised under this Act or under any rules made hereunder shall be paid into general revenue.

13. Any person thereto authorised in writing by the Chairman may enter any premises in which any pharmacist or licensed seller of poisons or medicines is carrying on business and may examine any books, papers, records or writings, drugs or medicines, whether patent or otherwise, or any article stored or offered for sale or used in the business.

14.—(1) The Minister may appoint from time to time a secretary to the Board.

Power of search

Secretary and inspectors
LN 46A of 1978

(2) The Minister may appoint inspectors for the purposes of enforcing the provisions of this Act or any rules made thereunder.

Powers of inspectors

15. For the purposes of enforcing the provisions of this Act or rules made hereunder, any inspector so appointed shall have the power at all reasonable times to enter upon the premises of any registered pharmacist or licensed seller of poisons or medicines and to inspect any books, papers, records or writings, drugs or medicines, whether patent or otherwise, or any article stored or offered for sale or used in the business; and shall have the power at all reasonable times to enter any premises in which he has reasonable cause to suspect that a breach of the law has been or is being committed, and to make such examination and inquiry and to do such other things (including the taking, on payment therefor, of samples) as may be necessary for the purpose of ascertaining whether the provisions aforesaid are being complied with.

PART III PHARMACISTSPART III

PHARMACISTS

16. The Board shall keep a register to be called the "Register of Pharmacists".

17.—(1) A person shall be registered by the entry in the register of his name and such other particulars relating to him as are prescribed.

Pharmacists how registered

(2) Every such entry in the register shall be signed by the Registrar of the Board.

(3) The Under Secretary (Health), Ministry of Health and Medical Services shall be the Registrar.

18.—(1) Subject to the provisions of this Act any person who is of good fame and character and who has passed the final examination of the Pharmaceutical Society of Great Britain or Northern Ireland, may be registered under the provisions of this Act.

Persons eligible for registration
7 of 1958, s. 2

(2) The Board may in its discretion, admit to the register any person who holds a pharmaceutical qualification other than that referred to in sub-section (1).

(3) No person shall be registered unless he has attained the age of twenty-one years.

19.—(1) The Board may direct that any person applying for registration as a pharmacist shall pass an examination and for that purpose may appoint an Examination Board consisting of the Under Secretary (Health), Ministry of Health and Medical Services as Chairman and of one or more members who shall be registered as pharmacists.

Board may direct examination of applicant
LN 46A of 1978

(2) The Board, by rules made under this Act with the approval of the Minister, may prescribe fees for such examination not exceeding ten dollars.

20. When any person has applied to be registered and has proved to the satisfaction of the Board—

(a) that he has attained the age of twenty-one years;

(b) that he is entitled to be registered by virtue of compliance with the requirements mentioned in sections 18 or 19; and

(c) that the certificate or diploma testifying to his qualification was, after examination, duly obtained by him from such a Society, Board or

the Board shall cause the person to be registered, by entering in the register his name and such other particulars as may be prescribed, and issue to him, upon payment of the prescribed fee, a certificate in the prescribed form.

21.—(1) If the Board refuses to register any person under this Act, the Board shall, if required by such person, state in writing the reasons for such refusal.

(2) Such person may thereupon appeal to the Court.

(3) An appeal under this section shall be by way of special case on any question of fact or law, and the Board shall, if the Court so orders, register the said person.

22.—(1) During the month of January in each year the Board shall cause to be published in the Gazette a true copy of the register.

Copy of register to be published

(2) A copy of the register so published shall be prima facie evidence of the registration of the persons named therein.

23. Any person who procures himself to be registered under this Act by means of any false or fraudulent representation or by the production of any false certificate or diploma shall be guilty of an offence and shall be liable to a penalty of two hundred dollars or to imprisonment for six months.

24. Any registered pharmacist who obtains or already possesses any higher degree, or any qualification other than the one qualification in respect of which he is registered, may have such higher degree or additional qualification entered in the register without payment of any additional fee.

25.—(1) Any registered pharmacist who changes his professional address shall forthwith give notice of the fact in writing to the Chairman.

Fraudulent representation

Amendments may be made in register

Notification of change of address or death LN 46A of 1978

(2) The Minister, upon receipt of a copy of the certificate of death of any pharmacist, shall cause notice thereof to be given in writing to the

Correction of register

26.—(1) The Board shall remove from the register the name of any registered pharmacist who has died and may make such alterations and amendments in the register as it thinks fit.

(2) The Board may, by notice in writing to any registered pharmacist addressed to him by registered post according to his address in the register, inquire whether he has changed his address or residence, and if an answer is not returned to such notice within six months after the date of the posting thereof, the Board may remove the name of such person from the register.

(3) The name of any registered pharmacist removed from the register under this Part of this Act be restored by the Board.

27.—(1) Subject to the provisions of this section, a body corporate carrying on a business which comprises the retail sale of drugs shall be an authorised seller of poisons, within the meaning of this Act, if the following conditions are complied with—

(a) the business shall, so far as concerns the keeping, dispensing and compounding of drugs and poisons, be under the management of a superintendent in relation to whom the following requirements are fulfilled—

(i) he shall be a registered pharmacist;

(ii) a statement in writing, signed by him on behalf of the body corporate, stating his name and stating whether or not he is a member of the board of directors shall have been sent to the Registrar;

(iii) he shall not be acting at the time in a similar capacity for any other body corporate; and

(b) in each set of premises where the business is carried on, the business shall, so far as concerns the retail sale of drugs, if not under the personal control of the superintendent, be carried on, subject to the directions of the superintendent under the personal control of a manager or assistant who is a registered pharmacist; and

(2) Notwithstanding the restrictions imposed by the provisions of this Act on the use of certain titles, emblems and descriptions, a body corporate which is an authorised seller of poisons may, if all the members of the board of directors are registered pharmacists, use the description of "chemist and druggist", or of "chemist", or of "druggist", or of "dispensing chemist", or of "dispensing druggist"; and may use the description of "pharmacy" in connection with the business:

Provided that nothing in this subsection shall authorise the use of any of the said descriptions in or upon any premises which are for the time being disqualified under this section from being registered in the register of premises, or in connection with any business so far as it is carried on in any premises so disqualified.

(3) If—

(a) a body corporate which is an authorised seller of poisons has been convicted of any offence under this Act; or

(b) any member of the board of directors, or any officer of that body, or any person employed by that body in carrying on the business, has been convicted of any such criminal offence or been guilty of any such misconduct as, in the opinion of the Board, renders him or would, if he were a registered pharmacist, render him unfit to be on the register,

the Board may inquire into the case and may, subject to the provisions of this Act, direct—

(i) that the body corporate shall cease to be an authorised seller of poisons and be disqualified for such period as may be specified in the direction from being an authorised seller of poisons; or

(ii) that any or all of the premises of the body corporate shall be removed from the register of premises and be disqualified, for such period as may be specified in the direction, from being registered therein.

(4) If the Board thinks fit in any case so to do, it may, either on its own motion or on the application of the body corporate concerned, direct that

Provided that where an appeal has been brought to the Court against a direction involving a period of disqualification, a direction under this subsection for a cesser of any disqualification subsisting by virtue of the direction as originally given, shall not take effect unless approved by the Minister.

(5) Any body corporate which has been disqualified in pursuance of this section may appeal by way of special case to the Court on any question of fact or law affecting the aforesaid disqualification, and the Board shall, if the Court so orders, set aside or modify the disqualification. LN 46A of 1978

(6) The body corporate shall pay for each separate set of premises a licence fee of ten dollars.

PART IV CONDUCT OF BUSINESS AS PHARMACISTPART IV

CONDUCT OF BUSINESS AS PHARMACIST

28.—(1) The Board shall remove from the register the name of any person—

Grounds of removal of name from register

(a) whose registration has been obtained by fraud or misrepresentation;

(b) who has ceased to possess, or does not possess, the qualifications in respect of which he was registered;

(c) who has been convicted in any part of Her Majesty's dominions, or elsewhere, of an indictable offence, or of any other offence which in the opinion of the Board renders him unfit to practise;

(d) who has been certified to be of unsound mind; or

(e) who is deemed by the Board guilty of—

(i) habitual drunkenness or habitual addiction to any drug;

(ii) such improper conduct as in the opinion of the Board renders him unfit to be allowed to continue to practise as a pharmacist.

(2) If the Board removes the name of any person from the register, it shall, if so required by him, state in writing the reason for the removal.

(3) Any person whose name has been removed from the register in pursuance of this section may appeal, by way of special case as aforesaid, to the Court to have his name restored to the register, and the Board shall, if the Court so orders, restore his name to the register.

Inquiry by the Board

29.—(1) Before removing from the register the name of any person, the Board shall make due inquiry, and such person may be represented by counsel, attorney or agent, who may examine witnesses and address the Board on his behalf.

(2) Pending the hearing of a charge against any person, the Board may suspend the registration of that person, who shall thereupon cease to practise.

30. Any person whose name is removed from the register under section 28 shall, within fourteen days after the posting of a notice demanding the return of his certificate of registration, surrender his certificate to the Board for cancellation; and any person who fails so to do shall be liable to a penalty of ten dollars for every day after the period of fourteen days during which the certificate is not returned.

31.—(1) Any person other than a registered pharmacist who carries on, or attempts to carry on, in any place or on any occasion, the business of a pharmacist, or pretends to be a pharmacist, or assumes or uses the title of pharmaceutical chemist, pharmacist, druggist, homeopathic chemist, dispensing chemist, or of member of any Pharmaceutical Society or Board, or takes or uses, in connection with the sale of goods, the title of chemist, shall be guilty of an offence, and shall be liable to a penalty of one thousand dollars.

Surrender of certificate of registration

Persons other than registered pharmacists not to carry on business

(2) No person shall use, in connection with any business, any title, emblem or description reasonably calculated to suggest that he, or anyone

For the purposes of this subsection the use of the description "pharmacy", in connection with a business carried on on any premises, shall be deemed to be reasonably calculated to suggest that the owner of the business and the person having the control of the business on these premises are registered pharmacists.

(3) If any person acts in contravention of the foregoing provisions of this section, he shall be liable, in respect of each offence, to a fine of one thousand dollars, and in the case of a continuing offence, to a further fine of fifty dollars for every day, subsequent to the day on which he is convicted of the offence, during which the offence continues.

32.—(1) Subject to the provisions of this section, if a registered pharmacist who is an authorised seller of poisons dies, or becomes of unsound mind, or is adjudged bankrupt, or enters into any arrangement with his creditors, any representatives who thereafter carry on his business in accordance with the conditions hereinafter specified and are persons in relation to whom the requirements of this section are satisfied, shall, for the purposes of that business and during the period specified in subsection (4), be authorised sellers of poisons within the meaning of this Acts, and be entitled to use, in conjunction with the business name of the pharmacist, such titles, emblems and descriptions as might have been used by the pharmacist.

(2) The conditions referred to in subsection (1) are as follows—

(a) in each set of premises where the business is carried on, the business, so far as concerns the retail sale of drugs, must be under the personal control of a registered pharmacist; and

(b) the name and certificate of registration of the person having the control of the business as aforesaid must be conspicuously exhibited in the premises.

(3) The requirements to be satisfied under subsection (1) in relation to the representatives are, that their names and addresses must be registered with the Registrar together with a statement of the name of the pharmacist whose representatives they are.

(4) The period referred to in subsection (1) shall be—

(a) in the case of the death of a pharmacist, a period of five years from the date thereof;

(b) in the case of the unsoundness of mind or bankruptcy of a pharmacist, a period of three years from the date when he became of unsound mind or was adjudged bankrupt;

(c) in the case of an arrangement with the creditors of a pharmacist, a period of three years from the date when the representatives became entitled thereunder to carry on his business,

or such longer period as, on the application of the representatives, the Board may, having regard to all the circumstances of the case, think fit to direct.

(5) If a representative, or a person employed by the representatives in the carrying on of the business, has been convicted of any such criminal offence or been guilty of any such misconduct as, in the opinion of the Board, renders him, or would, if he were a registered pharmacist, render him unfit to be on the register, the Board, after making inquiry into the case, may, subject to the provisions of this Act, direct that the representatives shall cease to be authorised sellers of poisons, and cease to be entitled to use the titles, emblems and descriptions which might have been used by the pharmacist.

(6) In this section the expression "representative" means an executor, administrator, trustee or committee, or a person authorised to exercise, in relation to a person of unsound mind not so found by inquisition, any of the powers of a committee, and, in respect of the period of three months after the death of a pharmacist leaving no executor who is entitled and willing to carry on his business, any person beneficially interested in the estate of the pharmacist.

33. Every pharmacist and every person or assistant under whose conduct or management the business of a pharmacist is carried on, shall have his name legibly painted or written and continually so maintained on a conspicuous place on the front of the building where the business is carried on.

34. Save as hereinafter provided, no person other than a registered

Name of pharmacist to be exhibited

Only pharmacists to dispense

Temporary licence

35.—(1) The Board may, upon the application of any registered pharmacist, issue a temporary permit to a pharmacist who possesses the qualifications mentioned in section 18, to act as locum tenens for such registered pharmacist for a period of three calendar months from the date of issue of the permit.

(2) The Board may renew any such permit for a further period of three months, but not for any longer period.

(3) The Board shall prescribe fees for such permit.

36.—(1) A medical practitioner shall not issue a prescription unless the prescription is signed by him with his usual signature, or is written on paper on which is printed his surname and the initials of his christian names, and bears the date on which the prescription was issued.

Prescriptions to be signed

(2) A prescription issued by a qualified veterinary surgeon shall, in addition to fulfilling the conditions laid down in the preceding subsection, bear the words "for veterinary purposes only".

(3) A prescription issued by a qualified dentist shall, in addition to fulfilling the conditions laid down in subsection (1), bear the words "for dental purposes only".

(4) A prescription which does not comply with the provisions of this section shall not be accepted by any pharmacist as authority for the sale or supply of any medicine or drug.

Record of prescriptions

37.—(1) Every pharmacist shall, as prescribed, record in a book (hereinafter called the "prescription book") to be kept by him for that purpose, every prescription of any medical practitioner dispensed, compounded or made up or supplied by him.

(2) Every prescription, whether issued by a qualified medical practitioner, qualified veterinary surgeon or qualified dentist, containing any of the drugs to which any Act as to the sale of dangerous drugs, for the time being in force, relates, shall be retained in the custody of the pharmacist dispensing the same for a period of two years and filed in the pharmacy.

(3) The prescription book shall be open for inspection by any inspector appointed under section 14.

38. A pharmacist shall not—

(a) keep or maintain any shop for selling or supplying medicines or drugs or for dispensing or compounding prescriptions, unless such shop is, while open for business, constantly under his own control or that of some other registered pharmacist as an assistant or agent of a registered pharmacist;

(b) permit any person, other than a bona fide assistant or apprentice in the course of his employment and under the actual personal supervision of a registered pharmacist, to sell, supply, compound or dispense medicines or drugs;

(c) permit any person, other than a registered pharmacist, to dispense or compound any prescription or supply any medicine or drugs containing any of the dangerous drugs to which subsection (2) of section 37 relates;

(d) carry on business except under the actual personal supervision of himself or some other registered pharmacist;

(e) practise pharmacy except under his own name;

(f) adopt the title "Consulting Chemist";

(g) give medical or surgical advice or aid, except in his place of business and—

(i) in the case of simple ailments of common occurrence;

(ii) in the administration of antidotes in the case of acute poisoning;

(iii) in the application of immediate aid in cases of accident or injury; or

(iv) in urgent cases under the direct instructions of a qualified medical practitioner;

(h) allow his name to be used in connection with the practice of pharmacy at any premises at which there is not a registered pharmacist in continual attendance; or

(i) aid or assist any person other than a registered pharmacist to practise pharmacy, except in accordance with the provisions of this Act.

39. Every medical practitioner, qualified veterinary surgeon or qualified dentist may dispense or compound any medicine or drugs for patients or animals without becoming a registered pharmacist, provided that a true and faithful record is made of every such prescription in the prescription book, which shall be open for inspection by any inspector or person duly authorised by the Board for that purpose.

40.—(1) Any person who—

Automatic machines for vending medicines prohibited

(a) installs any automatic machine for the sale or supply of any drug or medicine, or allows, permits or suffers any such automatic machine to be so installed;

(b) sells or supplies any drug or medicine by means of any such automatic machine;

(c) allows, permits or suffers any person to purchase or be supplied with or otherwise obtain any drug or medicine by means of any automatic machine,

shall be guilty of an offence, and shall be liable to a penalty of forty dollars, and in the case of a continuing offence, to a further penalty of ten dollars for every day, subsequent to the day on which he is convicted of the offence, during which the offence continues.

(2) For the purpose of the last preceding subsection, "automatic machine" means any machine or mechanical device used or capable of

41.—(1) Any person other than a qualified medical practitioner or a person acting under the direct instructions of such medical practitioner, who attends upon, prescribes for, or supplies any article as a drug, medicine, instrument or appliance to, any person for the alleviation, cure or treatment of any venereal disease, whether in fact such person is suffering from such disease or not, or of any disease affecting the generative organs or functions, or of sexual impotence, or of any complaint or infirmity arising or relating to sexual intercourse, or of female or menstrual irregularities, or for the purpose of terminating pregnancy, or of influencing the course of pregnancy, shall be guilty of an offence, and shall be liable to a penalty of two hundred dollars.

(2) Nothing in this section shall apply to—

(a) a registered pharmacist who dispenses to the patient of a qualified medical practitioner the prescription of such medical practitioner, if the prescription is dated and bears the address and the usual signature (including the surname) of the practitioner; or

(b) a registered pharmacist who, in the ordinary course of his business, sells or supplies any article as a drug, medicine, instrument or appliance (except such drugs, medicines, instruments or appliances as are prescribed), provided such drug, medicine, instrument or appliance is sold or supplied by such pharmacist for purposes other than those prescribed by this section.

42.—(1) No person shall publish any statement, whether by advertisement or otherwise, to promote the sale of any article as a medicine, instrument or appliance for the alleviation or cure of any venereal disease, or disease affecting the generative organs, or of sexual impotence, or of any complaint or infirmity arising from or relating to sexual intercourse, or of female or menstrual irregularities, or for terminating pregnancy, or for influencing the course of pregnancy, or for preventing conception.

(2) Any person who—

(a) affixes or inscribes any statement or any thing whatsoever so as to be visible to persons being in, or passing along, any street, road, highway, pathway, public place or public conveyance;

(b) delivers or offers or exhibits any statement to any person being

(c) throws any statement into or upon any street, road, highway, pathway, public place or public conveyance, or into the area, yard, garden or enclosure of any house;

(d) exhibits any statement to public view in any house, shop or place;

(e) prints or publishes any statement in any newspaper; or

(f) sells, offers or shows, or sends by post, any statement to any person,

shall be deemed to have published such statement.

(3) The word "statement" includes any document, book, or paper containing any statement.

(4) Any person who for himself or as assistant, servant, agent or manager does or permits any act, matter or thing contrary to this section or any part thereof, shall be guilty of an offence and shall be liable to a penalty of two hundred dollars.

(5) Nothing in this section shall apply to any books, documents or papers published in good faith for the advancement of medical or surgical science, or to any advertisement, notice or recommendation published by the authority of the Under Secretary (Health), Ministry of Health and Medical Services or to any publication sent only to qualified medical practitioners or registered pharmacists for the purpose of their business.

43. The British Pharmacopoeia as published in England under the direction of the General Council of Medical Education and Registration of the United Kingdom, in the edition for the time being in force, shall be the Pharmacopoeia in force in Solomon Islands as the standard of quality or composition for all drugs or medicines, and for the method of preparation of all drugs or medicines, and of compounding of all mixtures thereof; and for the purposes of this Act, the metre and the gramme shall be accepted respectively as legal units of measure and weight.

PART V SALE AND SUPPLY OF MEDICINES PART V

44.—(1) It shall not be lawful for any person who is not a registered pharmacist, or the assistant manager or bona fide apprentice of a registered pharmacist, to sell by retail any drug or medicines whatsoever, whether protected by letters patent, whether Imperial or Colonial, or not, except as prescribed by this Act.

(2) Nothing in this Act contained shall be construed to prohibit any licensed storekeeper from selling any of the articles mentioned in Schedule A.

(3) The Minister may, on the advice of the Board, by order add articles to or delete articles from Schedule A.

45.—(1) The Board may, on the application of any licensed storekeeper, grant such person a licence, to be called a Medicine Licence, to sell such articles as the Board deems fit:

Sale of drugs or medicines

Schedule A

LN 46A of 1978

Medicine licence 7 of 1958, s. 3

Provided that no such licence shall be granted to sell any of the drugs or medicines to which the provisions of subsection (2) of section 37 apply.

(2) Such licence shall be granted for a period not exceeding twelve months and may be renewed.

(3) The Board shall prescribe fees for such licence.

(4) The licence shall be in the form prescribed by rule hereunder and shall state clearly the names of all articles which the licensee is permitted to sell.

(5) Every application for a licence under this section shall be accompanied by a report by the Provincial Secretary of the province in which the business is carried on.

46. Immediately on the granting of a licence the Board shall so inform the Provincial Secretary or officer in charge of Police of that province in which the licence has been granted.

LN 88 of 1978

Police to be notified of issue of licence
LN 88 of 1978

47. A holder of such licence may sell or supply, or cause or suffer to be sold or supplied by his assistant or manager, only such drugs or medicines as, by virtue of such licence, he is entitled to sell or supply. Any person acting in contravention of this section shall be guilty of an offence and shall be liable to have his licence cancelled, and also to a penalty of forty dollars, and in the case of a continuing offence to a further penalty of ten dollars for every day, subsequent to the day on which he is found guilty of such offence, during which the offence continues.

48. It shall not be lawful for any person to sell any drug or medicine by wholesale to any person who does not possess a licence for the sale by retail of such drug or medicine.

49.—(1) It shall not be lawful for any person to import for sale by retail any drug or medicine which under his licence he is not entitled to sell or supply.

Only drugs mentioned in licence may be sold

Sale by wholesale of medicines or drugs

Importation of drugs or medicines

(2) Any drug or medicine imported in contravention of this section shall be liable to confiscation and shall be disposed of in such manner as the Comptroller of Customs and Excise may direct.

(3) Any person importing or attempting to import any drug or medicine in contravention of this section shall be guilty of an offence and shall be liable to a penalty of twenty dollars, and for a subsequent offence, to a penalty of two hundred dollars or to imprisonment for six months.

(4) The provisions of the Customs and Excise Act shall apply to proceedings under this section. Cap. 121

50. All medicines imported into Solomon Islands shall state on the label affixed to the container the percentage of proof spirit, if any, which the medicine contains; and in the case of a medicine containing a poison as one of the ingredients, such label shall state the proportion which the poison contained in the preparation bears to the total contents. In the case of such proportion being stated as a percentage, the statement shall indicate whether the percentage is weight in weight, weight in volume or volume in volume.

Labels on medicines imported
LN 88 of 1978

Provided that this section shall not apply to drugs, medicines, instruments or appliances imported by qualified medical practitioners, registered pharmacists, qualified veterinary surgeons or qualified dentists for bona fide medical, veterinary or dental treatment.

PART VI POISONSPART VI

POISONS

52.—(1) It shall not be lawful for any person to import any poison except under a licence issued by the Board:

Provided that this subsection shall not apply to the importation of poisons by qualified medical practitioners, registered pharmacists, qualified veterinary surgeons or qualified dentists for bona fide medical, veterinary or dental treatment.

(2) It shall not be lawful for any person to sell or deal in any of the several articles included in Schedule B hereto hereinafter referred to as the "Poisons List", except in the manner prescribed in this Act.

(3) The Minister may from time to time by order declare that any article named therein shall be deemed a poison within the meaning of this Act and be added to Part I or Part II of the Poisons List, as may be by such order directed.

(4) Any such order shall be published in the Gazette and on the expiration of three months from publication thereof, the article named therein shall be deemed to be added to such part of the said Schedule as may be directed in the order.

(5) Any person acting in contravention of this section shall be liable to a penalty of two hundred dollars, and in the case of a continuing offence, to a further penalty of ten dollars for each day, subsequent to the day on which he is convicted, during which the offence continues.

53. For the purposes of this Act all registered pharmacists shall be authorised sellers of poisons and may, subject to the provisions of this Act, sell and deal in poisons.

Pharmacists to be authorised sellers of poisons

54. On the application of any holder of a retail store licence, and on payment of the prescribed fee, the Board may issue to such person a licence to sell poisons, hereinafter referred to as a "Poisons Licence", provided that—

(a) such application is accompanied by a report, signed by the Provincial Secretary of the Province in which such retail store is situated, certifying that the applicant is a fit and proper person to hold such licence;

(b) such licence shall only apply to one place of business;

(c) no licence shall be granted empowering the holder thereof to sell or deal in any poisons included in Part I of the Poisons List;

(d) such licence shall be for a period of twelve calendar months and may be renewed; and

(e) such licence shall state specifically the poisons or class of poisons which the holder is licensed to sell or deal in.

55. The Board shall keep a book to be called the "Register of Premises", which shall be in the form prescribed by rules hereunder, and in which shall be entered the addresses of all premises where drugs, poisons or medicines are licensed to be sold, and such other particulars as may be prescribed by such rules.

56.—(1) Subject to the provisions of this Part of this Act it shall not be lawful—

Register of premises

Prohibition and regulations with respect to the sale of poisons

(a) for a person to sell any poison included in Part I of the Poisons List, unless—

(i) he is an authorised seller of poisons; and

(ii) the sale is effected on premises registered under section 55; and

(iii) the sale is effected by or under the supervision of a registered pharmacist;

(b) for a person to sell any poison included in Part II of the Poisons List, unless either—

(i) he is an authorised seller of poisons and the sale is effected on premises registered under section 55; or

(ii) he is the holder of a Poisons Licence and the sale is effected on premises registered under section 55;

(c) for a person to sell any poison, whether included in Part I or Part II of the Poisons List, unless the container of the poison is labelled in the prescribed manner—

(i) with the name of the poison; and

(ii) in the case of a preparation which contains a poison as one of the ingredients, with the prescribed particulars as to the proportion which the poison contained in the preparation bears to the total ingredients; and

(iii) with the word "poison" or other prescribed indication of the character of the article; and

(iv) with the name of the seller of the poison and the address of the premises on which it was sold.

(2) Subject to the provisions of this Part of this Act and to any rules made under this Act dispensing with or relaxing any of the requirements of this subsection—

(a) it shall not be lawful to sell any poison in Part I of the Poisons List to any person, unless that person is either—

(i) certified in the manner prescribed by rules and by a person authorised by rules to give a certificate for the purposes of this section; or

(ii) known by the seller or by some registered pharmacist in the employment of the seller at the premises where the sale is effected,

to be a person to whom the poison may properly be sold:

Provided that no poison shall be sold or delivered to any person under the age of twenty-one years;

(b) the seller of any such poison shall not deliver it until—

(i) he has made or has caused to be made an entry in a book to be kept for that purpose, hereinafter called the "Poisons Book", stating in the form prescribed by rules the date of the sale, the name and address of the purchaser and of the person, if any, by whom the certificate required under paragraph (a) of this subsection was given, the name and quantity of the article sold and the purpose for which it is stated by the purchaser to be required; and

(ii) the purchaser has affixed his signature to the entry aforesaid.

57.—(1) Nothing in the foregoing section shall apply—

Exemption with respect to medicines

(a) to a medicine which is supplied by a qualified medical practitioner for the purposes of medical treatment, by a qualified dentist for the purposes of dental treatment, or by a qualified veterinary surgeon for the purposes of animal treatment;

(b) to a medicine which is dispensed by a registered pharmacist at his place of business; or

(c) to a poison forming part of the ingredients of a medicine which is supplied by a registered pharmacist at his place of business:

Provided that the requirements contained in the following provisions of this section shall be satisfied in relation thereto.

(2) The medicine shall be distinctly labelled with the name and address of the person by whom it was supplied or dispensed.

(3) On the day on which the medicine was supplied or dispensed or, if that be not reasonably practicable, on the day next following that day, there shall be entered in the prescription book the following particulars—

(a) the date on which the medicine was supplied or dispensed;

(b) the ingredients of the medicine and the quantity thereof supplied;

(c) if the medicine was dispensed by a registered pharmacist the name or initials and, if it is known, the address of the person by whom, and the name and, if it is known, the address of the person to whom, and the date on which, the prescription was given;

(d) if the medicine was not so dispensed, the name and address of the person to whom it was supplied:

Provided that the provisions of this subsection shall, in the case of a medicine supplied on a prescription on which the medicine has been supplied by the seller on a previous occasion, be deemed to be complied with if the day on which the medicine is supplied and the quantity thereof supplied are entered in the prescription book on that day or, if that is not reasonably practicable, on the day next following that day, together with a sufficient reference to an entry in that book duly recording the dispensing of the medicine on the previous occasion.

(4) In the case of a medicine which is supplied or dispensed by a registered pharmacist and is compounded by the person supplying or dispensing it or by a person in his employment, the medicine shall have been compounded or dispensed by or under the immediate and personal supervision of a registered pharmacist.

(5) In the case of a medicine which is supplied or dispensed by a registered pharmacist, the supplying or dispensing of the medicine shall be effected by or under the immediate and personal supervision of a registered pharmacist.

58. Except as provided by rules made hereunder nothing in the foregoing provisions of this part of this Act shall extend to or interfere with—

Exemption with respect to sales wholesale and sales to certain persons

(a) the sale of poisons by wholesale dealing, provided that—

(i) such sale is to a registered pharmacist or to a holder of a poisons licence; or

(ii) such sale is to a person who requires the article—

(aa) for the purpose of his trade or business; or

(bb) for the purposes of enabling him to comply with any requirements made by or in pursuance of any Act with respect to the medical treatment of persons employed by that person in any trade or business carried on by him; or

(b) the sale of an article to a qualified medical practitioner, qualified dentist or qualified veterinary surgeon for the purposes of his profession.

59. It shall not be lawful for any holder of a poisons licence to use in connection with his business any title, emblem or description reasonably calculated to suggest that he is entitled to sell any poison other than a poison which he is under this Act entitled to sell; and if any person acts in contravention of this section, he shall be liable, in respect of each offence, to a fine of one hundred dollars, and in the case of a continuing offence, to a further penalty of ten dollars for each day, subsequent to the day on which he is convicted, during which the offence continues.

60. It shall not be lawful for a poison to be exposed for sale in or offered for sale by means of an automatic machine, and any person acting in contravention of this section shall be liable to a penalty of two hundred dollars, and in the case of a continuing offence, to a further penalty of ten dollars for each day, subsequent to the day on which he is convicted, during which the offence continues.

Use of titles, emblems and descriptions

Prohibition of sale of poisons by means of automatic machine

61.—(1) The Board, with the approval of the Minister, may make rules with respect to any of the following matters or for any of the following purposes—

(a) the manufacture of pharmaceutical preparations containing poisons;

(b) the sale, whether wholesale or retail, or the supply of poisons by or to any person or classes of persons, and in particular but without prejudice to the generality of the foregoing provisions—

(i) for regulating or restricting the sale or supply of poisons by holders of a poisons licence, and for prohibiting the sale of any specified poison or class of poisons by any class of such licensed sellers of poisons;

(ii) for prohibiting the sale by retail of poisons (being included in Part I of the Poisons List in Schedule B hereto) except on a prescription duly given by a duly qualified medical practitioner, qualified dentist or qualified veterinary surgeon, and for prescribing the form and regulating the use of prescriptions given for the purposes of rules made under this paragraph; Schedule B

(iii) for dispensing with or relaxing any of the provisions contained in Part VI of this Act relating to the sale of poisons;

(iv) the storage, transport and labelling of poisons;

(v) the containers in which poisons may be sold or supplied;

(vi) the additions to poisons of specified ingredients for the purposes of rendering them readily distinguishable as poisons;

(vii) the manufacture, compounding and dispensing of drugs and poisons;

(viii) the period for which any books required to be kept for the purposes of Part VI of this Act are to be preserved;

(ix) the period for which any certificate given under Part VI of this Act is to remain in force;

(x) for requiring persons in charge of the manufacture of pharmaceutical preparations containing poisons to be registered pharmacists;

(xi) for prescribing anything which by this Act is to be prescribed by rules;

(xii) the meetings and proceedings of the Board and the conduct of the business thereof and the duties of its officers;

(xiii) the forms to be used in pursuance of this Act;

(xiv) the manner of keeping the registers and the particulars to be entered therein;

(xv) the scale of fees to be charged and paid in respect of any application, registration, certificate or other proceedings, act or thing provided or required under this Act;

(xvi) the control of the professional conduct of registered pharmacists and the practice of the profession;

(xvii) the extent to which the British Pharmaceutical Codex, published by direction of the Pharmaceutical Society of Great Britain, or the Australasian Pharmaceutical Formulary, published by the Australasian Pharmaceutical Conference on behalf of the Pharmaceutical Societies of Australia and New Zealand, shall be accepted as a statement of official standards or quality or composition of drugs or medicines, and of the methods of preparation of drugs or medicines, and of compounding all mixtures thereof; and

(xviii) the qualifications of apprentices and assistants and the conditions under which apprentices or assistants may be employed;

(c) the conditions (including the keeping of records) to be observed

(2) The power to make rules under this section with respect to poisons includes the power to make rules with respect to any class of poisons or any particular poison.

General penalty

62.—(1) A person who acts in contravention of or fails to comply with any of the provisions of this Act, or any rule made under this Act, for which no specific penalty is prescribed, shall be liable to a penalty of not more than one hundred dollars, and in the case of a continuing offence, to a further penalty of twenty dollars for every day, subsequent to the day on which he is convicted, during which the offence continues.

(2) In the case of proceedings against a person under this section for or in connection with the sale, exposure for sale or supply of a poison effected by an employee—

(a) it shall not be a defence that the employee acted without the authority of the employer; and

(b) any material fact known to the employee shall be deemed to have been known to the employer.

(3) Notwithstanding any enactment prescribing the period within which proceedings may be commenced, proceedings for an offence under this Act may be commenced at any time within the period of twelve months next after the date of the commission of the offence; or, in the case of proceedings instituted by or by the direction of the Director of Public Prosecutions, either within the period aforesaid or within the period of three months next after the date on which evidence sufficient, in the opinion of the Director of Public Prosecutions, to justify a prosecution for the offence comes to his knowledge, whichsoever period ends on the later date. For the purposes of this subsection, a certificate purporting to be signed by the Director of Public Prosecutions as to the date on which such evidence as aforesaid came to his knowledge shall be conclusive evidence thereof.

63. Articles the importation of which is prohibited by this Act and, to the extent to which their importation is prohibited, articles the importation of which is restricted by this Act, shall be deemed to be goods the importation of which is prohibited under the Customs and Excise Act; and subject to the provisions of this Act, the said Act and any Act amending the same shall apply to such articles.

6 of 1953, Sched LN 46A of 1978

Application of Customs and Excise Act
Cap. 121

SCHEDULES

SCHEDULE A

LN 78 of 1973 LN 5 of 1988

(Section 44, subsection (2))

Tablets, capsules and lozenges

Asprin Tablets B.P. (in packs of not more than 25)

Asprin soluble tablets B.P. (in packs of not more than 25)

Benzalkonium lozenges B.P.C.

Disprin

Paracetamol B.P.

(in packs of not more than 25)

(in packs of not more than 25)

Ointments and Applications

Calamine lotion B.P.

Centrimide cream B.P.

Chlorozylenol Solution B.P.C.

Lanolin cream

Medicated powder (not containing antibiotics)

Medicated soap

Methylsalicylate liniment B.P.C.

Vick Vapour rub

Zinc cream B.P.

Zinc, starch and talc dusting powder B.P.C.

Dressings

Adhesive plasters

Bandages

Cotton wool, hospital quality

Lint, surgical

Surgical dressings, not containing any antibiotic

Surgical gauze, unmedicated

Miscellaneous

Cod liver oil

Dextrose

Enos (and similar effervescent antacids)

Eucalyptus

Lactogen

Sodium bicarbonate

Soda crystals (washing soda)

Vick inhaler.

Note: Proprietary preparations which, in the opinion of the Pharmacy and Poisons Board, are of substantially similar composition, pharmacoligical action and toxicity to any of the above preparations, may also be sold.

LN 60 of 1981

SCHEDULE B

(Section 52, subsection (2))

THE POISONS LIST

PART I

Acetanilide; alkyl acetanilides.
Alkali fluorides other than those specified in Part II of this List.
Alkaloids, the following; their salts, simple or complex —

- Acetyldihydrocodeinone; its esters.
- Aconite, alkaloids of.
- Aporphine.
- Atropine.
- Belladonna, alkaloids of.
- Benzylmorphine.
- Benzylmorphine.
- Brucine.
- Calabar bean; alkaloids of.
- Coca, alkaloids of
- Cocaine.
- Codeine.
- Colchicine.
- Coniine.
- Cotarnine.
- Curarine.
- Diacetylmorphine.
- Dihydrocodeinone; its esters.
- Dihydrohydroxycodeinone; its esters.
- Dihydromorphine; its esters.
- Dihydromorphinone; its esters.
- Ecgonine; its esters.
- Emetine.
- Ephedra; alkaloids of.
- Ergot; alkaloids of.
- Ethylmorphine.
- Gelsemium; alkaloids of.
- Homatropine.
- Hyoscine.
- Hyoscyamine.
- Jaborandi; alkaloids of
- Lobelia; alkaloids of.
- Morphine
- Papaverine.
- Pomegranate; alkaloids of.
- Quebracho; alkaloids of, other than the alkaloids of red quebracho.
- Sabadilla; alkaloids of.
- Solanaceous alkaloids not otherwise included in this List.
- Staphisagre; alkaloids of.
- Strychnine.
- Thebaine.
- Veratrum; alkaloids of.
- Yohimbe; alkaloids of.

Allylisopropylacetylurea.
Amidopyrine; its salts.
Amino-alcohols, esterified with benzoic acid, phenylacetic acid, phenylpropionic acid, cinnamic acid or the derivatives of these acids.
Amphetamines (beta-aminopropylbenzene and beta-amino-isopropylbenzene).
Amyl nitrite
Antimony, chlorides of, oxides of antimony, sulphides of antimony; antimonates; antimonides; organic compounds of antimony.
Arsenical substances, the following, except those specified in Part II of this List; arsenic; halides of; oxides of arsenic; arsenates, arsenites, organic compounds of arsenic.
Barbituric acid; its salts; derivatives of barbituric acid; their salts; compounds of barbituric acid, its salts, its derivatives, their salts with any other substance.
Barium, salts of, other than barium sulphate and the salts of barium specified in Part II of this List
Butyl chloral hydrate.

Cannabis (the dried flowering or fruiting tops of Cannabis sativa Linn.), the resin of cannabis; extracts of cannabis, tinctures of cannabis; cannabin tannate.

Cantharidin; cantharidates.

Chloral formamide.

Chloral hydrate.

Chloroform.

Creosote obtained from wood.

Croton; oil of.

Digitalis, glycosides of; other active principles of digitalis.

Dinitrocresols; dinitronaphthols, dinitrophenols, dinitrothymols.

Elaterin.

Ergot (the sclerotia of any species of Claviceps); extracts of ergot; tinctures of ergot.

Erythrityl tetranitrate.

Fluorocetamide

Fluorocetanilide

Glyceryl trinitrate.

Guanidines, the following: polymethylene diguanidines, dipara-anisylphenetyl guandine.

Hydrocyanic acid; cyanides; double cyanides of mercury and zinc.

Insulin.

Lead acetates; compounds of lead with acids from fixed oils.

Mannityl hexanitrate.

Mercury, oxides of; nitrates of mercury; mercuric ammonium chlorides; potassio-mercuric iodides; mercuric oxycyanides; mercuric thiocyanate.

Metanitrophenol; orthonitrophenol; paranitrophenol.

Monofluoracetic acid; or its salts

Nux Vomica.

Opium.

Orthocaine; its salts.

Ouabain.

Oxalic acid; metallic oxalates other than potassium quadroxalate.

Oxycinchroninic acid, derivatives of; their salts; their esters.

Para-amino-benzoic acid; esters of; their salts.

Phenetidylphenacetin.

Phenols (any member of the series of phenols of which the first member is phenol and of which the molecular composition varies from member to member by one atom of carbon and two atoms of hydrogen) except in substances containing less than sixty per cent, weight in weight, of phenols; compounds of phenol with a metal, except in substances containing less than the equivalent of sixty per cent weight in weight, of phenols.

Phenylcinchoninic acid; salicylcinchoninic acid; their salts; their esters.

Phenylethylhydantoin; its salts; its acyl derivatives; their salts.

Phosphorus, yellow.

Picric acid.

Picrotoxin.

Pituitary gland, the active principles of.

Savin; oil of.

Sodium monofluoroacetate syn: Sodium mononuoracetic acid; commonly known as compound 1080

Strophanthus; glycosides of strophanthus.

Sulphonal; alkyl sulphonals.

Suprarenal gland, the active principles of, their salts.

Thallium; salts of.

Thyroid gland, the active principles of, their salts.

Tribromethyl alcohol.

P-aminobenzenesulphonamide, Sulphonilamide, and preparations thereof and analagous compounds and derivatives and preparations thereof, whether described as Prontosil, Prontylin, Septasine, Soluseptasine, Sulphonamide-p or any other trade-name, trade-mark or designation.

PART II

Ammonia.

Arsenical substances, the following—

Arsenical substances, the following—

Arsenic sulphides.
Arsenious oxide.
Calcium arsenates.
Calcium arsenites.
Copper acetoarsenites.
Copper arsenates.
Copper arsenites.
Lead arsenates.
Potassium arsenites.
Sodium arsenates.
Sodium arsenites.
Sodium thioarsenates.

Barium, salts of, the following—

Barium carbonate.

Barium silicofluoride.

Formaldehyde.

Hydrochloric acid.

Hydrofluoric acid; potassium fluoride; sodium fluoride; sodium silicofluoride.

Mercuric chloride; mercuric iodide; organic compounds of mercury.

Methylated spirits.

Nicotine; its salts.

Nitric acid.

Nitrobenzene.

Phenols as defined in Part I of this List in substances containing less than sixty per cent weight in weight, of phenols; compounds of phenol with a metal in substances containing less than the equivalent of sixty per cent, weight in weight, of phenols.

Phenylene diamines; toluene diamines, their salts.

Potassium hydroxide.

Potassium quadroxalate.

Sodium hydroxide.

Sulphuric acid.

CHAPTER 105 PHARMACY AND POISONSCHAPTER 105

PHARMACY AND POISONS

Subsidiary Legislation APPOINTMENT OF INSPECTORSubsidiary Legislation

(Section 14)

The Medical Officer for the time being in charge of the Public Hospital at Gizo is appointed to be an inspector for the purposes of enforcing the provisions of the Pharmacy and Poisons Act. 107/133/1958

THE POISONS RULES

(Section 61)

Rules by the Pharmacy and Poisons Board

Rules by the Pharmacy and Poisons Board

1. These Rules may be cited as the Poisons Rules.

2.—(1) In these Rules, unless the context otherwise requires, the following expressions have the meanings hereby respectively assigned to them, that is to say—

Definitions

"animal" includes poultry;

"antimonial poisons" means chlorides of antimony, oxides of antimony, sulphides of antimony, antimonates, antimonites, and organic compounds of antimony;

"arsenical poisons" means halides of arsenic, oxides of arsenic, sulphides of arsenic, arsenates, arsenites, copper acetoarsenites, sodium thioarsenates, and organic compounds of arsenic;

"British Pharmacopoeia" and "British Pharmaceutical Codex" include supplements;

"food" includes a beverage;

"licensed seller of poisons" means a person entitled under Part VI of the Act to sell poisons included in Part II of the Poisons List;

"medicines for the internal treatment of human ailments" includes any medicine to be administered by hypodermic injection but does not include any mouth-wash, eye-drops, eye-lotion, ear-drops, douche or similar article;

"poisons list" means the Poisons List contained in Schedule B of the Act;

"sale exempted by section 58 of the Act" means a sale made in such circumstances as to be entitled except as provided by these Rules, to exemption under section 58 of the Act from the foregoing provisions of Part VI of the Act;

"transaction exempted by section 57 of the Act" means the supply of a medicine in such circumstances as to be entitled to exemption under section 57 of the Act from the provisions of section 56 of the Act.

(2) In these Rules any reference to an alkaloid shall include a reference to any salt of that alkaloid and in the case where the esters of an alkaloid are included in the Poisons List by virtue of the words "its esters" to any esters of that alkaloid.

(3) Any reference in the Schedules to these Rules to the percentage of a poison contained in any substance or preparation shall unless otherwise expressly provided be construed in the following manner, that is to say, a reference to a substance or preparation containing one per cent of any poison means—

(a) in the case of a solid that one gramme of the poison is contained in every hundred grammes of the substance or preparation;

(b) in the case of a liquid that one millilitre of the poison or if the poison itself is a solid one gramme of the poison is contained in every hundred millilitres of the substance or preparation;

and so in proportion for any greater or less percentage.

3. It shall not be lawful for any person to sell any poisons on any premises used for or in connection with his retail business notwithstanding that the sale is exempted by section 58 of the Act unless he complies with the provisions of paragraph (a) or paragraph (b), as the case may be, of section 56(1) of the Act.

4.—(1) Subject as hereinafter provided the provisions of paragraph (c) of section 56(1) of the Act and of rules 14 to 19 hereof (which provisions relate to the labelling of poisons) shall apply to sales exempted by section 58 of the Act and shall also apply to the supply of poisons (otherwise than on sale) in like manner as if references in the said provisions to the sale and seller of poisons included references to the supply and supplier of poisons respectively.

(2) The said provisions except the provisions of rule 18 and of paragraph (c) (iv) of section 56(1) as modified by rule 19 shall not apply to the sale or supply of any of the poisons included in the Second Schedule to these Rules to a person who requires the poison for the purpose of his trade or business if the outside of the package in which the poison is sold or supplied is labelled conspicuously with words indicating the dangerous properties of the poison.

5. The provisions of section 56(2) of the Act (which makes provision as to persons to whom poisons may be sold and to the keeping of records of sales) shall apply to all substances included in the First Schedule to these Rules whether or not the poison sold is a poison included in Part I of the Poisons List and shall not apply with respect to any other substance:

Prohibition against sale on retail premises

Application of provisions as to labelling

Second Schedule

Application of section 56 (2) of Act
First Schedule

Provided that paragraph (a) of section 56(2) shall in its application to sales as authorised sellers of Part II poisons be deemed to be satisfied if the person to whom the poison is sold is known by the person in charge of the premises on which the poison is sold or of the department of the business in which the sale is effected to be a person to whom the poison may properly be sold.

6.—(1) The provisions of section 56(2) of the Act as modified by the last foregoing rule shall apply to sales exempted by section 58 of the Act and shall also apply to the supply in the form of a commercial sample otherwise than on sale of any substance included in the First Schedule to these Rules in the like manner as if references in the said provisions to the sale and seller of poisons respectively included references to the supply and supplier of poisons in the form of commercial samples.

(2) Paragraph (a) of section 56(2) shall in its application to sales exempted by section 58 of the Act and to the supply in the form of commercial samples of substances included in the First Schedule to these Rules be deemed to be satisfied if the person to whom the poison or sample is sold or supplied is known by the person in charge of the department of the business through which the sale or supply is effected to be a person to whom the poison or sample may properly be sold or supplied.

(3) So much of paragraph (b) of section 56(2) as requires an entry in a book to be signed by the purchaser of a poison shall not as respects the sale of a poison to a person for the purposes of his trade, business or profession apply if the following requirements are satisfied—

(a) the seller must obtain before the completion of the sale an order in writing signed by the purchaser stating his name and address, trade, business or profession, the name and quantity of the article to be purchased and the purpose for which it is required;

(b) the seller must be reasonably satisfied that the signature is that of the person purporting to have signed the order and that that person carries on the trade, business, or profession stated in the order being one in which the poison to be purchased is used;

(c) if the article is sent by post it must be sent by registered post;

(d) the seller must insert in the entry prescribed by rule 34 hereof the words "signed order" and a reference number by which the order can be identified:

Provided that where a person represents that he urgently requires a poison for the purpose of his trade, business or profession the seller may if he is reasonably satisfied that the person so requires the poison and is by reason of some emergency unable before delivery to furnish to the seller an

If any purchaser by whom any such undertaking has been given fails to deliver to the seller a signed order in accordance with the undertaking or if any person for the purpose of obtaining delivery of any poison under the foregoing proviso makes a statement which is to his knowledge false he shall be deemed to have contravened the provisions of this rule.

7. The requirements mentioned in section 57 (3) of the Act (which requires particulars of medicines supplied or dispensed under that section to be entered in a book) need not be satisfied in the case of any medicine not being a substance included in the First Schedule to these Rules which is supplied by—

(a) a medical practitioner or qualified veterinary surgeon for the purpose of medical or animal treatment; or

(b) a registered pharmacist on and in accordance with a prescription given by a medical practitioner.

8. Nothing in these Rules shall apply except as is expressly provided therein to transactions exempted by section 57 of the Act.

9. Such of the provisions of these Rules and of Part VI of the Act as modified by these Rules as apply solely with respect to the substances included in the First Schedule to these Rules shall not apply with respect to—

Exemption of certain preparations
First Schedule

(a) machine spread plasters; or

(b) surgical dressings; or

(c) articles containing barium carbonate and prepared for the destruction of rats and mice; or

(d) corn paints in which the only poison is a poison included in the Poisons List under the heading of "Cannabis".

Exemption of articles,

(a) with respect to any article included in Group I of the Third Schedule to these Rules; or

(b) so far as any poison specified in the first column of Group II of that Schedule is concerned with respect to any of the articles or substances specified in the second column opposite the description of the poison.

11.—(1) It shall not be lawful to sell any poison included in the Fourth Schedule to these Rules except on and in accordance with the prescription given by a registered medical practitioner, registered dentist or a qualified veterinary surgeon in the form provided by this rule.

(2) This rule shall apply to the sale of any such poison notwithstanding that it is a transaction exempted by section 57 of the Act but shall not apply to any sale exempted by section 58 of the Act.

(3) For the purposes of this rule a prescription shall—

(a) be in writing and be signed by the person giving it with his usual signature and be dated by him;

(b) specify the address of the person giving it;

(c) specify the name and address of the person for whose treatment it is given or if the prescription is given by a qualified veterinary surgeon of the person to whom the medicine is to be delivered;

(d) have written thereon if given by a dentist the words "For Dental Treatment Only" or if given by a qualified veterinary surgeon the words "For Animal Treatment Only";

(e) indicate the total amount of the medicine to be supplied and the dose to be taken.

(4) The person dispensing the prescription shall comply with the following requirements—

(a) the prescription must not be dispensed more than once unless the prescriber has stated thereon that it may be dispensed more than once;

(b) if the prescription contains a direction that it may be dispensed a stated number of times or at stated intervals it must not be dispensed otherwise than in accordance with the directions;

(c) at the time of dispensing there must be noted on the prescription above the signature of the prescriber the name and address of the seller and the date on which the prescription is dispensed;

(d) except in the case of a prescription which may be dispensed again the prescription must for a period of two years be retained and kept on the premises on which it was dispensed in such manner as to be readily available for inspection.

12. It shall not be lawful for a registered pharmacist to sell any substance included in the First Schedule to these Rules notwithstanding that the poison is a poison included in Part II of the Poisons List unless the sale is effected by or under the supervision of a registered pharmacist.

13.—(1) No licensed seller of poisons shall be entitled by virtue of being a licensed seller of Part II poisons to sell—

(a) any poison other than ammonia, hydrochloric acid, methylated spirits, nitric acid, potassium quadroxalate, and sulphuric acid except in a closed container as closed by the manufacturer or other person from whom the poison was obtained;

(b) any substance included in the First Schedule to these Rules unless the sale is effected by himself or by a responsible deputy.

In this paragraph the expression "responsible deputy" means a person nominated as a deputy on the seller's form of application as hereinafter prescribed for licence as a licensed seller of Part II poisons or any person substituted by notice in writing to the Pharmacy and Poisons Board for a person so nominated and not more than two deputies shall be nominated at one time in respect of one set of premises.

(2) No person shall be entitled by virtue of being a licensed seller of poisons to sell any poison included in the first column of the Fifth Schedule to these Rules unless the article or substance sold is one of the articles or substances specified against the description of the poison in the second column of that Schedule and the container of the substance is, in addition to any other direction of the Act or of these Rules with respect to labelling, labelled clearly with a notice of the special purpose for which the article or substance is intended and a warning that it is only to be used for that purpose.

(3) It shall not be lawful to sell or supply strychnine except as an ingredient in medicine:

Provided that this rule shall not apply to the sale of strychnine—

(a) by way of wholesale dealing;

(b) for the purpose of being compounded in medicines prescribed or administered by a registered medical practitioner or qualified veterinary surgeon;

(c) to a person or institution concerned with scientific education or research or chemical analysis for the purposes of that education, research or analysis.

Labelling

14.—(1) Subject to the provisions of these Rules the particulars with which the container of a poison is required to be labelled under paragraph (c) of section 56(1) of the Act and under these Rules must appear in a conspicuous position on the container in which the poison is sold and on every box or other covering of whatever nature enclosing the container and the paticulars must be clearly and distinctly set out and not in any way obscured or obliterated.

(2) Where the poison is contained in an ampoule, cachet or similar article it shall not be necessary to label the article itself if every box or other covering in which the article is enclosed is duly labelled.

(3) Nothing in the said paragraph (c) or in Rules 14 to 19 shall require the labelling of any transparent cover or any wrapper, hamper, packing case, crate or other covering used solely for the purpose of transport or

15.—(1) Subject as hereinafter provided for the purpose of paragraph (c) (i) of section 56 (1) of the Act the name of a poison shall be the term under which it is included in the Poisons List:

Provided that when the said term describes a group of poisons and not the poison specifically, the name of the poison shall be—

(a) if the poison is the subject of a monograph in either the British Pharmacopoeia or the British Pharmaceutical Codex one or other of the names or synonyms or abbreviated names set out at the head of the monograph; and

(b) in any other case the accepted scientific name or name descriptive of the true nature and origin of the poison.

(2) For the purposes of the foregoing it shall in the case of a preparation in the British Pharmacopoeia or the Formulary to the British Pharmaceutical Codex or any dilution or admixture of such a preparation or any surgical dressing for which a standard is described in the British Pharmaceutical Codex be sufficient notwithstanding anything in the foregoing paragraph of this rule to state the name, synonym or abbreviated name used to describe the preparation, or surgical dressing in the British Pharmacopoeia or the Formulary to the British Pharmaceutical Codex with the addition of the letters B.P. or B.P.C. as the case may be.

16.—(1) For the purposes of paragraph (c) (ii) of section 56 (1) of the Act (which requires preparations containing poisons to be labelled with the prescribed particulars as to the proportions of a poison therein) the label of the container of any preparation containing a poison as one of its ingredients shall subject as hereinafter provided include a statement of the proportion which the poison bears to the total ingredients of the preparation.

(2) In the case of a preparation containing a poison specified in the first column of the Sixth Schedule to these Rules it shall be sufficient to state on the label the particulars specified in the second column of that Schedule against the description of the poison.

(3) In the case of a preparation or surgical dressing which is named in accordance with paragraph (2) of the last foregoing rule it shall not be

(4) Where the poison is in tablets, pills, capsules, cachets, lozenges or similar articles or in ampoules it shall be sufficient to state on the label of the box or other covering in which the articles are enclosed the number of the articles and the amount of the poison or in the case of such a preparation as is mentioned in the last foregoing paragraph the amount of the preparation contained in each article.

(5) Where any proportion is stated as a percentage the statement shall indicate whether the percentage is calculated on the basis of weight in weight, weight in volume, or volume in volume.

17.—(1) In pursuance of paragraph (c) (iii) of section 56 (1) of the Act (which requires the containers of poison to be labelled with the word "poison" or other prescribed indication of character) the container of any article specified in the Seventh Schedule to these Rules shall instead of being labelled with the word "poison" be labelled with the words specified in the said Schedule as applicable to that article.

(2) The said words or the word "Poison" as the case may be must not be modified in meaning by the addition of any other words or marks, and—

(a) in the case of a substance included in the First Schedule to these Rules must either be in red lettering or be set against a red background; and

(b) in all cases must either be on a separate label or be surrounded by a line within which there must be no other words except words with which the container of the poison is required to be labelled under the Act or these Rules.

18.—(1) It shall not be lawful to sell or supply any poison—

Prohibition of certain sales
Directions on container

(a) in the case of a liquid other than a medicine contained in a bottle of a capacity of not more than 120 fluid ounces unless the bottle is labelled with words "Not to be taken";

(b) in the case of an embrocation, liniment, lotion, liquid antiseptic

(2) It shall not be lawful to sell or supply any compressed hydrocyanic acid unless the container is labelled with the words "Warning. This container holds poisonous gas and should only be opened and used by persons having expert knowledge of the precautions to be taken in its use."

(3) This rule shall be in addition to the other requirements of the Act and these Rules with respect to labelling and shall apply to transactions exempted by section 57 of the Act.

19.—(1) The provisions of paragraph (c) (iv) of section 56 (1) of the Act (which requires the container of a poison to be labelled with the name of the seller and the address of the premises on which it was sold) shall not apply in the case of an article sold for the purpose of being sold again in the same container.

Name and address of seller on label

(2) The requirements of the said paragraph shall be deemed to be satisfied in the case of a poison supplied from a warehouse or depot if the container of the poison is labelled with the address of the supplier's principal place of business or in the case of a limited company of the registered office of the company.

(3) When any poison (other than a substance included in the First Schedule to these Rules) is sold in a container and outer covering, being the container and covering in which it was obtained by the seller, it shall be sufficient if the name of the seller and the address of the premises on which it was sold appear only on the outer covering. First Schedule

(4) Where the names of more than one person or more than one address appear on any label there must also be words on the label indicating clearly which person is the seller and at which of the addresses the poison was sold.

20.—(1) It shall not be lawful to sell, whether by wholesale or retail, or supply any poison unless—

(a) it is contained in a container impervious to the poison and sufficiently stout to prevent leakage arising from the ordinary risks of handling and transport; and

(2) Sub-paragraph (b) of the foregoing paragraph shall apply to transactions exempted by section 58 of the Act.

21.—(1) It shall not be lawful to store any poison except in a container impervious to the poison and sufficiently stout to prevent leakage from the container arising from the ordinary risks of handling.

(2) It shall not be lawful to store any substance included in the First Schedule to these Rules in any retail shop or premises used in connection therewith unless the container is stored—

(a) in a cupboard or drawer reserved solely for the storage of poisons;

(b) in a part of the premises which is partitioned off or otherwise separated from the remainder of the premises and to which the customers are not permitted to have access;

(c) on a shelf reserved solely for the storage of poisons and—

(i) no food is kept directly under the shelf; and

(ii) the container of the substance is rendered distinguishable by touch from the containers of articles and substances other than poisons stored upon the same premises:

Provided that in the case of any such substance to be used in agriculture or horticulture, it shall not be lawful to store the substance in any such part of the premises as aforesaid if food is kept in that part, or in any cupboard or drawer unless the cupboard or drawer is reserved solely for the storage of poisons to be used as aforesaid.

22. It shall not be lawful to consign any poison for transport unless it is sufficiently stoutly packed to avoid leakage arising from the ordinary risks of handling and transport.

23.—(1) It shall not be lawful to consign for transport by carrier any

Transport of poisons

Restrictions on carriage Eighth Schedule

(2) It shall not be lawful for any person knowingly to transport any such poison as aforesaid either on his own behalf or for another person in any vehicle in which food is being transported unless the food is carried in a part of the vehicle effectively separated from that containing the poison or is otherwise adequately protected from the risk of contamination.

(3) This rule shall not apply with respect to medicines.

24. In all establishments in which pharmaceutical preparations containing any poison are manufactured for the purpose of the internal treatment of human ailments the preparation must be manufactured by or under the supervision of a registered pharmacist:

Provided that this rule shall not apply to the manufacture by or under the supervision of a qualified medical practitioner of preparations containing pituitary, suprarenal or thyroid glands, the active principles of any of those glands or the salts of the active principles of thyroid gland.

25. Every application for a licence to sell poisons included in Part II of the Poisons List shall be as set out in the Ninth Schedule hereto.

26. Every licence to a storekeeper to sell such Part II poison as is permitted by these Rules shall be as set out in the Tenth Schedule hereto.

27. The fees payable for each licence or certificate granted or premises registered under the Act shall be as set out in the Eleventh Schedule hereto.

28. The form of Certificate of Registration as a Pharmacist shall be as set out in the Twelfth Schedule hereto.

29. The Register of Pharmacists shall be in the form as set out in the Thirteenth Schedule hereto.

30. The register of premises where drugs, poisons or medicines are

Application to sell poison
Ninth Schedule

Storekeeper's poison licence
Tenth Schedule

Fees
Eleventh Schedule

Form of certificate Twelfth Schedule

Form of Register of Pharmacists Thirteenth Schedule

Form of register of

31. The form of application for a licence to sell medicines shall be as set out in the Fifteenth Schedule hereto.

32. (1) A licence to sell medicines shall be in the hereto form prescribed in the Sixteenth Schedule hereto and shall be subject to such terms and conditions as may therein be specified.

(2) Any person who, being the holder of such a licence, contravenes any of its terms or conditions or causes or permits any of its terms or conditions to be contravened shall be guilty of an offence and liable to a fine of fifty dollars.

33.—(1) A certificate given for the purposes of paragraph (a) of section 56 (2) of the Act being a certificate certifying a person to be a person to whom a poison may properly be sold shall be in the form and shall contain the particulars set out in the Seventeenth Schedule hereto.

(2) All householders are hereby authorised to give such certificates as aforesaid:

Provided that a certificate given by a householder who is not known to the seller of the poison to be a responsible person of good character shall not be a sufficient certificate for the purposes of the said paragraph unless it is endorsed in the manner specified in the Seventeenth Schedule by a police officer of or above the rank of Inspector.

Certificate in accordance with section 56 (2)

Seventeenth Schedule

11 of 1970, Sched

(3) On any sale of a poison on such a certificate as aforesaid the certificate shall be retained by the seller.

34. The particulars of sales of poisons which are required by paragraph (b) of section 56(2) of the Act to be entered in a book shall be entered in the form set out in the Eighteenth Schedule hereto.

35. All books kept for the purpose of Part VI of the Act shall be preserved on the premises on which the sales recorded therein were made for a period of two years from the date on which the last entry was made therein.

Form of entry of sales

Eighteenth Schedule Preservation of books

SCHEDULES

FIRST SCHEDULE

SUBSTANCES FALLING WITHIN THE POISONS LIST TO WHICH SPECIAL RESTRICTIONS APPLY

Alkaloids, the following, their salts simple or complex—

Acetyldihydrocodeinone.

Aconite, alkaloids of, except substances containing less than 0.02 per cent of the alkaloids of aconite.

Apomorphine, except substances containing less than 0.2 per cent of apomorphine.

Atropine, except substances containing less than 0.15 per cent of atropine.

Belladonna, alkaloids of, except substances containing less than 0.15 per cent of the alkaloids of belladonna calculated as hyoscyamine.

Benzoylmorphine.

Benzylmorphine.

Brucine, except substances containing less than 0.2 per cent of brucine.

Calabar bean, alkaloids of.

Coca, alkaloids of, except substances containing less than 0.1 per cent of the alkaloids of coca.

Cocaine, except substances containing less than 0.1 per cent of cocaine.

Codeine, except substances containing less than 0.1 per cent of codeine.

Colchicine, except substances containing less than 0.5 per cent of colchicine.

Coniine, except substances containing less than 0.1 per cent of coniine.

Cotarnine, except substances containing less than 0.2 per cent of cotarnine.

Curarine.

Diacetylmorphine.

Dihydrocodeinone.

Dihydrohydroxycodeinone.

Dihydromorphine.

Dihydromorphinone.

Ecgonine, except substances containing less than 0.1 per cent of ecgonine.

Emetine, except substances containing less than 1 per cent of emetine.

Ergot, alkaloids of.

Ethylmorphine, except substances containing less than 0.2 per cent of ethylmorphine.

Gelsemium, alkaloids of, except substances containing less than 0.1 per cent of the alkaloids of gelsemium.

Homatropine, except substances containing less than 0.15 per cent of homatropine.

Hyoscine, except substances containing less than 0.15 per cent of hyoscine.

Hyoscyamine, except substances containing less than 0.15 per cent of hyoscyamine.

Jaborandi, alkaloids of, except substances containing less than 0.5 per cent of the alkaloids of jaborandi.

Lobelia, alkaloids of, except substances containing less than 0.5 per cent of the alkaloids of lobelia.

Morphine, except substances containing less than 0.2 per cent of morphine calculated as anhydrous morphine.

Nicotine.

Papaverine, except substances containing less than 1 per cent of papaverine.

Pomegranate, alkaloids of, except substances containing less than 0.5 per cent of the alkaloids of pomegranate.

Quebracho, alkaloids of.

Sabadilla. alkaloids of, except substances containing less than 1 per cent of the alkaloids of sabadilla.

Solanaceous alkaloids, not otherwise included in this Schedule except substances containing less than 0.15 per cent of solanaceous alkaloids calculated as hyosyamine.

Stavesacre, alkaloids of, except substances containing less than 0.2 per cent of the alkaloids of stavesacre.

Strychnine, except substances containing less than 0.2 per cent of strychnine.

Thebaine, except substances containing less than 1 per cent of thebaine.

Veratrum, alkaloids of, except substances containing less than 1 per cent of the alkaloids of veratrum.

Yohimba, alkaloids of.

Allylisopropylacetylurea.

Amidopyrine, its salts.

Amino-alcohols, esterified with benzoic acid, phenylacetic acid, propylpropionic acid, cinnamic acid or the derivatives of those acids, excepting substances containing less than 10 per cent of the esterified amino-alcohols.

Antimonial poisons, excepting substances containing less than the equivalent of 1 per cent of antimony trioxide.

Arsenical poisons, except substances containing less than the equivalent of 0.01 per cent of arsenic trioxide.

Barbituric acid, its salts, derivatives of barbituric acid, their salts, compounds of barbituric acids, its salts, its derivatives, their salts, with any other substance.

Barium, salts of.

Cannabis, the resins of cannabis, extracts of cannabis, tinctures of cannabis, cannabin tannate.

Cantharidin, except substances containing less than 0.01 per cent of cantharidin.

Cantharidates, except substances containing less than the equivalent of 0.01 per cent of cantharidin.

Digitalis, glycosides of, except substances containing less than one unit of activity (as defined in the British Pharmacopoeia) in two grammes of the substance.

Dinitrocresols, Dinitronaphthols, Dinitrophenols, Dinitrothymols.

Ergot, extracts of ergot, tinctures of ergot.

Fluorocetamide LN 61/1981

Fluorocetanilide

Guanidines, the following, polymethylene diguanidine, dipara-anisyl-phenetyl guanidine.

Hydrocyanic acid, except substances containing less than 0.1 per cent of hydrocyanic acid (HCN), cyanides except substances containing less than the equivalent of 0.1 per cent weight in weight of hydrocyanic acid (HCN) double cyanides of mercury and zinc.

Lead, compounds of, with acids from fixed oils.

Mercuric Chloride, except substances containing less than 1 per cent of mercuric chloride, mercuric iodide, except substances containing less than 2 per cent of mercuric iodide, nitrates of mercury except substances containing less than the equivalent of 3 per cent weight in weight of mercury (Hg) potassio-mercuric iodides excepting substances containing less than the equivalent of 1 per cent of mercuric iodide, organic compounds of mercury excepting substances less than the equivalent of 0.2 per cent weight in weight of mercury (Hg).

Metanitrophenol, orthonitrophenol, paranitrophenol.

Monofluoracetic acid; its salts LN 61/1981

Nux Vomica, except substances containing less than 0.2 per cent of strychnine.

Opium, except substances containing less than 0.2 per cent of morphine calculated as anhydrous morphine.

Ouabain.

Oxycinchoninic acid, derivatives of their salts, their esters.

P-aminobenzenesulphonamide, Sulphanilamide, and preparations thereof and analogous compounds and derivatives and preparations thereof, whether described as Prontosil, Prontylin, Septasine, Soluseptasine, Sulphonamide-P or any other trade-name, trade-mark or designation.

Phenetidylphenacetin.

Phenylcinchoninic acid, salicyl-cinchoninic acid, their salts, their esters.

Phenylethylhydantoin, its salts, its acyl derivatives, their salts.

Picrotoxin.

Savin, oil of.

Sodium monofluoroacetate syn: Sodium monofluorocetic acid; commonly known as compound 1080 LN 61/1981

Strophanthus, glycosides of.

Thallium, salts of.

Tribromethyl alcohol.

SECOND SCHEDULE

POISONS EXEMPTED BY RULE 4(2) FROM LABELLING PROVISIONS WHEN SOLD IN CERTAIN CIRCUMSTANCES

Alkali fluorides.

Ammonia.

Antimony, chlorides of, oxides of antimony, sulphides of antimony, antimonates, antimonites.

Chloroform.

Dinitrocresols, dinitronaphthols, dinitrophenols.

Formaldehyde.

Glyceryl trinitrate.

Hydrochloric acid.

Hydrofluoric acid, sodium silicofluoride.

Lead acetate, compounds of lead with acids from fixed oils.

Mercuric chloride, mercuric iodide, organic compounds of mercury.

Mercury, oxides of, nitrates of mercury.

Metanitrophenol, orthonitrophenol, paranitrophenol.

Nitric acid.

Nitrobenzene.

Oxalic acid, metallic oxalates.

Phenols, compounds of phenol with a metal.

Phosphorus, yellow.

Picric acid.

Potassium hydroxide.

Sodium hydroxide.

Sulphuric acid.

THIRD SCHEDULE

ARTICLES EXEMPTED BY RULE 10 FROM THE PROVISIONS OF THE ACT AND OF THESE RULES

Group I

GENERAL EXEMPTIONS

Adhesives.

Anti-fouling compositons.

Builders' materials.

Ceramics.

Distempers.

Electrical valves.

Enamels.

Explosives.

Fillers.

Fireworks.

Glazes.

Glue.

Lacquer solvents.

Loading materials.

Marking inks.

Matches.

Motor fuels and lubricants.

Paints other than pharmaceutical

paints.

Photographic paper.

Pigments.

Plastics.

Polishes.

Printers' ink

Propellants.

Rubber varnishes.

Group II

SPECIAL EXEMPTIONS

Poison

Acetanilide, alkyl acetanilides

Substance or article in which exempted

Substances not being preparations for the treatment of human ailments.

Alkaloids—

Emetine.

Ephedra, alkaloid of.

Jaborandi, alkaloids of.

Lobelia, alkaloids of.

Nicotine.

Ipecacuanha, extracts and tincytures of ipecacuanha; substances containing less than 0á5 per cent of emetine.

Substances containing less than 1 per cent of the alkaloids of ephedra.

Substances containing less than 0á025 per cent of the alkaloids of jaborandi.

Preparations for the relief of asthma in the form of cigarettes, smoking mixtures or fumigants; substances containing less than 0.1 per cent of the alkaloids of lobelia.

Tobacco.

Pomegranate, alkaloids of. Peomegranate bark.

Solanaceous alkaloids.

Stavesacre, alkaloids.

Ammonia.

Stramonium contained in preparations for the relief of asthma in the form of cigarettes, smoking mixtures, or fumi-gants.

Soaps, ointments, lotions for external use.

Substances not being solutions of ammonia or preparations containing solutions of ammonia, liquids containing les than 5 per cent weight in eight of ammonia (NH_3), refrigerators, smelling bottles.

Arsenical poisons.

Chloroform.

Pyrites ores, or sulphuric acid containing arsenical poisons as natural impurities.

Substances containing les than 10 per cent of chloroform.

Creosote obtained from wood. Substances containing less than 50 per cent of creosote obtained from wood.

Formaldehyde.

Hydrochloric acid.

Lead acetate.

Lead, compounds of.

Mercuric chloride.

Substances containing less than 5 per cent weight in weight of formaldehyde (H.CHO) photographic glazing or hardening solutions.	
Substances contaiing less than 9 per cent weight in weight of hydrochloric acid (HCl).	
Substances containing less than 4 per cent of lead acetate.	
Machine spread plasters.	
Batteries.	
Mercuric chloride, mercuric iodide, organic compounds of mercury.	Dressings on seeds or bulbs.

Poison
Mercury, nitrates of.
Nitric acid
Nitrobenzene.

Substance or article in which exempted
Ointments containing less than the quivalent of 3 per cent weight in weight of mercury (Hg).
Substances containing less than 9 per cent weight in weight of nitric acid (HNO_3).
Substances containing less than 0.1 per cent of nitro-benzene, soaps containing less than 1 per cent of nitro-benzene.

Phenols.	Carvacrol; coal tar, crude or refined; creosote obtained from coal tar; essential oils in which phenols occur naturally; medicines, containing less than 1 per cent of phenols; nasal sprays, mouthwashes, pastilles, lozenges, capsules, pessaries, ointments or suppositories containing less than 2.5 per cent of phenols; smelling bottles; soaps for washing; solid substances containing less than 60 per cent of phenols; tertiary butyl cresol; thymol.

Phenylenediamines, toluene diamines; their salts.

Substances other than preparations for the dyeing of hair.

Picric acid.

Potassium hydroxide.

Sodium fluoride.

Sodium hydroxide.

Sodium silicofluoride.

Sulphuric acid.

Substances containing less than 5 per cent of picric acid.

Substances containing less than 12 per cent of potassium hydroxide.

Substances containing less than 3 per cent of sodium fluoride as a preservative.

Substances containing less than 12 per cent ofsodium hydroxide.

Substances containing less than 3 per cent of sodium silicofluoride as a preservative.

Substances containing less than 9 per cent weight in weight of sulphuric acid acid (H_2SO_4), accumulators, batteries, fire extinguishers.

LN 102/1987
LN 63/1988

FOURTH SCHEDULE

LN 102/1987
LN 61/1988

SUBSTANCES REQUIRED BY RULE 17 TO BE SOLD BY RETAIL ONLY ON A PRESCRIPTION GIVEN BY A REGISTERED MEDICAL PRACTITIONER, REGISTERED DENTIST OR QUALIFIED VETERINARY SURGEON

Acebutolol Hydrochloride
Acepromazine
Acepromazine Maleate
Acetamide
Acetarsol
Acetazolamide
Acetazolamide Sodium
Acethydrocodone
Acetohexamide
Acetorphine
Acetylcholine Chloride; but if in preparations for external use and ms 0.2% Schedule 1
Acetylcysteine
Acetyldihydrocodeine
Acetyldigitoxin
Acetylmethadol
Acetylstrophanthidin
Acetyl Sulphafurazole
Acetyl Sulphamethoxypyridazine
Aconite Root; but if in preparations for external use and ms 1.3% of the crude drug Schedule 1
Aconitine; but if in preparations for external use and ms 0.02% Schedule 1
Aconitine Hydrobromide, but if in preparations for external use and ms 0.02% (calculated as base) Schedule 1
Aconitine Hydrochloride; but if in preparations for external use and ms 0.02% (calculated as base) Schedule 1
Aconite Nitrate; but if in preparations for external use and ms 0.02% (calculated as base) Schedule 1
Acrosoxasin
ACTH preparations
Actinomycin C
Actinomycin D
Acyclovir
Adrolin
Adiphenine Hydrochloride
Adrenaline; but if:
(a) in inhalers Schedule 1; or
(b) in preparations for external use Schedule 1
Adrenaline Acid Tartrate; but if:
(a) in inhalers Schedule 1; or
(b) in preparations for external use Schedule 1
Adrenaline Hydrochloride; but if:
(a) in inhalers Schedule 1; or
(b) in preparations for external use Schedule 1
Adreno-cortical Extract
Adriamycin
Aklomide
Alclofenac
Alcuronium Chloride
Aldosterone
Alevac Preparations
Albumin
Alfacalcidol
Algestone
Algestone Acetonide
Algestone Acetophenide
Allobarbitone
Allylprodine
Allopurinol
Allyloestrenol
Alphadolone Acetate
Alphameprodine
Alphamethadol
Alphaprodine
Alphaxalone
Alprenolol
Alprenolol Hydrochloride
Alprostadil
Alseroxylon
Altizide
Amantadine Hydrochloride
Ambenonium Chloride
Ambusilk
Ambutonium Bromide
Amcinonide
Amelazole Hydrochloride
Amethocaine, but if in preparations for non parenteral use (Schedule 1) (except preparations for local ophthalmic use)

Amethocaine Gentisate: but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Amethocaine Hydrochloride, but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Amidopyrine
Amikacin Sulphate
Amiloride Hydrochloride
Aminocaproic Acid
Aminoglutethimide
Aminophylline Injection
Aminopterin Sodium
Aminosalicylic Acid
Amiodarone Hydrochloride
Amiphenazole Hydrochloride
Amitryptyline
Amitryptyline Embonate
Amitryptyline Hydrochloride
Ammonium Bromide
Amoxapine
Ampicillin
Amoxycillin Trihydrate
Amphetamine
Amphomycin
Amphotericin
Ampicillin
Ampicillin Sodium
Ampicillin Trihydrate
Amygdalin
Amylobarbitone
Amylobarbitone Sodium
Amylocaine Hydrochloride: but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Anerod
Androsterone
Angiotensin Amide
Anileridine
Anterior Pituitary Extract
Antimony Barium Tartrate
Antimony Dimercaptosuccinate
Antimony Lithium Thiomalate
Antimony Pentasulphide
Antimony Potassium Tartrate
Antimony Sodium Tartrate
Antimony Sodium Thioglycollate
Antimony Sulphate
Antimony Trichloride
Antimony Trioxide
Antimony Trisulphide
Apiol
Apomorphine
Apomorphine Hydrochloride
Apramycin
Apramycin Sulphate
Aprobarbitone
Aprobarbitone Sodium
Aprotinin
Arecoline
Arecoline-Acetarsol
Arecolin Hydrobromide
Argipressin
Arsanilic Acid
Arsenic
Arsenic Triiodide
Arsenic Trioxide
Arsphenamine
Atenolol
Atenolol
Atropine: but if:
(a) not combine with Hyoscine or Hyoscyamine or their salts and:
(i) in inhalers Schedule 1
(ii) in preparations for internal use with not 300 micrograms and not 1 mg Schedule 1; or
(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)
(b) combined with Hyoscine or Hyoscyamine or their salts and:
(i) in inhalers Schedule 1 or
(ii) in preparations for internal use and not 1 mg of the total alkaloids (calculated as base) Schedule 1; or
(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use) Atropine Methobromide: but if:
(a) not combined with Hyoscine or Hyoscyamine or their salts and:
(i) in inhalers Schedule 1; or
(ii) in preparations for internal use with mg 400 micrograms

and mdd 1.3 mg Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

(c) combined with Hyoscine or Hyoscyamine or their salts and:

(i) in inhalers Schedule 1; or

(ii) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

Atropine methonitrate; but if:

(a) not combine with Hyoscine or Hyoscyamine or their salts and:

(i) in inhalers Schedule 1; or

(ii) in preparations for internal use with mdd 400 micrograms and mdd 1.3 mg Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

(b) combined with Hyoscine or Hyoscyamine salts and:

(i) in inhalers Schedule 1; or

(ii) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

Atropine oxide hydrochloride, but if:

(a) not combined with Hyoscine or Hyoscyamine or their salts and:

(i) in inhalers Schedule 1; or

(ii) in preparations for internal use with mdd 360 micrograms and mdd 1.2 mg Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

(b) combined with Hyoscine or Hyoscyamine or their salts and:

(i) in inhalers Schedule 1; or

(ii) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

Atropine sulphate; but if:

(a) not combined with Hyoscine or Hyoscyamine or their salts and:

(i) in inhalers Schedule 1; or

(ii) in preparations for internal use with mdd 360 micrograms and mdd 1.2 mg Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

(b) combined with Hyoscine or Hyoscyamine or their salts and:

(i) in inhalers Schedule 1; or

(ii) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

Azacyclonol

Azacyclonol Hydrochloride

Azaperone

Azapropazone

Azathioprine

Azathioprine Sodium

Azidocillin Potassium

Bacampicillin hydrochloride

Bacitracin

Bacitracin methylene disalicylate

Bacitracin Zinc

Baloleten

Bambermycin

Barbitone

Barbitone Sodium

Barium Carbonate

Barium Chloride

Barium Sulphide

Beclamide

Beclomethasone
Beclomethasone Dipropionate
Belladonna Herb; but if:
(a) in preparations for internal use and mdd 1 mg of the alkaloids Schedule 1; or
(b) in preparations for external use Schedule 1
Belladonna Root; but if:
(a) in preparations for internal use and mdd 1 mg of the alkaloids Schedule 1; or
(b) in preparations for external use Schedule 1
Bemegride
Bemegride Sodium
Benactyzine Hydrochloride
Benactyzine Hydrochloride
Bendrofluazide
Benethamine Penicillin
Benoxaprofen
Benperidol
Benserazide
Benzamine Lactate, but if in preparations for non-parenteral use Schedule 1
Benzathine Penicillin
Benzbromarone
Benzhexol Hydrochloride
Benzilonium Bromide
Benzocaine; but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Benzoctamine Hydrochloride
Benzoestrol
N-Benzol Sulphanilamide
Benzquinamide
Benzquinamide Hydrochloride
Benzthiazide
Benztrophine Mesylate
Benzylmorphine
Benzylpenicillin
Benzylpenicillin Calcium
Benzbromarone
Benzoyl Peroxide (Hunts)
Benzpheramine
Benoxinate
Beroperidol
Benserazide
Benethamine Penicillin
Benacetylmethadol
Betamiprodine
Betamethadol
Betahistine Hydrochloride
Betamethasone
Betamethasone Adamantoate
Betamethasone Benzoate
Betamethasone Sodium Phosphate
Betamethasone Valerate
Betaprodine
Bethanechol Chloride
Bethanidine Sulphate
Bezafibrate
Bezitramide
Bicillin
BiCNU
Biperiden Hydrochloride
Biperiden Lactate
Bismuth Glycollylarsanilate
Bleomycin Sulphate
Boldenone Undecylenate
Bretylium Tosylate
Bromhexine Hydrochloride
Bromazepam
Bromocriptine Mesylate
Bromvalerone
Brotizolam
Budesonide
Bufexamac
Bumetanide
Buphenine Hydrochloride; but if in Preparations for internal use with mdd 6 mg and mdd 18 mg Schedule 1
Bupivacaine; but if in preparations for non parenteral use Schedule 1 (except preparations for local ophthalmic use)
Bupivacaine hydrochloride; but if in preparations for non parenteral use Schedule 1 (except preparations for local ophthalmic use)
Buprenorphine
Buprenorphine Hydrochloride
Busulphan
Butacaine Sulphate; but if in preparations for non parenteral use Schedule 1 (except preparations for local ophthalmic use)
Butalbital
Butalbital Sodium

Bupivacaine Phosphate; but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Butobarbitone
Butobarbitone Sodium
Butriptyline Hydrochloride
Butyl Aminobenzoate; but if in preparations for non-parenteral use Schedule 1
Butylchloral Hydrate

Calcitonin
Calcitriol
Calcium 5-Allyl-5-N-Butylbarbiturate
Calcium Aminosalicylate
Calcium Amphomycin
Calcium Benzamidosalicylate
Calcium Bromide
Calcium Bromidolactobionate
Calcium Carbimide
Calcium Folinate
Calcium Sulphadoxate
Candicidin
Canrenoic Acid
Cantharidin; but if in preparations for external use and not 0.01% Schedule 1
Capreomycin Sulphate
Caramiphen Hydrochloride; but if:
(a) in tablet preparations and not 7.5 mg (calculated as base) Schedule 1
(b) in liquid preparations and not 0.1% (calculated as base) Schedule 1
Carbachol
Carbalon
Carbamazepine
Carbenicillin Sodium
Carbenoxolone sodium; but if:
(a) in pellets with not 5 mg acid and not 25 mg Schedule 1; or
(b) in gels and not 2% Schedule 1
Carbimazole
Carbocaine
Carbocysteine
Carbon Tetrachloride
Carcromal
Carbuterol
Carfecillin Sodium
Carindacillin
Carisoprodol
Carmustine
Carperidine
CCNU
Cefadezone
Cefaclor
Cefadroxil
Cefoxitin Sodium
Cephalexin
Cephalexin Sodium
Cephaloglycin
Cephaloram
Cephaloridine
Cephalosporin C
Cephalosporin P
Cephalosporin N
Cephalothin Sodium
Cephazolin Sodium
Cephradine
Cerium Oxalate
Chloral Antipyrine
Chloral Betaine
Chloral Formamide
Chloral Glycerolate
Chloral Hydrate; but if in External preparations for human use Schedule 1
Chloralose
Chloralcrethane
Chlorambucil
Chloramphenicol
Chloramphenicol Cinnamate
Chloramphenicol Palmitate
Chloramphenicol Sodium Succinate
Chlordiazepoxide
Chlordiazepoxide Hydrochloride
Chlorhexadol
Chlorphentermine
Chlorisondamine Chloride
Chlormadinone Acetate
Chlormerodrin
Chlormethiazole
Chlormethiazole Edisylate
Chlormezanone
Chloroform
Chloroquine Phosphate; but if for the prophylaxis of malaria Schedule 1
Chloroquine Sulphate; but if for the prophylaxis of malaria Schedule 1

Chlorothiazide
Chlorotrianisene
Chlorphenoxamine Hydrochloride
Chlorpromazine
Chlorpromazine Embonate
Chlorpromazine Hydrochloride
Chlorpropamide
Chlorprothixene
Chlortetracycline
Chlortetracycline Calcium
Chlortetracycline Hydrochloride
Chlorthalidone
Chlorzoxazone
Cholestyramine
Chorionic Gonadotrophin
Cicloacillin
Cimetidine
Cimetidine Hydrochloride
Cinchocaine, but if preparations for non-parenteral use and ms 3% Schedule 1 (except preparations for local ophthalmic use)
LN 93/1988
Cinchocaine Hydrochloride: but if in preparations for non-parenteral and ms 3% (calculated as base) Schedule 1 (except preparations for local ophthalmic use)
Cinchophen
Cinoxacin
Cisplatin
Clenbuterol Hydrochloride
Clidinium Bromide
Clindamycin
Clindamycin Hydrochloride Hydrate
Clindamycin Palmitate Hydrochloride
Clindamycin Phosphate
Clioquinol; but if:
(a) in external preparations for human use Schedule 1; or
(b) in preparations for internal human use for treatment of mouth ulcers and ms 35 mg and mdd 350 mg Schedule 1
Clobazam
Clobetasol 17 Propionate
Clobetasone Butyrate
Clofazimine
Clofibrate
Clomiphene Citrate
Clomipramine
Clomipramine Hydrochloride
Clonidine
Clonidine Hydrochloride
Clonitazine
Cloparide
Cloprostenol Sodium Salt
Clopenthixol
Clorexolone
Clorprenaline Hydrochloride
Closterbol Acetate
Clotrimazole; but if:
(a) in creams for external use Schedule 1; or
(b) in solutions for external use Schedule 1
Cloxacillin Benzathine
Cloxacillin Sodium
Cocaine
Coccules Indicus
Codeine; but if:
(a) in linctus for treatment of cough ms 0.3% Schedule [illegible]; or
(b) in compound analgesic preparations ms 10 mg as phosphate per unit dose, mdd 80 mg Schedule 1.
Colaspase
Colchicine
Colestipol Hydrochloride
Colistin Sulphate
Colistin Sulphomethate
Colistin Sulphomethate Sodium
Conine
Conium leaf; but if in preparation for external use and ms 7% of the crude drug Schedule 1
Corticotorphin
Cortisone
Cortisone Acetate
Cortodoxone
Cotarnine Chloride
Co-trimoxazole
Cropropamide
Crotethamide
Croton Oil
Croton Seed
Curare
Cyclobarbitone
Cyclobarbitone Calcium
Cyclobendazole

Cyclofenil
Cyclopenthiazide
Cyclopentolate Hydrochloride
Cyclophosphamide
Cycloprostin
Cycloserine
Cyclothiazide
Cyproterone Acetate
Cytarabine
Cytarabine Hydrochloride
Dacarbazine
Danazol
Dantrolene Sodium
Dapsone
Dapsone Ethane Ortho Sulphonate
Daunorubicin Hydrochloride
Deanol Bitartrate; but if in preparations for internal use and mdd 26 Schedule 1
Debrisoquine Sulphate
Dehydroemetine Hydrochloride
Delcycroepiandrosterone
Delmadinone Acetate
Demecarium Bromide
Demeclocycline
Demeclocycline Calcium
Demeclocycline Hydrochloride
Deoxycortone Acetate
Deoxycortone Pivalate
Deptropine Citrate
Dequalinium Chloride; but if:
(a) in throat lozenges or throat pastilles and ms 0.25 mg Schedule 1; or
(b) in external paint preparations and ms 1% Schedule 1
Desarpidine
Desdeethioxantone Lesylate
Desfluorotriamcinolone
Desipramine Hydrochloride
Deslanoside
Desmopressin
Desonide
Desoxymethasone
Dexamphetamine
Dexamethasone
Dexamethasone Acetate
Dexamethasone 21 Isonicotinate
Dexamethasone Phenylpropionate
Dexamethasone Pivalate
Dexamethasone Sodium m-Sulphobenzoate
Dexamethasone Sodium Phosphate
Dexamethasone Trioxaundecanoate
Dexetimide
Dexedrine
Dextromethorphan Hydrobromide; but if in preparations for internal use with mdd 15 mg (calculated as base) and mdd 75 mg (calculated as base) Schedule 1
Dextromethorphan Hydrochloride; but if in preparations for internal use with mdd 15 mg (calculated as base) and mdd 75 mg (calculated as base) Schedule 1
Dextromoramide
Dextropropoxyphene Hydrochloride
Dextropropoxyphene Napsylate
Dextrothyroxine Sodium
Diacetylmorphine
Diamorphine
Diamorphomide
Diazepam
Diboxide
Dibenzepin Hydrochloride
Diclofenac Sodium
Dichlorulphenazone
Dichlorophenarsine Hydrochloride
Dichlorphenamide
Dicyclamine Hydrochloride; but if in preparations for internal use with mdd 10 mg and mdd 60 mg Schedule 1
Diethanolamine Fusidate
Diethylarsine Acetarsol
Diethylenbutene
Diethylpropion Hydrochloride
Difenoxin
Difluocortolone Valerate
Diflurasal
Digitalin
Digitalis Leaf
Digitalis Prepared
Digitoxin
Digoxin
Dihydrallazine Sulphate
Dihydrocodeine
Dihydroergotamine Mesylate
Dihydroergotoxine Mesylate
Dihydromorphine
Dihydrostreptomycin
Dihydrostreptomycin Sulphate
Diloxanide Furoate

Dimenoxadole
Dimepheptanol
Dimepregen
Dimercaprol
Dimethisoquin Hydrochloride; but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Dimethisterone
Dimethothiazine Mesylate
Dimethylthiambutene
Dimethyl Sulphoxide
Dimethyltubocurarine Bromide
Dimethyltubocurarine Chloride
Dimethyltubocurarine Iodide
Dimetridazole
Dinitrodiphenylsulphonylethylenediamine
Dinoprost
Dinoprostone
Dioxaphetyl
Dipipanone
Diphetarsone
Diprenorphine Hydrochloride
Dipyridamole
Dipyrone
Disopyramide
Disopyramide Phosphate
Distigmine Bromide
Disulphine Blue
Disulfiram
Disulphamide
Dithiazol (limits)
Dobutamine Hydrochloride
Domperamide
Dopamine Hydrochloride
Dothiepin
Dothiepin Hydrochloride
Doxapram Hydrochloride
Doxepin Hydrochloride
Doxorubicin
Doxycycline
Doxycycline Calcium Chelate
Doxycycline Hydrochloride
Droperido
Drostano one
Drostanolone Propionate
Drotelsuicol
Dydrogesterone
Dyflos

Ecgonine
Econazole
Econazole Nitrate
Econtropate Iodide
Ekgestrone
Edrophonium Chloride
Embutramide
Emepronium Bromide
Emetine; but if in preparations for internal or external use and ms 1% Schedule 1
Emetine Bismuth Iodide; but if in preparations for internal or external use and ms 1% (calculated as emetine) Schedule 1
Emetine Hydrochloride; but if in preparations for internal or external use and ms 1% (calculated as emetine) Schedule 1
Ephedrine Hydrochloride; but if:
(a) in preparations for internal use (except nasal sprays or nasal drops) with not 30 mg (calculated as base) Schedule 1; or
(b) in nasal sprays or nasal drops and ms 2% (calculated as base) Schedule 1 or
(c) in preparations for external use Schedule 1
Ephedrine Sulphate; but if:
(a) in preparations for internal use (except nasal sprays or nasal drops) with not 30 mg (calculated as base) and not 60 mg (calculated as base) Schedule 1, or
(b) in nasal sprays or nasal drops and ms 2% (calculated as base) Schedule 1; or
(c) in preparations for external use Schedule 1
Epicillin
Epiestriol
Epitiazide
Epoprostenol
Ergometrine Maleate
Ergometrine Tartrate
Ergot, Prepared
Ergotamine Tartrate

Ergotoxine Esylate
Erythromycin
Erythromycin Estolate
Erythromycin Ethyl Carbonate
Erythromycin Ethyl Succinate
Erythromycin Lactobionate
Erythromycin Phosphate
Erythromycin Stearate
Erythromycin Thiocyanate
Estramustine Phosphate
Etafedrine Hydrochloride
Ethacrynic Acid
Ethambutol Hydrochloride
Ethamivan
Ethamsylate
Ethchlorvynol
Ethebenecid
Ethiazide
Ethinyloestradiol
Ethionamide
Ethisterone
Ethoglucid
Ethoheptazine Citrate
Ethopropazine Hydrochloride
Ethosuximide
Ethotoin
Ethyl Acetanilide
Ethyl Biscoumacetate
Ethylmethylthiambutene
Ethylmorphine
Ethyloestrenol
Ethylstibamine
Ethynodiol Diacetate
Etidronate Disodium
Etomidate
Etonitazene
Etoposide
Etorphine
Etoxeridine

Famprofazone
Fazadinium Bromide
Fenbufen
Fencamfamin Hydrochloride
Fenclofenac
Fenfluramine Hydrochloride
Fenoprofen
Fenoprofen Calcium
Fenoterol Hydrobromide
Fenpipramide Hydrochloride
Fenpiprane Hydrochloride
Fentin Compounds
Feprazol
Feprazone
Ferrous Arsenate
Flavoxate Hydrochloride
Fluanisone
Flubendazole
Fluclorolone Acetonide
Flucloxacillin Sodium
Flucytosine
Flubendazole
Fludrocortisone Acetate
Flufenamic Acid
Flugestone
Flugestone Acetate
Flumedroxone Acetate
Flumethasone
Flumethasone Pivalate
Flumethiazide
Flunisolide
Flunitrazepam
Fluocinolone Acetonide
Fluocinonide
Fluocortolone
Fluocortolone Hexanoate
Fluocortolone Pivalate
Fluopromazine Hydrochloride
Fluorometholone
Fluorouracil
Flourouracil Trometamol
Fluoxymesterone
Flupenthixol Decanoate
Flupenthixol Dihydrochloride
Fluperolone Acetate
Fluphenazine Decanoate
Fluphenazine Enanthate
Fluphenazine Hydrochloride
Fluprednidene Acetate
Fluprednisolone
Fluprostenol Sodium Salt
Flurandrenolone
Flurazepam Hydrochloride
Flurazepam Monohydrochloride
Flurbiprofen
Fluspirilene
Folic Acid; but if in preparations for internal use and mdd 200 micrograms Schedule I.
Follicle Stimulating Hormone
Formocortal
Formosulphathiazole
Fosfestrol Tetrasodium
Framycetin Sulphate

Frusemide
Fumagillin
Fumagillin Bicyclohexylamine
Furahadone
Furazolidone
Furethidine
Fusafungine
Fusidic Acid

Gallamine Triethiodide
Gammaglobulin
Gastrozepin
Gelsemine; but if in preparations for internal or external use and ms 0.1% Schedule 1
Gelsemium; but if in preparations for internal use with mld 25 mg of the crude drug and mskd 75 mg of the crude drug Schedule 1
Gentamicin
Gentamicin Sulphate
Gestronol
Gestronol Hexanoate
Glibenclamide
Glibornuride
Gliperide
Gliquidone
Glucagon
Glutethimide
Glycopyrronium Bromide; but if in preparations for internal use with mld 1 mg and mskd 2 mg Schedule 1
Glymidine
Gonadorelin
Griseofulvin; but if in preparations for external use and ms 0.02% Schedule 1
Griseofulvin
Growth Hormone
Guanethidine Monosulphate
Guanoclor Sulphate
Guanoxan Sulphate

Hachimycin
Haldimide
Halofuginone
Haloperidol
Heparin
Heparin Calcium; but if in preparations for external use Schedule 1
Heptabarbitone
Heptaminol Hydrochloride
Hexachlorophane; but if in human preparations for external use and
(a) in soaps
(i) with ms more than 0.1% but not more than 2% Schedule 1; or
(ii) with ms 0.1% General Sale
(b) in preparations in aerosol dispensers with ms 0.1% General sale
(c) in preparations other than soaps or aerosol dispensers
(i) with ms more than 0.1% but not more than 0.75% Schedule 1
(ii) with ms 0.1% General Sale
Hexamine Phenylethicholate
Hexobarbitone
Hexobarbitone Sodium
Hexoestrol
Hexoestrol Dipropionate
L-Histidine Hydrochloride; but if for use as an ingredient in dietary or nutritional products as an amino-acid Schedule 1
Homatropine, but if:
(a) in preparations for internal use with mld 0.15 mg and mdd 0.45 mg Schedule 1
(b) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)
Homatropine Hydrobromide; but if:
(a) in preparations for internal use with mld 0.2 mg and mdd 0.6 mg Schedule 1
(b) in preparations for external use Schedule 1 (except preparations for local opthalmic use)
Homatropine Methylbromide, but if:
(a) in preparations for internal use with mld 2 mg and mdd 6 mg Schedule 1; or
(b) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)
Hydrallazine Hydrochloride
Hydrargaphen; but if in preparations for local application to the skin Schedule 1
Hydrobromic Acid
Hydrochlorothiazide

Hydrocodone
Hydrocortamate Hydrochloride
Hydrocortisone
Hydrocortisone Acetate
Hydrocortisone 17-Butyrate
Hydrocortisone Caprylate
Hydrocortisone Hydrogen Succinate
Hydrocortisone Sodium Phosphate
Hydrocortisone Sodium Succinate
Hydroflumethiazide
Hydrogen Cyanide; but if in preparations for internal or external use and not more than 0.1% Schedule 1
Hydromorphinol
Hydromorphone
Hydroxypethidine
Hydroxychloroquine Sulphate; but if for the prophylaxis of malaria Schedule 1
25-Hydroxy-cholecalciferol
Hydroxymethylgramicidin; but if in throat lozenges or throat pastilles Schedule 1
4-Hydroxy-3-Nitrophenylarsonic Acid
Hydroxyprogesterone
Hydroxyprogesterone Enanthate
Hydroxyprogesterone Hexanoate
Hydroxyurea
Hydroxyzine Embonate
Hydroxyzine Hydrochloride
Hygromycin B.
Hyoscine; but if:
(a) not combined with Atropine or Hyoscyamine or their salts and:
(i) in preparation for internal use and not more than 0.15% Schedule 1; or
(ii) in preparation for external use Schedule 1 (except preparations for local ophthalmic use)
(b) combined with Atropine or Hyoscyamine or their salts and;
(i) in preparation for internal use and not more than 1 mg of the total alkaloids (calculated as base) Schedule 1; or
(ii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)
Hyoscine Butybromide; but if:
(a) not combined with Atropine or Hyoscymine or their salts and;
(i) in preparations for internal use with not more than 5 mg and not more than 9 mg Schedule 1; or
(ii) in inhalers Schedule 1; or
(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)
(b) combined with Atropine or Hyoscyamine or their salts and:
(i) in preparations for internal use and not more than 1 mg of the total alkaloids (calculated as base) Schedule 1; or
(ii) in inhalers Schedule 1; or
(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)
Hyoscine Hydrobromide; but if
(a) not combined with Atropine or hyoscyamine or their salts and:
(i) in preparations for internal use with not more than 300 micrograms and not more than 900 micrograms Schedule 1; or
(ii) in inhalers Schedule 1; or
(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)
(b) combined with Atropine or Hyoscyamine or their salts and:
(i) in preparations for internal use and not more than 1 mg of the total alkaloids (calculated as base) Schedule 1; or
(ii) in inhalers Schedule 1; or
(iii) in preparations for external use Schedule 1 (except preparations for local ophthamic use)
Hyoscine Methobromide; but if:
(a) not combined with Atropine or Hyoscyamine or their salts and:
(i) in preparations for internal use with not more 2.5 mg and not more 7.5 mg Schedule 1; or
(ii) in inhalers Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

(b) combined with Atropine or Hyoscyamine or their salts and:

(i) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(ii) in inhalers Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

Hyoscine Methonitrate; but if:

(a) not combined with Atropine or Hyoscyamine or their salts and;

(i) in preparations for internal use with mdd 2.5 mg and mdd 7.5 mg Schedule 1; or

(ii) in inhalers Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

(b) combined with Atropine or Hyoscymine or their salts and:

(i) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(ii) in inhalers Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

Hyoscyamine; but if:

(a) not combined with Atropine or Hyoscine or their salts and:

(i) inpreparations for internal use with mdd 300 micrograms and madd 1 mg Schedule 1; or

(ii) in inhalers Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

(b) present as an alkaloid of Stramonium in products for the relief of asthma in the form of cigarettes, smoking mixtures or fumigants Schedule 1

(c) combined with Atropine or its salts and;

(i) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(ii) in inhalers Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

Hyoscyamine; but if:

(a) not combined with Atropine or Hyoscine or their salts and;

(i) in preparations for internal use with mdd 300 micrograms (calculated as base) and mdd 1 mg (calculated as base) Schedule 1, or

(ii) in inhalers Schedule 1; or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthlamic use)

(b) present as an alkaloid of Stramonium in products for the relief of asthma in the form of cigarettes, smoking mixtures or fumigants Schedule 1

(c) combined with Atropine or its salts and:

(i) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(ii) in inhalers Schedule 1; or

(iii) in preparations for external use (except preparations for local ophthalmic use)

(d) combined with Hyoscine or its salts and:

(i) in preparations for internal use and mdd 1 mg of the total alkaloids (calculated as base) Schedule 1; or

(ii) in inhalers schedule 1, or

(iii) in preparations for external use Schedule 1 (except preparations for local ophthalmic use)

Hyoscyamine Hydrobromide but if:
(a) not combined with Atropine or Hyoscine or their salts and:
(i) in preparations for internal use with not 300 micrograms (calculated as base) and not 1 mg (calculated as base) Schedule 4; or
(ii) in inhalers Schedule 4; or
(iii) in preparations for external use Schedule 4 (except preparations for local ophthlamic use)
(b) combined with Atropine or its salts and:
(i) in preparations for internal use and not 1 mg of the total alkaloids (calculated as base) Schedule 4; or
(ii) in preparations for external use Schedule 4 (except preparations for local ophthlamic use)
(c) combined with Hyoscine or its salts and:
(i) in preparations for internal use and not 1 mg of the total alkaloids (calculated as base) Schedule 4; or
(ii) in inhalers Schedule 4; or
(iii) in preparations for external use Schedule 4 (except preparations for local ophthalmic use)
Hyoscyamine Sulphate: but if:
(a) not combined with Atropine or Hyoscine or their salts and;
(i) in preparations for internal use with not 200 micrograms (calculated as base) and not 1 mg (calculated as base) Schedule 4; or
(ii) in inhalers Schedule 4; or
(iii) in preparations for external use Schedule 4 (except preparations for local ophthalmic use)
(b) combined with Atropine or its salts and:
(i) in preparations for internal use and not 1 mg of the total alkaloids (calculated as base) Schedule 4; or
(ii) in inhalers Schedule 4; or
(iii) in preparations for external use Schedule 4 (except preparations for local ophthalmic use)
(c) combined with Hyoscine or its salts and:
(i) in preparations for internal use and not 1 mg of the total alkaloids (calculated as base) Schedule 4
(ii) in inhalers Schedule 4; or
(iii) in preparations for external use Schedule 4 (except preparations for local ophthalmic use)

Ibuprofen
Idoxuridine
Ignatius Bean
Ilosfamide
Imipramine
Imipramine Hydrochloride
Imipramine Ion Exchange Resin Bound Salt or Complex
Indapamide Hemihydrate
Indomethacin
Indoprofen
Indonamin Hydrochloride
Ipratropium Bromide
Iprindole Hydrochloride
Isoniazid Phosphate
Isoaminile
Isoaminile Citrate
Isocarboxazid
Isocharine
Isocharine Hydrochloride
Isocharine Mesylate
Isoflurane
Isoniazid
Isoprenaline Hydrochloride
Isoprenaline Sulphate
Isopropamide Iodide: but if in preparations for internal use with not 2.5 mg (calculated as base) and not 5 mg (calculated as base) Schedule 4
Isopyrin

Jaborandi: but if in preparations for external use and:
(a) not more than 0.025% of the

alkaloids in the medicinal product Schedule 1; or
(b) nis 0.625% of the alkaloids in the medicinal product Schedule 1

Kanamycin Sulphates
Ketobemidone
Ketamine Hydrochloride
Ketazolam
Ketaconazole
Ketoprofen
Khellin

Labetolol Hydrochloride
Lactogenic Hormone
Lanatoside C
Lanatoside Complex A, B, and C
Latamoxef
Lead Arsenate
Levallorphan Tartrate
Levodopa
Levomethorphan
Levomoramide
Levophenacylmorphan
Levorphanol
Lidoflazine
Lignocaine, but if:
(a) in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use); or
(b) in preparations for external use and nis 0.6% – Schedule 1 (except preparations for local ophthalmic use)
Lignocaine Hydrochloride; but if
(a) in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use); or
(b) in preparations for external use and nis 0.7% (except preparations for local ophthalmic use)
Lincomycin
Lincomycin Hydrochloride
Liothyronine Sodium
Lippes Loops
Lithium Carbonate; but if in preparations for internal use with: mdd 5 mg (calculated as base) and mdd 15 mg (calculated as base) Schedule 1
Lithium Sulphate; but if in preparations for internal use with mdd 5 mg (calculated as base) and mdd 15 mg (calculated as base) Schedule 1
Lobeline; but if:
(a) in preparations for internal use with md 3 mg and mdd 9 mg Schedule 1; or
(b) in preparations for external use Schedule 1
Lobeline Hydrochloride; but if:
(a) in preparations for internal use with md 3 mg (calculated as base) and mdd 9 mg (calculated as base) Schedule 1; or
(b) in preparations for external use Schedule 1
Lobeline Sulphate; but if:
(a) in preparations for internal use with md 3 mg (calculated as base) and mdd 9 mg (calculated as base Schedule 1; or
(b) in preparations for external use Schedule 1
Loferamine
Lormetine
Loperamide Hydrochloride; but if for treatment of acute diarrhoea Schedule 1
Lorazepam
Luteinising Hormone
Lymecycline
Lynoestrenol
Lypressin

Mafenide
Mafenide Acetate
Mafenide Hydrochloride
Mafenide Propionate; but if in eye drops nis 5% Schedule 1
Magnesium Bromide
Magnesium Fluoride
Mandragora Autumnalis
Mannomustine Hydrochloride
Maprotiline Hydrochloride
Mazindol
Mebanazine
Mebendazole
Mebeverine Hydrochloride; but if in

preparations for internal use with not 100 mg and not 300 mg
Schedule 1
Mebezonium Iodide
Mebhydrolin
Mecamylamine Hydrochloride
Mecillinam
Medrofenoxate Hydrochloride
Medazepam
Medigoxin
Medroxyprogesterone Acetate
Mefenamic Acid
Mefruside
Megestrol
Megestrol Acetate
Melarsonyl Potassium
Melarsoprol
Melengestrol
Melengestrol Acetate
Melphalan
Melphalan Hydrochloride
Menotrophin
Mepenzolate Bromide; but if in preparations for internal use with not 25 mg and not 75 mg
Schedule 1
Mephenesin
Mephenesin Carbamate
Mephentermine
Mepivacaine Hydrochloride; but if in use preparations for non-parenteral use Schedule 1 (except for local ophthalmic use)
Meprobamate
Meptazinol
Mequitazine
Mercaptopurine
Mercuderamide
Mersalyl
Mersalyl Acid
Mesoridazine
Mestanolone
Mesterolone
Mestranol
Metabutethamine Hydrochloride; but if in preparations for non-parenteral use Schedule 1 (except preparations for ophthalmic use)
Metaraminol
Metazocine
Metformin Hydrochloride
Methacycline
Methacycline Citrate
Methadone
Methadyl
Methallenoestril
Methandienone
Methandriol
Methaqualone
Methaxolone
Methdilazine Hydrochloride
Methenolone Acetate
Methenolone Enanthate
Methicillin Sodium
Methimazole
Methinidizate Hydrochloride
Methixene
Methixene Hydrochloride
Methocarbamol
Methohexitone Sodium
Methotrin
Methoserpidine
Methotrexate
Methotrexate Sodium
Methotrimeprazine
Methotrimeprazine Hydrochloride
Methotrimeprazine Maleate
Methoxamine Hydrochloride, but if in nasal sprays or nasal drops not containing liquid paraffin as a vehicle and not 0.25% Schedule 1
Methsuximide
Methyclothiazide
N-Methyl Acetanilide
Methylamphetamine
Methylbenzoquate
Methyldesorphine
Methyldihydromorphine
Methyldihydromorphinone
Methyldopate Hydrochloride
Methylephedrine Hydrochloride; but if in preparations for internal use with not 30 mg not more than 60 mg
Schedule 1;
Methylergometrine Maleate
Methylpentynol
Methylpentynol Carbamate
Methylphenobarbitone
Methylphenidate
Methylprednisolone
Methylprednisolone Acetate
Methylprednisolone Sodium Succinate
Methylsulphonal

Methyltestosterone
Methylthiouracil
Methyprylone
Methysergide Maleate
Metirosine
Metoclopramide Hydrochloride
Metolazone
Metomidate Hydrochloride
Metopon
Metoprolol Tartrate
Metronidazole
Metyrapone
Mexiletine Hydrochloride
Mezlocillin
Mianserin Hydrochloride
Miconazole; but if for external use Schedule 1 (except for vaginal use)
Miconazole Nitrate; but if for external use Schedule 1 (except for vaginal use)
Minocycline
Minocycline Hydrochloride
Mithramycin
Mitobronitol
Mitomycin C
Mitopodozide
Molindone Hydrochloride
Morpheridine
Morphine
Morazone Hydrochloride
Moxalactam
Mustine Hydrochloride
Myrophine

Nadelol
Nalbuphine Oxalate
Nalbuphine
Nalidixic Acid
Nalorphine Hydrobromide
Naloxone Hydrochloride
Nandrolone Decanoate
Nandrolone Laurate
Nandrolone Phenylpropionate
Naphazoline Hydrochloride; but if:
(a) in nasal sprays or nasal drops not containing liquid paraffin as a vehicle and not 0.05% Schedule 1; or
(b) in eye drops and not 0.015% Schedule 1
Naphazoline Nitrate; but if in nasal sprays or nasal drops not containing liquid paraffin as a vehicle and not 0.05% Schedule 1
Naproxen
Naproxen Sodium
Natamycin
Nealbarbitone
Nefopam
Neoarsphenamine
Neomycin
Neomycin Oleate
Neomycin Palmitate
Neomycin Sulphate
Neomycin Undecanoate
Neostigmine Bromide
Neostigmine Methylsulphate
Netilmycin
Niclamide
Nicardipine
Niceocdinate
Nicomorphine
Nicotinaldehyde Thio-Semicarbazone
Nicotinalkone
Nifedipine
Nifenazone
Nikethamide
Niridazole
Nitrazepam
Nitrofurantoin
Nitrofurazone
Nitroxoline
Nomifensine Hydrogen Maleate
Norcyclmethadol
Noradrenaline
Noradrenaline Acid Tartrate
Norcodeine
Norethandrolone
Norethisterone
Norethisterone Acetate
Norethynodrel
Norgestrel
d-Norgestrel
Norlevorphanol
Normethadone
Normorphine
Norpipanone
Nortriptyline Hydrochloride
Novobiocin Calcium
Novobiocin Sodium
Nux Vomica Seed
Nux Vomica Tincture

Nystatin

Octacosactrin
Oestradiol
Oestradiol Benzoate
Oestradiol Cypionate
Oestradiol Dipropionate
Oestradiol Dundecanoate
Oestradiol Enanthate
Oestradiol Phenylpropionate
Oestradiol Undecanoate
Oestradiol Valerate
Oestriol
Oestriol Di-Hemi Succinate
Oestrogenic Substances Conjugated
Oestrone
Ofloxacin
Ofloxacin Hydrochloride
Opium Tincture
Orciprenaline Sulphate
Orphenadrine Citrate
Orphenadrine Hydrochloride
Orthocaine, but if in preparations for non parenteral use Schedule 1 (except preparations for local ophthalmic use)
Ouabain
Ovarian Gland Dried
Oxamniquine
Oxandrolone
Oxantel Pamoate
Oxaprozin
Oxazepam
Oxedrine Tartrate
Oxethazaine
Oxethazine; but if in preparations for non-parenteral use Schedule 1
Oxolinic Acid
Oxophenarsine Hydrochloride
Oxophenarsine Tartrate
Oxpentifylline
Oxprenolol Hydrochloride
Oxybuprocaine Hydrochloride; but if in preparations for non-parenteral use Schedule 1 (except preparations for local Ophthalmic use)
Oxycodone
Oxymestrone
Oxymetholone
Oxymorphone
Oxypertine
Oxyperitn Hydrochloride
Oxyphenabutazone
Oxyphencyclimine Hydrochloride
Oxyphenonium Bromide, but if in preparations for internal use with mid 5 mg and mdd 15 mg Schedule 1
Oxytetracycline
Oxytetracycline Calcium
Oxytetracycline Dihydrate
Oxytetracycline Hydrochloride
Oxytocin, natural
Oxytocin, synthetic

Pancuronium Bromide
Papaverine, but if (i) in inhalers Schedule 1; or
(ii) in preparations for internal use with mid 50 mg and mdd 150 mg Schedule 1
Papverine Hydrochloride; but if (i) in inhalers Schedule 1; or
(ii) in preparations for internal use with mid 50 mg (calculated as base) and mdd 150 mg (calculated as base) Schedule 1
Papaverine Nitrite, but if (i) in in inhalers Schedule 1; or
(ii) in preparations for internal use with mid 50 mg (calculated as base) and mdd 150 mg (calculated as base) Schedule 1
Papaveroline
Papaveroline 2-Sulphonic Acid Paraldehyde
Paramethadione
Paramethasone Acetate
Parathyroid Gland
Pargyline Hydrochloride
Paromomycin Sulphate
Pecilocin
Pemoline
Pempidine Tartrate Penaltero.
Penamecillin
Penbutol
Penethamate Hydriodide
Penicillamine
Penicillamine Hydrochloride
Pentacosactride
Pentazocine Hydrochloride

Pentazocine Lactate
Penthienate Methobromide; but if in preparations for internal use with mid 5 mg and midd 15 mg Schedule 1
Penthobarbitone
Pentobarbitone Sodium
Pentolinium Tartrate
Perhexiline Hydrogen Maleate
Pericyazine
Perifusin
Perphenazine
Pethidine
Phenacaine; but if in Preparations for non-parenteral use Schedule 1 (except preparations for local opthalmic use)
LN 63/1988 Phenacemide
Phenacetin; but if in preparations as a stabiliser and ms 0.1% Schedule 1
Phenadoxone
Phenapromide
Phenarsone Sulphoxylate
Phenazocine
Phenazone; but if in preparations for external use Schedule 1
Phenazone and Caffeine Citrate
Phenazone Salicylate
Phenbenicillin Potassium
Phenbutrazate Hydrochloride
Pheneyclicine
Pheneyclidine Hydrochloride
Phenelzine Sulphate
Phenmetrazine
Phendimetrazine
Phenethicillin Potassium
Phenfluridone
Phenformin Hydrochloride
Phenglutarimide Hydrochloride
Phenidione
Phenobarbitone
Phenobarbitone Sodium
Phenomorphan
Pheneperidine
Phenoxybenzamine Hydrochloride
Phenoxymethylpenicillin
Phenoxymethylpenicillin Calcium
Phenoxymethylpenicillin Potassium
Phenprocoumon
Phensuximide
Phentermine Hydrochloride
Phentermine Resin Complex
Phentolamine Hydrochloride
Phentolamine Mesylate
Phenyl Aminosalicylate
Phenylbutazone
Phenylbutozone Sodium
Phenylmethylbarbituric Acid
Phenylpropanolamine Hydrochloride; but if:
(a) in preparations for internal (except nasal sprays or nasal drops) with mid 50 mg and midd 150 mg Schedule 1; or
(b) in nasal sprays or nasal drops and ms 2% Schedule 1
Phenytoin
Phenytoin Sodium
Pholcodine; but if in Preparations for treatment of cough ms 0.1% w/v Schedule 1
Phthalylsulphacetamide
Phthalylsulphathiazole
Physostigmine
Physostigmine Aminoxide Salicylate
Physostigmine Salicylate
Physostigmine Sulphate
Picrotoxin
Pilocarpine
Pilocarpine Hydrochloride
Pilocarpine Nitrate
Pimindine
Pimozide
Pindol
Pipenzolate Bromide; but if in preparations for internal use with mid 5 mg and midd 15 mg Schedule 1
Piperacillin
Piperazine Oestrone Sulphate
Piperidolate Hydrochloride; but if in preparations for internal use with mid 50 mg and midd 150 mg Schedule 1
Pipothiazine Palmitate
Piracetam
Pirprofen
Pirroxicam
Pituitary Gland (Whole Dried)
Pituitary Powdered (Posterior Lobe)
Pivampicillin Hydrochloride
Pivmecillinam
Pivmecillinam Hydrochloride

Pizotifen
Pizotifen Hydrogen Maleate
Podophyllum
Podophyllum Indian
Podophyllum Resin; but if in preparations for external use and not 20% Schedule 1
Poldine Methylsulphate; but if in preparations for internal use with not 2 mg and not 6 mg Schedule 1
Polidexide
Polidexide Hydrochloride
Polidexide Sulphate
Polymyxin B Sulphate
Polyoestradiol Phosphate
Polythiazide
Poppy Capsule
Potassium Aminosalicylate
Potassium Arsenite; but if in preparations for internal or external use and not 0.0127% Schedule 1
Potassium Bromide
Potassium Canrenoate
Potassium Clorazepate
Potassium Perchlorate
Practolol
Pralidoxime Chloride
Pralidoxime Iodide
Pralidoxime Mesylate
Prazepam
Prazosin Hydrochloride
Prednisolone Prednisolone Acetate
Prednisolone Butylacetate
Prednisolone Hexanoate
Prednisolone Pivalate
Prednisolone Sodium Phosphate
Prednisolone Sodium m Sulphobenzoate
Prednisolone 21-Stearate
Prednisolone m-Sulphobenzoate
Prednisone
Prednisone Acetate
Prenylamine Lactate
Prilocaine Hydrochloride; but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Primidone
Pripadol
Probenecid
Probucol
Procainamide Hydrochloride
Procaine Hydrochloride, but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Procaine Penicillin
Procarbazine Hydrochloride
Prochlorperazine Edisylate
Prochlorperazine Maleate
Prochlorperazine Mesylate
Procyclidine Hydrochloride
Progesterone
Proligestone
Prolintane Hydrochloride
Promazine Embonate
Promazine Hydrochloride
Propandid
Propantheline Bromide; but if in preparations for internal use with not 15 mg and not 45 mg Schedule 1
Propicillin Potassium
Propiram
Propiomazine Hydrogen Maleate
Propranolol Hydrochloride
Propylhexedrine; but if in inhalers Schedule 1
Propylhexedrine Hydrochloride; but if in inhalers Schedule 1
Propylthiouracil
Propyphenazone
Proquamezine Fumarate
Proquazone
Prostaglandin F2 Alpha Trometh amine
Protamine Sulphate
Protriptyline
Protriptyline Hydrochloride
Proxymetacaine Hydrochloride; but if in preparations for non-parenteral use Schedule 1 (except preparations for local ophthalmic use)
Pseudoephedrine Hydrochloride, but if in preparations for internal use with not 60 mg and not 180 mg Schedule 1

Pyrantel Embonate
Pyrantel Tartrate
Pyrazinamide
Pyridostigmine Bromide
Pyrimethamine; but if for (1) Human use Schedule 1
L-Pyroglutamyl-1 Histidyl-l-Proline Amide

Quinalbarbitone
Quinalbarbitone Sodium
Quinestradol
Quinestrol
Quinethazone
Quingestanol
Quinidine
Quinidine Bisulphate
Quinidine Phenylethybarbiturate
Quinidine Polygalacturonate
Quinidine Sulphate
Quinine; but if:
(a) in preparations for internal use with not 100 mg and not 300 mg Schedule 1; or
(b) in preparations for internal use and not 35 mg Schedule 1
Quinine Bisulphate; but if:
(a) in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1; or
(b) in preparations for internal use and not 35 mg (calculated as base) Schedule 1
Quinine Dihydrochloride; but if in preparations for internal use with not 100 mg (Calculated as base) and not 300 mg (calculated as base) Schedule 1
Quinine Ethyl Carbonate, but if in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1
Quinine Glycerophosphate; but if in preparations for internal use with not 100 mg (Calculated as base) and not 300 mg (calculated as base) Schedule 1
Quinine Hydrobromide; but if in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1
Quinine Hydrochloride; but if:
(a) in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1; or
(b) in preparations for internal use and not 35 mg (calculated as base) Schedule 1
Quinine Iodobismuthate, but if in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1
Quinine Phenylcinchoninate, but if in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1
Quinine Phosphate; but if in preparations for internal use with not 100 mg (Calculated as base) and not 300 mg (calculated as base) Schedule 1
Quinine Salicylate; but if:
(a) in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1; or
(b) in preparations for internal use and not 35 mg (calculated as base) Schedule
Quinine Sulphate; but if:
(a) in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1; or
(b) in preparations for internal use and not 35 mg (calculated as base) Schedule 1
Quinine Tannate; but if in preparations for internal use with not 100 mg (calculated as base) and not 300 mg (calculated as base) Schedule 1
Quinine and Urea Hydrochloride
Quinuretium Sulphate

Racephedrine hydrochloride: but it:
(a) in preparations for internal use (except nasal sprays or nasal drops) with not 30 mg and not 60 mg Schedule 1; or
(b) in nasal sprays or nasal drops and not 2% Schedule 1; or
(c) in preparations for external use Schedule 1
Racemethorphan
Racemoramide
Racemorphan
Ragwort: but if in preparations for external use and not 10% of the crude drug Schedule 1
Ranitidine
Rauwolfia (Serpentina and Vomitoria)
Razoxane
Reproterol Hydrochloride
Resennamine
Reserpine
Rifamide
Rifampicin
Rifamycin
Rimiterol Hydrobromide
Ritodrine Hydrochloride
Rolitetracycline Nitrate
Ronidazole

Sabadilla
Salazosulphadimidine
Salbutamol
Salbutamol Sulphate
Salcatonin
Salcatonium Hydrated Polyacetate
Salmefamol
Salsalate
Secbutobarbitone
Secbutibarmitone Sodium
Sera and Antisera:
(1) Human:
Botulin Antitoxin
Gas gangrene Antitoxin (Oedematiens)
Gas gangrene Antitoxin (Perfringens)
Gas-gangrene Antitoxin (Septicum)
Leptospira Antiserum
Mixed Gas-gangrene
Rabies Antiserum
Scorpion Venom Antiserum
Snake Venom Antiserum
Tetanus Antitoxin
Serum Gonadotrophin
Silver Sulphadiazine
Sissomycin
Sodium Aminosalicylate
Sodium Antimony Gluconate
Sodium Apolate; but if preparations for external use Schedule 1
Sodium Arsanilate
Sodium Arsenate
Sodium Arsenite; but if in preparations for internal or external use and not 0.012% Schedule 1
Sodium Bromate
Sodium Bromide
Sodium Cacodylate
Sodium Cromoglycate; but if in preparations for human use by being administered through the nose Schedule 1
Sodium Ethacrynate
Sodium Fluoride: but if in
(a) dentifrices and not 0.33% Schedule 1
(b) other preparations for use in the prevention of dental caries in the form of:
(i) tablets or drops and not 2.2 mg Schedule 1
(ii) mouth rinses other than those for daily use and not 0.2% Schedule 1
(iii) mouth rinses for daily use and not 0.05% Schedule 1
Sodium Fusidate
Sodium Methylarsinate
Sodium Monofluorophosphate, but if in dentifrices and not 1.14% Schedule 1
Sodium Stibogluconate
Sodium Valproate
Solapsone
Sotalol Hydrochloride
Spectinomycin
Spiramycin
Spiramycin Adipate
Spironolactone
Stannous Fluoride: but if in dentifrices and not 0.62% Schedule 1

Stanolone
Stanozolol
Stibocaptate
Stibophen
Stilboestrol
Stilboestrol Dipropionate
Streptodornase; but if in preparations for external use Schedule 1
Streptokinase; but if in preparations for external use Schedule 1
Streptomycin
Streptomycin Sulphate
Strontium Bromide
Strophanthin K
Strychnine
Strychnine Arsenate
Strychnine Hydrochloride
Styramate
Succinylsulphathiazole
Sucralfate
Sufentanil
Sulfabenz
Sulfacytine
Sulfadiazamide
Sulfadoxine
Sulfamethpyrazine
Sulfamonomethoxine
Sulfapyrazole
Sulindac
Sulphabromomethazine
Sulphacetamide
Sulphacetamide Sodium
Sulphachlorpyridazine
Sulphadiazine
Sulphadiazine Sodium
Sulphadimethoxine
Sulphadimidine
Sulphadimidine Sodium
Sulphaethidole
Sulphafurazole
Sulphafurazole Diethanolamine
Sulphaguanidine
Sulphalexic Acid
Sulphamerazine
Sulphamerazine Sodium
Sulphamethizole
Sulphamethoxazole
Sulphamethoxydiazine
Sulphamethoxypyridazine
Sulphamethoxypyrazine Sodium
Sulphamethylphenazole
Sulphamoxole
Sulphanilamide
Sulphanitran
Sulphaphenazole
Sulphapyridine
Sulphapyridine Sodium
Sulphaquinoxaline
Sulphaquinoxaline Sodium
Sulpharsphenamine
Sulphasalazine
Sulphasomidine
Sulphasomidine Sodium
Sulphathiazole
Sulphathiazole Sodium
Sulphathiourea
Sulphatolamide
Sulphaurea
Sulphinpyrazone
Sulphomyxin Sodium
Sulphonal
Sulpiride
Sulthiame
Suxamethonium Bromide
Suxamethonium Chloride
Suxe hominum Bromide

Taenia Hydrochloride
Talampicillin
Talampicillin Hydrochloride
Talampicillin Napsylate
Tamoxifen
Tamoxifen Citrate
Teclothiazide Potassium
Temazepam
Terbutaline
Terbutaline Sulphate
Testosterone
Testosterone Acetate
Testosterone 17B Chloral Hemiacetal
Testosterone Cyclohexylpropionate
Testosterone Cypionate
Testosterone Decanoate
Testosterone Enanthate
Testosterone Isocaproate
Testosterone Phenylpropionate
Testosterone Propionate
Tetrabenazine
Tetracosactrin
Tetracosactrin Acetate
Tetracycline
Tetracycline Hydrochloride

Tetracycline Phosphate Complex
Thallium Acetate
Thallous Chloride
Theobromine
Theobromine
Thiamylal
Thiamylal Sodium
Thiamylal Sodium
Thiethylperazine
Thiethylperazine Di-(Hydrogen Malate)
Thiocarlide
Thioguanine
Thiopentone Sodium
Thiopropazate Hydrochloride
Thiopropazate Mesylate
Thioridazine
Thioridazine Hydrochloride
Thiosinamine and Ethyl Iodide
Thiostrepton
Thiotepa
Thiothixene
Thiouracil
Thymoxamine Hydrochloride
Thymol
Thyrotrophin
Thyrotrophin Releasing Hormone
Thyroxine Sodium
Timolin Hydrogen Furarate
Ticarcillin
Tiglotidine Hydrobromide
Tilidate
Timolol Maleate
Tinidazole
Tobramycin Sulphate
Tofenacin Hydrochloride
Tolazamide
Tolazoline Hydrochloride
Tolbutamide
Tolbutamide Sodium
Tolmetin Sodium Hydrate
Tolperisone
Totaquine
Tranexamic Acid
Tranylcypramine sulphate
Trazodone
Treosulphan
Tretamine
Trethonin
Triacetyloleandomycin
Triamcinolone
Triamcinolone Acetonide
Triamcinolone Diacetate
Triamcinolone Hexacetonide
Triamterene
Triaziquone
Triazolam
Tribromoethyl Alcohol
Trichlofos Sodium
Tricyclamol Chloride
Trienbolone Acetate
Trifluoperazine
Trifluoperazine Hydrochloride
Trifluperidol
Trilostane
Trimeperidine
Trimeprazine
Trimeprazine Tartrate
Trimetaphan Camsylate
Trimetazidine
Trimetazidine Hydrochloride
Trimethoprim
Trimipramine Maleate
Trimipramine Mesylate
Trimustine Hydrochloride
Trispiramide
Troxidone
L-Tryptophan: but it
(a) for external use Schedule 3; or
(b) used as an ingredient in dietary or nutritional products as an essential amino-acid Schedule 1
Tubocurarine Chloride
Tybamate
Tylosin
Tylosin Phosphate
Tylosin Tartrate
Tyrothricin: but it in throat lozenges or throat pastilles Schedule 1

Uramustine
Urea Stibamine
Uredofos
Urethane
Uridine-5-Triphosphoric Acid
Urokinase
Ursodeoxycholic Acid
Vaccines:
(1) Human: Bacillus Calmette-Guerin Vaccine Percutaneous Bacillus Calmette-Guerin Vaccine
Cholera Vaccine
Diphtheria Vaccine

Adsorbed Diphtheria Vaccine
Diphtheria and Tetanus Vaccine
Adsorbed Diphtheria and Tetanus Vaccine
Diphtheria, Tetanus and Pertussis Vaccine
Adsorbed Diphtheria, Tetanus and Pertussis Vaccine
Ebtor Vaccine
Diphtheria, Tetanus and Poliomyelitis Vaccine
Diphtheria, Tetanus, Pertussis and Poliomyelitis Vaccine
Influenza Vaccine
Measles Vaccine (Live Attenuated)
Pertusis Vaccine
Plague Vaccine
Poliomyelitis Vaccine (Inactivated)
Poliomyelitis Vaccine (Oral)
Rabies Vaccine
Rubella Vaccine (Live Attenuated)
Schick Control
Schick Test Toxin
Smallpox Vaccine
Dried Smallpox Vaccine
Tetanus Vaccine
Adsorbed Tetanus Vaccine
Tetanus and Pertussis Vaccine
Tuberculin Purified Protein Derivative
Old Tuberculin
Typhoid Vaccine
Typhoid and Tetanus Vaccine
Typhoid paratyphoid A and B Vaccine
Typhoid-paratyphoid A and B and Cholera Vaccine
Typhoid-paratyphoid A and B and Tetanus Vaccine
Typhus Vaccine
Yellow Fever Vaccine

Valproic Acid
Vancomycin Hydrochloride
Vasopressin Injection
Vasopressin Tannate
Verapamil Hydrochloride
Veratrine
Veratrum (Green and White)
Vidarabine
Viloxazine Hydrochloride
Vinbarbitone
Vinbarbitone Sodium
Vinblastine Sulphate
Vincristine Sulphate
Vindesine
Viomycin Pantothenate
Viomycin Sulphate
Virginiamycin

Warfarin
Warfarin Sodium

Xipamide
Xylazine Hydrochloride

Yohimbine Hydrochloride

Zeranol
Zimeldine
Zotepirac

FIFTH SCHEDULE

FORMS TO WHICH THE SUBSTANCES SPECIFIED ARE RESTRICTED WHEN SOLD BY LICENSED SELLERS OF PART II POISONS—RULE 13 (2)

Poison	Form to which sale is restricted
Arsenical substances—	
Arsenious oxide.	Sheep dips, sheep washes.
Arsenic sulphides.	" "
Calcium arsenates.	Agricultural and horticultural insecticides or fungicides.
Calcium arsenites.	" " " "
Copper acetoarsenites.	" " " "
Copper arsenates.	" " " "
Copper arsenites.	" " " "
Lead arsenates.	" " " "
Potassium arsenites.	Sheep dips sheep washes.
Sodium arsenates.	" "
Sodium arsenites.	" "
Sodium theoarsenates.	" "
Barium carbonate.	Preparations for the destruction of rats and mice.
Mercurial substances—	

Mercuric chloride.

Mercuric iodide.

Alkaloids

Aconite, alkaloids of.	The proportion of any one alkaloic of aconite that the preparation would be calculated to contain on the assumption that all the alkaloids of aconite in the preparation were that alkaloid.
Belladonna, alkaloids of.	The same as above with the substitution for the reference to aconite, of a reference to belladonna, calabar bean, or such other of the said poisons as the case may require.
Calabar Bean, alkaloids of.	
Coca, alkaloids of	
Ephedra, alkaloids of.	
Ergot, alkaloids of.	

Agricultural and horticultural fungicides, seed and bulb dressings, insecticides.	
Agricultural and horticultural fungicides, seed and bulb dressings.	
Organic compounds of mercury.	" " " "
Nitrobenzene.	Agricultural and horticultural insecticides, substances for the treatment of bee disease.

SIXTH SCHEDULE

STATEMENT OF PARTICULARS AS TO PROPORTIONS OF THE POISON IN CERTAIN CASES PERMITTED BY RULE 16 (2)

Name of poison	Particulars

Name of poison	*Particulars*
Gelsemium, alkaloids of. Jaborandi, alkaloids of. Lobelia, alkaloids of. Pomegranate, alkaloids of. Quebracho, alkaloids of, other than the alkaloids of red quebracho. Sabadilla, alkaloids of. Solanaceous alkaloids not otherwise included in the Poisons List. Staves acre, alkaloids of. Veratrum, alkaloids of. Yohimba, alkaloids of.	
Antimonial poisons.	The proportion of antimony trioxide (Sb_2O_3) or antimony pentoxide (Sb_2O_5) that the preparation would be calculated to contain on the assumption that the antimony (Sb) in the poison has been wholly converted into antimony trioxide or antimony pentoxide as the case may be.
Arsenical poisons.	The proportion of arsenic trioxide (As_2O_3) or arsenic pentoxide (As_2O_5) that the preparation would be calculated to contain on the assumption that all the arsenic (As) had been wholly converted into arsenic trioxide or arsenic pentoxide as the case may be.
Barium, salts of.	The proportion of one particular Barium salt that the preparation would be calculated to contain on the assumption that all the barium (Ba) in the poison had been wholly converted into that salt
Digitalis, glycosides of, other active principles of digitalis.	The number of units of activity as defined in the British Pharmacopoeia contained in a specified quantity of the preparation
Hydrocyanic acid, cyanides, double cyanides of mercury and zinc.	The proportion of hydrocyanic acid (HCN) that the preparation would be calculated to contain on the assumption that the cyanides in the poison had been wholly converted into hydrocyanic acid.
Lead, compound of, with acids from fixed oils.	The proportion of lead oxide (PbO) that the preparation would be calculated to contain on the assumption that all the lead in the poison has been wholly converted into lead oxide.

Particulars

The proportion of organically combined mercury (Hg) contained in the preparation.

The proportion of phenols (added together) contained in the preparation.

The proportion of phenols (added together) that the preparation would be calculated to contain on the assumption that the compounds of phenol with a metal had been wholly converted into the corresponding phenol.

Either—

(a) the number of units of activity as defined in the British Pharmacopoeia contained in a specified quantity of the preparation, or

(b) the proportion of pituitary gland or of anterior or posterior lobe of the gland as the case may be contained in the preparation; or

(c) the amount of pituitary gland or of anterior or posterior lobe of the gland as the case may be, from which a specified quantity of the preparation was obtained together with an indication whether the amount relates to fresh or dried gland substance.

The proportion of potassium monoxide (K_2O) that the preparation would be calculated to contain on the assumption that all the potassium hydroxide in the preparation had been wholly converted into potassium monoxide.

The proportion of sodium monoxide (Na_2O) that the preparation would be calculated to contain on the assumption that all the sodium hydroxide in the preparation had been wholly converted into sodium monoxide.

The amount of Standard Tincture of Strophanthus as defined in the British Pharmacopoeia which possesses the same activity as a specified quantity of the preparation when assayed by the same method as described in the said Pharmacopoeia.

Either—

(a) the proportion of suprarenal gland or of the cortex or of the medulla of the gland as the case may be, contained in the preparation; or

Name of poison	*Particulars*
	(b) the amount of suprarenal gland or of the cortex or of the medulla of the gland as the case may be, from which a specified quantity of the preparation was obtained, together with an indication whether the amount relates to fresh or dried gland substance.
Thyroid gland, the active principles of, their salts.	Either— (a) the proportion of thyroid gland contained in the preparation; or (b) the amount of thyroid gland from which a specified amount of the preparation was obtained together with an indication whether the amount relates to fresh or dried gland.

SEVENTH SCHEDULE

INDICATION OF CHARACTER PRESCRIBED BY RULE 17(1) FOR THE PURPOSE OF SECTION 56(1) (c) (iii) of the Act LN 61/1981

1. To be labelled with the words "Caution. It is dangerous to take this preparation except under medical supervision."

Medicines made up ready for the treatment of human ailments if the poison is one of the following—

Allylisopropylacetylurea.

Insulin.

Phenylethylhydantoin; its salts; its acyl derivatives; their salts.

Pituitary gland, the active principles of.

Thyroid gland, the active principles of; their salts.

2. To be labelled with the words "Caution. It is dangerous to exceed the stated dose."

Medicines (other than medicines mentioned in paragraph 1 of this Schedule)

3. To be labelled with the words "Poison. For animal treatment only."

Medicines made up ready for the treatment of animals.

4. To be labelled with the words "Caution. This preparation may cause serious inflammation of the skin in certain persons and should only be used in accordance with expert advice."

Preparations for the dyeing of hair containing phenylene diamines or toluene diamines or their salts.

5. To be labelled with the words "Caution. This substance is caustic."

Potassium hydroxide, sodium hydroxide and articles containing either of these substances.

LN 61/1981

6. To be labelled with the words "Poison. Highly dangerous."

Fluorocetamide

Fluorocetanilide

Monoflourocetic acid or its salt

Sodium monofluoroacetate syn: Sodium monofluorocetic acid; commonly known as compound 1080

EIGHTH SCHEDULE

POISON TO WHICH RULE 23 (TRANSPORT) APPLIES

Arsenical poisons.

Barium, salts of.

Hydrocyanic acid; cyanides.

Nicotine.

Strychnine.

Thallium, salts of.

NINTH SCHEDULE

FORM OF APPLICATION TO BE FILLED BY A LICENSED STOREKEEPER FOR A LICENCE TO SELL PART II POISONS (SECTION 54 of the Act).

Pharmacy and Poisons Act.

APPLICATION FOR A LICENCE TO SELL PART II POISONS.

I, being a licensed storekeeper carrying on business at hereby apply for a licence to sell such Part II poisons as may be permitted by the Pharmacy and Poisons Act, Part VI, and the Poisons Rules.

The application refers only to the premises situated at the above address.

I hereby nominate to act as my deputy (deputies) for the sale of poisons in accordance with Rule 13 (1) of the Poisons Rules.

Date: Signature:

I hereby certify that to the best of my knowledge and belief the applicant of is of good character and is a fit and proper person to be a Licensed Seller of Part II Poisons.

Police Officer.

Rank:

Police Station:

Date:

TENTH SCHEDULE

(Rule 26)

FORM OF LICENCE TO BE ISSUED TO A LICENSED STOREKEEPER LICENSING HIM TO SELL CERTAIN PART II POISONS

Pharmacy and Poisons Act

LICENCE

This Licence cannot be transferred and is available for one place of business only.

To [name] [address] you are hereby licensed to sell such Part II Poisons as are enumerated at the foot hereof.

Your particular attention is drawn to the Poisons Rules, especially those portions dealing with the sale, supply, storage, labelling and transport of poisons.

By order of the Pharmacy and Poisons Board.

Secretary.

ELEVENTH SCHEDULE

LN 68/1987

(Rule 27)

SCALE OF FEES TO BE PAID IN RESPECT OF REGISTRATION, LICENCE, ETC.

Licence to sell PART II poison.....$ 15.00 per year

Licence to sell methylated spirits.....$ 7.50 per year

Licence to sell medicens.....$ 30 per year

Licence to sell medicines & methylated spirits.....$ 37.50 per year

Licence to sell medicines & Part II Poisons.....$ 45.00 per year

Import Licence (for each item & on every

occasion of importation).....$ 5.00

Registration as a Pharmacist.....$ 20.00 per year

Registration of premises.....$ 50.00 per year

TWELFTH SCHEDULE

(Rule 28)

FORM OF CERTIFICATE OF REGISTRATION AS PHARMACIST SOLOMON ISLANDS

Pharmacy and Poisons Act

This is to certify that of who has duly passed the Qualifying Examination for Pharmacists of has been registered as a Pharmacist and is an authorised seller of poisons under the Pharmacy and Poisons Act.

This certificate expires on the 31st December, 19.

Registrar, Pharmacy and Poisons Board.

Date:

THIRTEENTH SCHEDULE

(Rule 29)

FORM OF REGISTER OF PHARMACISTS
REGISTER OF PHARMACISTS

Year:

		Date Name	Address qualified		

Where qualified

Date registered

Signature of Registrar

FOURTEENTH SCHEDULE

(Rule 30)

FORM OF REGISTER OF PREMISES

REGISTER OF PERSONS ENTITLED TO SELL DRUGS, MEDICINES, and Poisons.

Licence No.	Name

Address of premises

Class of business

Name of Deputy or Deputies permitted to sell

FIFTEENTH SCHEDULE

(Rule 31)

FORM OF APPLICATION FOR LICENCE TO SELL MEDICINES

Pharmacy and Poisons Act

To the Chairman,

Pharmacy and Poisons Board.

I, of hereby apply for a licence to sell medicines.

My place of business is situated miles from the nearest registered pharmacist at

Date: Signature of applicant

I hereby certify that to the best of my knowledge and belief the distance of the applicant's place of business is situated the stated distance from the nearest pharmacists' business at and that the applicant is a fit and proper person to sell medicines.

Police Officer

Rank:

Police Station:

Date:

(Rule 32)

FORM OF LICENCE TO SELL MEDICINES.

Solomon Islands Pharmacy and Poisons Act

LICENCE TO SELL MEDICINES

/name/ address/ is hereby licensed to sell at/premises/ the medicines specified in the First Schedule below subject to the conditions specified in the Second Schedule below.

This licence is not transferable and is available for only the place of business specified above.

This licence is valid for a period of one year and shall expire on the day of ,19

FIRST SCHEDULE

Medicines authorised to be sold

SECOND SCHEDULE

Conditions

By order of the Pharmacy and Poisons Board.

Secretary.

SEVENTEENTH SCHEDULE

(Rule 33)

CERTIFICATE FOR THE PURCHASE OF A POISON

For the purposes of subsection (2) (a)(i) of section 56 of the Pharmacy and Poisons Act, I, the undersigned, a house-holder of (a) from my knowledge of (b) of (a) that he is a person to whom (c) may properly be supplied.

I hereby certify that (d) is the signature of the said (b).

Signature of householder giving certificate.

Date:

(a) Insert full postal address.

(b) Insert full name of intending purchaser.

(c) Insert name of poison.

(d) Intending purchaser to sign his name here.

Endorsement required when the householder giving the certificate is not known to the seller of the poison, as a responsible person of good character.

I hereby certify that in so far as is known to the police of the province in which resides he is a responsible person of good character.

Police Officer

Rank:

Police Station:

Date:

EIGHTEENTH SCHEDULE

FORM OF ENTRY TO BE MADE UNDER RULE 34 IN THE BOOK TO BE KEPT BY SELLERS OF POISONS IN ACCORDANCE WITH SECTION 56 (2) (B) OF THE ACT

THE POISONS (AGRICULTURAL AND SILVICULTURAL USE OF ARSENICAL POISONS) RULES

(Section 61)

[1st March 1964]

Rules by the Pharmacy and Poisons Board

1. These Rules may be cited as the Poisons (Agricultural and Silvicultural Use of Arsenical Poisons) Rules.

2. In these Rules, except where the context otherwise requires—

Title

Interpretation

"arsenical poisons" means any substance containing any form of arsenic and prepared, intended or likely to be used for agricultural or silvicultural purposes;

"inspector" means an inspector appointed under section 14 (2) of the Act; LN 69/1964

"licensing officer" means the Director of Agriculture and includes a licensing officer appointed under rule 3 (1).

3.—(1) The Director of Agriculture may, for the purposes of these Rules, in writing appoint any person to be a licensing officer either generally or for such areas as he shall specify.

(2) On application therefor in Form A in the Schedule to these Rules, a licensing officer may, in his discretion, issue to any person, subject to such

Appointment and powers of licensing officers

Form A, Schedule Form B, Schedule

4.—(1) No person shall purchase or otherwise acquire any arsenical poison except under and in accordance with the terms and conditions of a written permit first had and obtained under rule 3 (2).

Prohibition of sale and purchase of arsenical poisons without a permit

(2) No person shall sell or otherwise supply any arsenical poison to any other person save a person who is authorised so to acquire that poison by a permit issued under rule 3 (2).

5.—(1) No person shall transport, or cause to be transported, any arsenical poison in any compartment of any vessel, aircraft or vehicle in which food, drink, medicine, or any other thing for human or animal consumption is being transported:

Provided that nothing in this paragraph shall apply to any arsenical poison imported into Solomon Islands until such arsenical poison has been discharged from the vessel or aircraft in which it has been transported to Solomon Islands.

(2) Where any compartment of any vessel, aircraft or vehicle that has been used to transport any arsenical poison retains any trace of such poison, no person shall use, cause or permit that compartment to be used for any other purpose until it has been so treated as to remove all traces of the poison.

6. Any person who has in his possession or under his control any arsenical poison shall comply with the following provisions—

(a) arsenical poisons shall be stored in a secure locked room which shall be at a distance of not less than 24 feet from any room or building of whatever sort containing food, drink, medicine or other thing for human or animal consumption;

(b) no room shall be used for the storage of arsenical poisons unless it is constructed of concrete blocks, wooden planking, galvanised iron, fibre board or similar durable materials;

(c) the store shall only be opened in the presence of the person who has possession or control of the arsenical poison or by such other responsible person as may be deputed by him in writing to be in

(d) every container of arsenical poisons shall be painted with a blue band, centrally placed and not less than 18 inches wide, with a white band, not less than 6 inches wide, in the centre of the blue band and the word "POISON" shall be painted in the white band in blue letters not less than 2 inches high:

Provided that, if the container is less than 18 inches high, the whole container shall be painted blue with a white band, centrally placed, not less than one-third of the height of the container within which shall be painted the word "POISON" in blue letters the height of which shall be not less than one-third of the height of the white band;

(e) the store shall be thoroughly washed and cleaned before being used for any other purpose.

7. Every person who uses or causes or permits to be used any arsenical poison shall comply with the following provisions—

(a) The following records shall be maintained and produced to an inspector on demand, that is to say—

(i) the date, quantity and description of arsenical poisons acquired;

(ii) the name and address of the vendor;

(iii) the quantity issued for use on each day;

(iv) the method of use;

(v) the dilution proportions used;

(vi) the area treated; and

(vii) the manner and place of disposal of any arsenical solution remaining at the end of each day's operations;

(b) apparatus and equipment for the use of arsenical poisons

(i) be painted blue with a white band or patch on which the word "POISON" in blue letters shall be painted as clearly and prominently as practicable;

(ii) when not in use, be kept in a secure locked store; and

(iii) be destroyed or thoroughly cleaned and washed when no longer required;

(c) there shall be displayed in conspicuous places at reasonable intervals on the boundaries of areas which have been treated with arsenical poisons notices with a warning that the areas within the boundaries have been treated with arsenical poisons. Such notices shall be in the English language and shall not be removed until the area treated has been cleansed by a heavy rainfall;

(d) roads, paths and rights of way, or the verges thereof, and the banks of rivers, streams or other water courses and any area where there is a risk of contaminating water supplies used for human consumption shall not be treated with arsenical poisons.

8.—(1) No person shall employ, engage or direct any other person (hereafter in this rule referred to as an employee) to perform any work involving exposure to or use of arsenical poisons unless he complies, and ensures the compliance of his employees, with each of the following provisions, that is to say—

(a) every employee shall be provided with, and when exposed to or engaged in the use of arsenical poisons, shall use long sleeved dungarees, rubber gloves and boots and barrier cream;

(b) adequate facilities to the satisfaction of an inspector shall be provided to enable every employee using, or exposed to, arsenical poisons to wash his body and clothing daily, and every such employee shall daily wash his body and clothing as soon as practicable after the completion and termination of the day's use of or exposure to arsenical poisons;

(c) every employee, being before exposed to or permitted to engage in the use of arsenical poisons, shall be warned of the dangers of contact with such poisons and shall be examined by a

(d) no employee shall be exposed to or permitted to engage in the use of arsenical poisons while suffering from any cut, wound or abrasion;

(e) every person exposed to or engaged in the use of arsenical poisons shall be examined by a medical practitioner registered under the Medical and Dental Practitioners Act or a person registered under the Nursing Council Act, at intervals of not more than one week and a record of all such examinations shall be entered in a book kept for the purpose;

(f) there shall at all times be available for immediate use such equipment and drugs for the treatment of arsenical poisoning as the Under Secretary (Health), Ministry of Health and Medical Services may from time to time approve;

(g) all employees shall be warned of the danger of walking in areas which have been treated with arsenical poisons and of the danger of contact with arsenical poisons:

Provided that no person shall be responsible for his employee's failure to comply with any of the provisions of this rule if he proves that he took all reasonable steps to procure the employee's compliance with such provisions.

(2) No person shall against his will be required to perform any work involving exposure to or use of arsenical poisons.

Construction of Rules

9. These Rules shall be in addition to and not in derogation of the Poisons Rules.

SCHEDULE

FORM A

(Rule 3)

I (name).....
of (address).....occupation).....
apply for a permit to obtain arsenical poisons for the following agricultural/ silvicultural purposes—

at (state place or area).....
I certify that I am able to comply with the provisions of the Poisons (Agricultural and Silvicultural Use of Arsenical Poisons) Rules.

.....

(date)

.....

(signature)

FORM B

(Rule 3)

PERMIT TO PURCHASE OR ACQUIRE ARSENICAL POISONS

Permission is hereby granted to (name) of (address) to obtain arsenical poisons for agricultural/silvicultural purposes to be used at (place or area) Special terms and conditions.

1.

2.

3.

.....

(date)

.....

(Licensing Officer)

Note.—Persons wishing to import arsenical poisons should note that a licence to import under section 52 (1) of the Pharmacy and Poisons Act must be obtained from the Pharmacy and Poisons Board to whom this permit should be produced.

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