

Uganda

National Drug Policy and Authority Act Chapter 206

Legislation as at 15 February 2016

FRBR URI: /akn/ug/act/statute/1993/13/eng@2016-02-15

There may have been updates since this file was created.

PDF created on 21 February 2024 at 19:15.

Collection last checked for updates: 15 February 2016.

[Check for updates](#)



About this collection

The legislation in this collection has been reproduced as it was originally printed in the Government Gazette, with improved formatting and with minor typographical errors corrected. All amendments have been applied directly to the text and annotated. A scan of the original gazette of each piece of legislation (including amendments) is available for reference.

This is a free download from the Laws.Africa Legislation Commons, a collection of African legislation that is digitised by Laws.Africa and made available for free.

www.laws.africa
info@laws.africa

There is no copyright on the legislative content of this document.
This PDF copy is licensed under a Creative Commons Attribution 4.0 License (CC BY 4.0). Share widely and freely.

National Drug Policy and Authority Act
 Contents

Part I – Interpretation 1

 1. Interpretation 1

Part II – National drug policy and national drug authority 2

 2. National drug policy 2

 3. Establishment of the National Drug Authority 3

 4. Application of the seal 4

 5. Functions of the drug authority 4

 6. Commission and other bodies of the authority 4

 7. Meetings of the drug authority 6

Part III – Control of the drug supply 6

 8. National list of essential drugs 6

 9. Selection of drug items 6

 10. Estimation of drug needs 6

 11. Drug nomenclature 6

 12. Restricted drugs 6

 13. Supply and dispensing of restricted drugs 7

 14. Licensed persons 8

 15. Licensed sellers 8

 16. Places from which restricted drugs may be supplied 8

 17. Certificates of suitability of premises 9

 18. Loss of class A or B drugs 9

Part IV – Special provisions relating to classified drugs 9

 19. Classified drugs 9

 20. Need for prescription for classified drugs 9

 21. Action to be taken in relation to prescription 10

 22. Classified drugs to be supplied to responsible persons 10

 23. Supply to conform to prescription 10

 24. Classified Drugs Book 10

 25. Containers and labels 11

 26. *** 11

 27. Possession of classified drugs 11

 28. Withdrawal of authority 12

 29. *** 12

 30. Impure drugs not to be supplied 12

31. Power to call for information as to proprietary drugs	12
32. Power to prohibit retail sale of proprietary drugs	13
33. Control of publication of descriptive matter	13
34. Return of details of pharmacy business	13
35. Drug regulation and registration of specialities	13
36. Drug quality	14
37. Licence required for wholesale supply of restricted drugs	14
38. Restrictions on manufacture of classified drugs	15
39. Further restrictions on the manufacture of drugs	15
40. Clinical trials	15
41. Local research and production	15
42. Storage	15
Part V – Control of transport, import and export of drugs	16
43. Transportation of drugs	16
44. Importation of pharmaceuticals	16
45. Exportation of drugs	16
46. Import and export licences	16
Part VI – Further restrictions on narcotics	16
47. ***	16
48. ***	16
49. ***	17
Part VII – Powers of entry and investigation	17
50. Powers of entry	17
51. Powers of investigation	17
52. Authority to be shown	18
53. Obstruction	18
Part VIII – The secretariat and financial provisions	18
54. Secretariat	18
55. Funds of the drug authority	18
56. Estimates	18
57. Accounts	19
58. Audits	19
Part IX – Miscellaneous provisions	19
59. Rational use of drugs	19
60. Offences and penalties	19

61. Vicarious criminal responsibility	20
62. Evidence	20
63. Drugs bureau	21
64. Regulations	21
65. Amendment of Schedules	22
First Schedule (Sections 12, 29)	22
Second Schedule (Sections 12, 29)	25
Third Schedule (Section 12)	38
Fourth Schedule (Section 12)	40
Fifth Schedule (Section 33)	41
Sixth Schedule (Section 38)	43
Seventh Schedule (Section 42)	43
Eighth Schedule (Section 43)	43

Uganda

National Drug Policy and Authority Act Chapter 206

Commenced on 3 December 1993

[This is the version of this document from 15 February 2016.]

[Note: The version of the Act as at 31 December 2000 was revised and consolidated by the Law Reform Commission of Uganda. All subsequent amendments have been researched and applied by Laws.Africa for ULII.]

[Amended by [Narcotic Drugs and Psychotropic Substances \(Control\) Act, 2016 \(Act 3 of 2016\)](#) on 15 February 2016]

An Act to establish a national drug policy and a national drug authority to ensure the availability, at all times, of essential, efficacious and cost-effective drugs to the entire population of Uganda, as a means of providing satisfactory health care and safeguarding the appropriate use of drugs.

Part I – Interpretation

1. Interpretation

In this Act, unless the context otherwise requires—

- (a) “**advertisement**” includes any notice, circular, label, wrapper or other document, and any announcement made orally or by means of producing or transmitting light or sound;
- (b) “**approved institution**” includes gazetted hospitals, health centres, dispensaries, aid posts, registered medical clinics and nursing homes;
- (c) “**authorised person**” means a person authorised under this Act;
- (d) “**authorised pharmacopoeia**” means the current edition for the time being of any of the following, namely, the International Pharmacopoeia, the British Pharmacopoeia, the British Pharmaceutical Codex, the European Pharmacopoeia, the United States Pharmacopoeia and the British Veterinary Codex;
- (e) “**class A drug**”, “**class B drug**” and “**class C drug**” shall be construed in accordance with [section 12](#);
- (f) “**classified drug**” means a class A, B or C drug;
- (g) “**commission**” means the National Drug Authority Commission;
- (h) “**descriptive matter**” means any statement, whether written or oral, which purports to describe the composition or effect of any drug; and references to the publication of descriptive matter shall be references to its publication by way of advertisement, or on or with the container in which the drug is supplied or in any other manner;
- (i) “**disease**” includes injury and bodily or mental deficiency or abnormality;
- (j) “**dispense**”, in relation to a medicine or poison, means to supply a medicine or poison on and in accordance with a prescription duly given by a duly qualified medical practitioner, dentist or veterinary surgeon;
- (k) “**drug**” means any substance or preparation used or intended to be used for internal or external application to the human or animal body either in the treatment or prevention of disease or for improving physiological functions, or for agricultural or industrial purposes;
- (l) “**drug authority**” means the National Drug Authority;

- (m) “**duly qualified**”, used in relation to a medical practitioner, dentist or veterinary surgeon, means a person recognised by law to practise medicine, surgery, dentistry and midwifery or, as the case may be, veterinary surgery;
- (n) “**generic name**” means the International Nonproprietary Name (INN) established by a body of the World Health Organisation;
- (o) “**Indian hemp**” includes the dried flowering or fruiting tops of the pistillate plant known as cannabis sativa or cannabis indica from which the resin has not been extracted, by whatever name the tops are called, and resins obtained from those tops, all preparations of which those resins form the base and all extracts or tinctures obtained from those tops;
- (p) “**inspecting officer**” means a person empowered under Part VII of this Act to enter any premises;
- (q) “**international control**” means the international conventions on the control of narcotic drugs and psychotropic substances;
- (r) “**International Nonproprietary Name (INN)**” means the official name of a drug, regardless of the manufacturer;
- (s) “**licensed person**” means a person licensed under [section 14](#);
- (t) “**licensed seller**” means a person licensed under [section 15](#);
- (u) “**manufacture**” includes any treatment of a plant, mineral or other substance for the purpose of extracting a drug;
- (v) “**Minister**” means Minister responsible for health;
- (w) “**narcotic drug**” means a class A drug or preparation;
- (x) “**pharmacist**” means pharmacist under the Pharmacy and Drugs Act;
- (y) “**prepared opium**” means opium prepared for smoking and includes dross and any other residues remaining after opium has been smoked, and also includes any opium, for whatever purpose prepared, which is capable of being smoked;
- (z) “**proprietary drug**” means a drug distributed for sale by retail under a brand name or other proprietary description and in a form ready for use;
- (aa) “**register**” means the register of specialties maintained under the drug authority;
- (bb) “**restricted drug**” means a classified drug or any other drug which is not an exempted drug;
- (cc) “**substance**” includes a preparation;
- (dd) “**supply**”, with its grammatical variations and cognate expressions, includes, in relation to a drug, the administration of any such drug.

Part II – National drug policy and national drug authority

2. National drug policy

- (1) The national drug policy shall be—
 - (a) to ensure that essential, safe, efficacious and cost-effective drugs are made available to the entire population of Uganda to provide satisfactory health care;
 - (b) to make a continuous review of the needs, knowledge and resources of essential drugs;
 - (c) to promote the rational use of drugs both in the public and private sector;
 - (d) to improve Government regulation and control on manufacture, production, importation, exportation, marketing and use of drugs;

- (e) to provide systematic public information and professional training and retraining of health workers;
 - (f) to improve the registration of drugs and licensing of pharmaceutical premises;
 - (g) to intensify research in all types of drugs, including traditional medicines;
 - (h) to comply with the international regulations on drugs, including the conventions on narcotic drugs and psychotropic substances under international control; and
 - (i) to fight against drug and substance abuse.
- (2) The national drug policy shall relate to the regulation of the importation, production, distribution, marketing, exportation and use of pharmaceuticals in the public as well as in the private sector and to any matter related to the above.

3. Establishment of the National Drug Authority

- (1) There is established a National Drug Authority which shall be a body corporate with perpetual succession and a common seal and may sue or be sued in its corporate name.
- (2) The drug authority shall consist of the chairperson and the following other persons—
- (a) the director of medical services;
 - (b) the commissioner for veterinary services;
 - (c) the commissioner for commissioner for trade;
 - (d) the director, criminal investigation department;
 - (e) the chief of medical services, Ministry of Defence;
 - (f) the chief of pharmaceuticals and health supplies;
 - (g) the head of the Natural Chemotherapeutics Laboratory;
 - (h) the director, Mulago Hospital;
 - (i) a representative of each of the following—
 - (i) the National Medical Stores;
 - (ii) the Uganda Medical Association;
 - (iii) the Pharmaceutical Society of Uganda;
 - (iv) the Uganda Veterinary Association;
 - (v) the head of the School of Pharmacy, Makerere University;
 - (vi) the Uganda herbalists;
 - (vii) the Uganda Dental Association; and
 - (viii) the Joint Medical Stores;
 - (j) the director general of the Uganda AIDS Commission;
 - (k) two other persons appointed from the public.
- (3) The chairperson and the members appointed under subsection (2)(k) shall be appointed by the Minister.
- (4) The members appointed under subsection (3) shall be in office for three years but shall be eligible for reappointment.

4. Application of the seal

- (1) The common seal of the drug authority shall be as the drug authority may determine and shall be kept by the secretary.
- (2) The common seal shall, when affixed into any document, be authenticated by any two signatures of the chairperson, the secretary and any other member of the commission as may be authorised by the drug authority.
- (3) A contract or instrument which if entered into or executed by a person not being a body corporate would not be required to be under seal may be entered into or executed without seal on behalf of the drug authority by the secretary or any other person authorised by the drug authority.
- (4) Every document purporting to be—
 - (a) an instrument issued by the drug authority and sealed with the common seal of the drug authority and authenticated in the manner prescribed in subsection (2); or
 - (b) a contract or instrument entered into or executed by the drug authority shall be received in evidence without further proof as that instrument duly issued or a contract duly entered into or executed unless the contrary is proved.

5. Functions of the drug authority

The drug authority shall be charged with the implementation of the national drug policy and, in particular, but without derogation of the foregoing, shall—

- (a) deal with the development and regulation of the pharmacies and drugs in the country;
- (b) approve the national list of essential drugs and supervise the revisions of the list in a manner provided by the Minister;
- (c) estimate drug needs to ensure that the needs are met as economically as possible;
- (d) control the importation, exportation and sale of pharmaceuticals;
- (e) control the quality of drugs;
- (f) promote and control local production of essential drugs;
- (g) encourage research and development of herbal medicines;
- (h) promote rational use of drugs through appropriate professional training;
- (i) establish and revise professional guidelines and disseminate information to health professionals and the public;
- (j) provide advice and guidance to the Minister and bodies concerned with drugs on the implementation of the national drug policy; and
- (k) perform any other function that is connected with the above or that may be accorded to it by law.

6. Commission and other bodies of the authority

- (1) There shall be a National Drug Authority Commission which shall consist of the chairperson and four other members appointed by the drug authority from among themselves.
- (2) The chairperson of the drug authority shall be the chairperson of the commission.
- (3) The functions of the commission shall be—
 - (a) to exercise the functions of the drug authority which may require exercising when the drug authority is not sitting;

- (b) to monitor and supervise the implementation of the decisions of the drug authority;
 - (c) to establish and revise from time to time, the working procedure of the drug authority;
 - (d) to perform any other function relating to the functions of the drug authority as the authority may direct.
- (4) There shall be the following committees of the drug authority—
- (a) the committee on essential drugs; and
 - (b) the committee on the national formulary.
- (5) The membership of the committee on essential drugs shall be as follows—
- (a) a chairperson appointed by the drug authority;
 - (b) the commissioner of curative services of the Ministry of Health;
 - (c) the chief of pharmaceuticals and health supplies;
 - (d) the chief of medical services, Ministry of Defence;
 - (e) the head of the School of Pharmacy;
 - (f) a representative of each of the following specialities—
 - (i) physician;
 - (ii) paediatrician;
 - (iii) gynaecologist/obstetrician;
 - (iv) surgeon;
 - (v) psychiatrist;
 - (g) a member from the Private Medical Practitioners Association;
 - (h) a non-government organisation pharmacist from the Joint Medical Stores.
- (6) The committee on essential drugs shall have power to co-opt members deemed necessary.
- (7) The membership of the committee on the national formulary shall be as follows—
- (a) a chairperson appointed by the drug authority on the recommendation of the appropriate professional bodies;
 - (b) a member of the faculty of medicine of the universities in Uganda;
 - (c) a member of the faculty of veterinary sciences;
 - (d) a member from the School of Pharmacy;
 - (e) a member from the Pharmaceutical Society of Uganda;
 - (f) a member from the Private Medical Practitioners Association;
 - (g) a member from the Uganda Medical Association;
 - (h) the executive director of the National Bureau of Standards;
 - (i) a representative of each of the following specialities—
 - (i) physician;
 - (ii) surgeon;
 - (iii) paediatrician;

- (iv) gynaecologist/obstetrician;
- (v) psychiatrist.

7. Meetings of the drug authority

- (1) The drug authority shall meet for the discharge of its functions at least six times a year.
- (2) The National Drug Authority Commission shall establish the working procedure for the drug authority.

Part III – Control of the drug supply

8. National list of essential drugs

- (1) There shall be a national list of essential drugs which shall be revised from time to time.
- (2) There shall be a national formulary made of the national list of essential drugs and such other drugs as the authority may, from time to time, approve.
- (3) No person shall import or sell any drug unless it appears on the national formulary.
- (4) Notwithstanding subsection (3), a drug not appearing on the national formulary may be imported and sold after authorisation by the drug authority to meet emergency or extraordinary circumstances.

9. Selection of drug items

The drug authority shall receive from the committee on essential drugs the proposals of the revised list which shall be made in accordance with the available resources and existing diagnostic and therapeutic capacity.

10. Estimation of drug needs

- (1) The commission shall ensure regular assessment and estimation of the national drug needs both in the public and private sectors.
- (2) Estimates of the national drug needs shall be expressed both in unit (quantity) and financial cost.
- (3) For the purposes of providing accurate estimates of drug needs, the commission shall promote and encourage investigations, including studies of current morbidity patterns, drug utilisation and available diagnostic and therapeutic resources.

11. Drug nomenclature

All drugs imported in Uganda shall be labelled, known and prescribed by their International Nonproprietary Names (generic names) except where no such name has been allocated and no satisfactory nonproprietary alternative exists.

12. Restricted drugs

- (1) For the purpose of this Act and subject to this section—
 - (a) the drugs specified in the First, Second and Third Schedules to this Act shall be classified drugs;
 - (b) the drugs and articles specified in the Fourth Schedule to this Act shall be exempted drugs and articles; and

- (c) any classified drug or any other drug which is not exempted shall be deemed to be a restricted drug.
- (2) Subject to subsection (3), where a preparation contains any quantity of a drug which is included in the First, Second or Third Schedule, the preparation shall be deemed to be a classified or restricted drug of the same class as the drug which it contains.
- (3) Where an entry in the First, Second or Third Schedule to this Act defines the proportions of a drug which bring a preparation containing it within the list of restricted drugs, subsection (2) shall not apply to that preparation.
- (4) Where, apart from this subsection, a preparation would fall to be treated as a class A drug and also as a class B or class C drug or both, it shall be treated as a class A drug only.
- (5) Where, apart from this subsection, a preparation would fall to be treated as a drug of both class B and class C, it shall be treated as a class B drug only.

13. Supply and dispensing of restricted drugs

- (1) Subject to this section, no person shall mix, compound, prepare, supply or dispense any restricted drug unless that person is a registered pharmacist, medical practitioner, dentist or veterinary surgeon or a licensed person.
- (2) Subsection (1) shall not prevent—
 - (a) the supply of any drug, other than a drug of class A or B, by a licensed seller;
 - (b) the mixing, compounding or preparing of a drug under the immediate supervision of a registered pharmacist;
 - (c) the supply or dispensing of a restricted drug by a member of the staff of a hospital, dispensary or similar institution which has been authorised to do so by a general or special order of the drug authority;
 - (d) the supply of restricted drugs subject to regulations made by the Minister after consultation with the drug authority, by a representative of a person engaged in the sale and supply of pharmaceutical goods for the purposes of giving free samples of the drugs to persons who may lawfully possess restricted drugs.
- (3) A person registered or enrolled under the Nurses and Midwives Act or any other authorised person may supply or dispense restricted drugs in accordance with regulations made by the Minister in that behalf.
- (4) The supply or dispensing of restricted drugs under subsections (2) and (3) shall be subject to the following—
 - (a) the restricted drug shall be distinctly labelled with the name and address of the person by whom it is supplied or dispensed;
 - (b) the following particulars shall, within twenty-four hours after the restricted drug has been supplied or dispensed, be entered in a book used regularly for the purpose, which shall be known as the Prescription Book—
 - (i) the date on which the restricted drug was supplied or dispensed;
 - (ii) the ingredients and quantity supplied;
 - (iii) the name and address of the person to whom the restricted drug was supplied;
 - (iv) the name and address of the person by whom the prescription was given, except that paragraph (a) shall not apply in any case where any restricted drug is administered by a medical practitioner, dentist, veterinary surgeon or midwife, or under his or her direct supervision and in his or her presence.

- (5) Any record kept under this section shall be open to inspection by an inspector of drugs.

14. Licensed persons

- (1) If, on application made in the prescribed form by any person, the authority is satisfied—
- (a) that the applicant is fit to carry on a business of mixing, compounding and preparing and supplying restricted drugs by retail;
 - (b) that the business, so far as concerns the restricted drugs, will be carried on under the immediate supervision of a pharmacist in each set of premises where the business is to be carried on;
 - (c) in the case of a body corporate, that at least one of the directors is a pharmacist resident in Uganda; and
 - (d) in the case of a partnership, that at least one of the partners is a pharmacist resident in Uganda,

the authority may, on payment of a prescribed fee, issue a licence to the applicant to carry on the business required at the premises and on conditions specified in the licence.

- (2) A licence issued under this section shall be valid for a period specified in the licence, but the drug authority may revoke the licence if, at any time, it is satisfied that the licensed person has contravened any provision of this Act or any condition specified in the licence, or has ceased to be fit to carry on the business.
- (3) A person who carries on the business of a pharmacist without a licence issued under this section commits an offence and is liable to a fine not exceeding one million shillings or to imprisonment not exceeding five years or to both.

15. Licensed sellers

- (1) If, on application made in the prescribed form by a person other than a pharmacist or a licensed person, the authority is satisfied—
- (a) that the applicant is fit to carry on a business of supplying by retail restricted drugs, other than drugs of class A or B;
 - (b) that the area in which the applicant proposes to carry on that business is not sufficiently served by existing facilities for the retail supply of the drugs; and
 - (c) that the applicant is an authorised person,

the authority may issue to the applicant a licence authorising him or her, subject to any conditions specified in the licence, to carry on the business required from the premises specified in the licence.

- (2) A licence issued under this section shall be valid for a period specified in the licence, but the authority may revoke the licence if, at any time, it is satisfied that the holder of the licence has contravened any provision of this Act or any condition specified in the licence, or has ceased to be fit to carry on the business.

16. Places from which restricted drugs may be supplied

- (1) No person shall carry on the business of supplying restricted drugs from any premises—
- (a) if restricted drugs including drugs of class A or B are supplied, unless either a general or a limited certificate is issued under this Act for the purpose;
 - (b) if restricted drugs not including drugs of class A or B are supplied, unless either a general or a limited certificate issued under this Act is in force.

- (2) No person shall supply any drug by means of an automatic machine.

17. Certificates of suitability of premises

- (1) If on application made in the prescribed form for a certificate in relation to any premises, the authority is satisfied that the accommodation, fixtures, equipment and other physical attributes of those premises render those premises suitable for the supply of restricted drugs or for the supply of restricted drugs excluding drugs of classes A and B, it may issue in respect of those premises either a general or limited certificate.
- (2) Every person carrying on the business of supplying restricted drugs from the premises in respect of which a certificate issued under this section is in force shall notify the authority of any alteration in the physical attributes of the premises, or if no alteration occurs in any calendar year, shall notify the authority of that fact before the end of January in the following year.
- (3) A certificate issued under this section shall remain in force until a date specified in the certificate, but the authority may revoke the certificate if, at any time, it is satisfied, on the recommendation of the inspector of drugs, that, owing to an alteration or deterioration in the physical attributes of the premises, the premises have ceased to be suitable for the supply of the restricted drugs, or of restricted drugs other than drugs of classes A and B, as the case may be.
- (4) The authority shall keep a register in the prescribed form of the premises in respect of which a certificate is issued under this section.

18. Loss of class A or B drugs

- (1) Any person entitled under this Act to supply or dispense a class A or B drug shall, upon the loss of that drug in his or her possession or control or of any records kept under this Act in relation to that drug, report that loss to the inspector of drugs, within seven days of the loss, giving particulars of the ingredients and quantities of the drug or the particulars of the records lost.
- (2) A person contravening any provision of this section commits an offence and is liable to a fine not exceeding one million shillings or to a term of imprisonment not exceeding five years or to both.

Part IV – Special provisions relating to classified drugs

19. Classified drugs

The Minister on the advice of the authority may, by statutory instrument, declare a drug to be a classified drug.

20. Need for prescription for classified drugs

- (1) A pharmacist or licensed person shall not supply a class A or class B Group I drug unless it is under prescription reasonably believed by the person supplying the drug to be valid.
- (2) A prescription shall be valid only if—
 - (a) it is in indelible writing, dated and signed with the usual signature of a registered medical practitioner, dentist or veterinary surgeon;
 - (b) it states the name, qualification and address of the person signing it;
 - (c) it states the name and address of the person for whose treatment it is given or, if signed by a veterinary surgeon, of the person in charge of the animal to which the drug is to be administered;
 - (d) it is signed by a dentist, and bears the words “for dental treatment only” or, if signed by a veterinary surgeon, and bears the words “for animal treatment only”;

- (e) it indicates the total amount of the drug to be supplied and the dose to be taken or the manner of its application or use; and
 - (f) it has not previously been fully dispensed.
- (3) A prescription shall be fully dispensed if the drug prescribed has been supplied once, unless it clearly states—
- (a) the number of times it may be dispensed; and
 - (b) the intervals at which it may be dispensed, and shall in that case, be fully dispensed if the drug prescribed has been supplied the stated number of times.
- (4) This section shall not apply—
- (a) if the drug is supplied, whether personally or on a signed order, to a medical practitioner, dentist, veterinary surgeon, pharmacist or licensed pharmacy for the purpose of being subsequently dispensed or supplied or used for purposes of scientific education or research; or
 - (b) if the drug is supplied from the dispensing department of an approved institution in accordance with regulations made by the Minister in that behalf.

21. Action to be taken in relation to prescription

Where a classified drug is supplied under a prescription—

- (a) the person supplying the drug shall enter on the prescription in indelible writing the date on which it is supplied and the name and address of the supplier;
- (b) if the prescription is fully dispensed, it shall be retained by the supplier and, for two years thereafter, shall be kept on the premises at which it was dispensed in such a manner as to be readily available for inspection.

22. Classified drugs to be supplied to responsible persons

A pharmacist or licensed pharmacy shall not supply a class A or B drug to a person who is not reasonably believed by the supplier to be a person to whom the drug may properly be supplied.

23. Supply to conform to prescription

No person shall supply any classified drug which does not conform to the prescription or order under which it is supplied.

24. Classified Drugs Book

- (1) Every person who supplies class A, B or C Group II drugs shall keep in all premises from which the drugs are supplied by him or her a book of the prescribed description to be known as the Classified Drugs Book.
- (2) Subject to subsection (3), before any person supplies class A, B or C Group II drugs, he or she shall enter or cause to be entered in the Classified Drugs Book the following particulars—
 - (a) the name and quantity of the drug to be supplied;
 - (b) the name and address of the person who requires the drug;
 - (c) the purpose for which the drug is stated to be required;
 - (d) the signature of the person to whom the drug is delivered; and
 - (e) the date of the delivery.

- (3) Where any classified drug is sold in the presence of an agent or servant of the person by whom it is to be used or where sale is effected by post, the following provisions shall apply—
- (a) before the sale is completed, the seller shall obtain an order in writing, signed by the purchaser showing—
 - (i) the purchaser's name, address and occupation;
 - (ii) the name and the quantity of drug to be purchased; and
 - (iii) the purpose for which it is required,but where a person represents that he or she urgently requires a classified drug for the purpose of his or her trade, business or profession, and satisfies the seller that, by reason of some emergency, he or she is unable before delivery to furnish the order in writing, the seller may deliver the drug to the purchaser who shall, within twenty-four hours of the sale, furnish the seller with a written order;
 - (b) before the sale is completed, the seller shall satisfy himself or herself that the signature on the order is that of the person by whom it is supposed to be signed and that that person carries on the occupation stated in that order, being an occupation for which the drug is properly required;
 - (c) the requirements of subsection (2) as to the making of entries in the Classified Drugs Book shall be complied with except that in place of the signature of the person to whom the drug is delivered, it shall be sufficient to record "signed order" giving a reference by which the particular signed order may be readily identified;
 - (d) all signed orders and prescribed records of transactions to which this subsection applies shall be retained on the premises where the sales were made for two years.
- (4) Any person who contravenes any of the provisions of this section commits an offence and is liable to a fine not exceeding two million shillings or to imprisonment for a term not exceeding five years or to both.

25. Containers and labels

No person shall supply any classified or restricted drug unless—

- (a) the drug is in a container of the prescribed description; and
- (b) the container bears a label giving the prescribed particulars of its contents.

26. ***

[section 26 repealed by section 93 of [Act 3 of 2016](#)]

27. Possession of classified drugs

- (1) The following persons may be in possession of classified drugs, but to the extent only and subject to the limitations prescribed below —
- (a) any person specified in [section 14](#) for the purposes of that section;
 - (b) a licensed person or seller of classified drugs, on premises registered under this Act;
 - (c) a wholesale dealer licensed under this Act for the purposes of the licence and on the premises so licensed;
 - (d) any person, institution or department to whom a classified drug has been lawfully sold in accordance with this Act, for the purpose for which the sale was made;

- (e) any person for whom the classified drug has been lawfully supplied or dispensed by a duly qualified medical practitioner, dentist or veterinary surgeon or by an approved institution.
- (2) Any person who is in possession of a classified drug otherwise than in accordance with this section commits an offence and is liable to a fine not exceeding two million shillings or to imprisonment for a term not exceeding five years or to both.

28. Withdrawal of authority

- (1) Where any person authorised to obtain or supply narcotics under this Act is convicted of any offence under this Act, if the Minister is of the opinion that that person ought not to be allowed to obtain, possess or supply drugs, he or she may, acting in accordance with the recommendation of the authority by notice published in the *Gazette*, withdraw the authority of that person.
- (2) Where the person whose authority is withdrawn under subsection (1) is a registered or licensed medical practitioner or dentist or a duly qualified veterinary surgeon, the Minister may, by notice published in the *Gazette*, direct that it shall not be lawful for that person to give prescriptions or orders for the purposes of this Act.

29. ***

[section 29 repealed by section 93 of [Act 3 of 2016](#)]

Drugs generally

30. Impure drugs not to be supplied

Any person who—

- (a) sells any drug, medical appliance or similar article which is not of the nature, substance and quality demanded or which, unless otherwise agreed at the time of demand, does not conform to the standards laid down in the authorised pharmacopoeia; or
- (b) supplies any drug which is unwholesome or adulterated or which does not conform to the prescription under which it is supplied,

commits an offence and is liable to a fine not exceeding five million shillings or to imprisonment for a term not exceeding ten years or to both.

31. Power to call for information as to proprietary drugs

- (1) Where the authority has reason to believe that any person is proposing to sell any proprietary drug by retail or to procure, whether directly or indirectly, its sale by retail, the authority may require that person to furnish to it—
 - (a) details of the composition of the drug;
 - (b) copies of any descriptive matter published or proposed to be published in relation to the drug; and
 - (c) any other information that the authority may require.
- (2) No disclosure of information furnished under this section shall be made without the consent of the person by whom it was furnished.

32. Power to prohibit retail sale of proprietary drugs

The authority may prohibit the sale by retail of a proprietary drug if, in the opinion of the authority—

- (a) claims are made for the drug, whether or not in a statement furnished under [section 31](#), which are unjustified;
- (b) the use of the drug may endanger the health of the user or there may be other undesirable effects in the use of the drug;
- (c) details of the composition of the drug furnished under [section 31](#) differ substantially from those disclosed on an analysis of samples of the drug obtained from retail suppliers; or
- (d) descriptive matter published in relation to the drug differs substantially from that, whether or not in the same language, contained in copies furnished to the authority in relation to the drug under [section 31](#).

33. Control of publication of descriptive matter

- (1) Subject to this section, no person shall, by way of advertisement, publish, in whatever manner, in relation to any drug, descriptive matter calculated to lead to the use of that drug—
 - (a) for prevention or treatment of any disease specified in the Fifth Schedule to this Act;
 - (b) for the purpose of termination or influencing the course of human pregnancy; or
 - (c) for any purpose relating to enhancing human potency.
- (2) Subject to this section, the authority may, with the approval of the Minister, serve on any person a notice prohibiting him or her from publishing in relation to any drug descriptive matter referred to in the notice.
- (3) This section shall not apply to the publication of descriptive matter —
 - (a) by direction of the Minister;
 - (b) in a document intended for persons whose profession or employment calls for a knowledge either of drugs generally or of drugs of the description to which the matter in question relates; or
 - (c) for the purposes of an application for the grant of a patent.

34. Return of details of pharmacy business

- (1) Every person carrying on a pharmacy business on any premises shall, within twenty-one days after the commencement by him or her of that business on those premises and annually in the month of January thereafter, send to the authority returns in the prescribed manner, stating—
 - (a) the location and postal address of the premises;
 - (b) the name and principal postal address of the person carrying on the business; and
 - (c) the name of the pharmacist supervising the sale of drugs at those premises.
- (2) If any alteration occurs in the particulars stated in the last return made, the person carrying on the business shall, within twenty-one days of the alteration, send notice in writing to the authority.

35. Drug regulation and registration of specialities

- (1) The drug authority—
 - (a) may scientifically examine any drug for the purposes of ascertaining efficacy, safety and quality of that drug;

- (b) shall institute a system for the approval of drugs or drug combinations not included in the national list of essential drugs.
- (2) The drug authority shall keep a register of specialities in the prescribed form.
- (3) If, on application made in the prescribed manner and on payment of the prescribed fee, the authority is satisfied—
 - (a) that the drug or preparation in respect of which the application is made has not previously been registered; and
 - (b) that the use of the drug or preparation is likely to prove beneficial,the authority shall register the name and description of that drug or preparation.
- (4) Where, on application so made, the authority is not satisfied as aforesaid, it shall notify the applicant that the application is dismissed on the grounds which shall be specified.
- (5) The authority may direct at any time for the deletion of any drug or preparation from the register.
- (6) The register shall, at all reasonable times, be open for public inspection on payment of such fee as may be prescribed.

36. Drug quality

- (1) The drug authority shall advise the Minister on measures to be taken to ensure the quality of drugs imported into or held in stock in the country.
- (2) The execution of the measures prescribed shall be entrusted to bodies charged with the importation and distribution of drugs.
- (3) The inspection of drugs and measures prescribed may be delegated to the chief of pharmaceuticals and health supplies or any other person properly qualified in pharmaceuticals and health supplies.

Wholesale trade

37. Licence required for wholesale supply of restricted drugs

- (1) No person shall carry on a business of supplying restricted drugs by wholesale unless he or she is authorised to carry on that business by a licence granted under this section.
- (2) The authority may, on application made in the prescribed form and upon payment of the prescribed fee, grant a licence for the carrying out of a business of supplying restricted drugs by wholesale, if the authority is satisfied —
 - (a) that the applicant is a person to whom the licence can properly be granted;
 - (b) that the business will be carried on in separate premises apart from any other business;
 - (c) that the business will be carried on in premises under the immediate supervision of a pharmacist;
 - (d) in the case of a company, that at least one of the directors is a pharmacist resident in Uganda; and
 - (e) in the case of a partnership, that at least one of the partners is a pharmacist resident in Uganda.
- (3) A licence granted under this section may include a condition prohibiting or limiting the supply of restricted drugs of a description specified in the condition, and shall be deemed to include a condition prohibiting the supply of any prepared opium or Indian hemp which is prepared for smoking.

- (4) A licence granted under this section shall be valid for a period specified in the licence; but the authority may revoke the licence if, at any time, it is satisfied that the holder of the licence has contravened any provision of this Act or any condition contained in the licence or has ceased to be fit to carry on the business.

Control of manufacture and storage of drugs

38. Restrictions on manufacture of classified drugs

- (1) No person shall manufacture any drug or preparation which is not included on the national formulary unless the drug or preparation is approved by the authority.
- (2) No person, unless approved by the authority in that behalf, shall manufacture a speciality.
- (3) No person shall manufacture any classified drug unless the processes of manufacture are carried out or supervised by a pharmacist.
- (4) Subsection (3) shall not apply to the manufacture of preparations mentioned in the Sixth Schedule to this Act if the processes of manufacture are carried out or supervised by a medical practitioner.

39. Further restrictions on the manufacture of drugs

- (1) The Minister may, by statutory instrument, make regulations further limiting the persons who may manufacture any drug or preparation and the premises in which they may be manufactured, and otherwise controlling their manufacture.
- (2) No person shall manufacture any narcotic drug or psychotropic substances under international control for purposes other than for medical, dental or veterinary use.

40. Clinical trials

- (1) The authority may issue a certificate to any person for the purpose of carrying out clinical trials in respect of a drug that may be specified in the certificate.
- (2) No person may carry out any clinical trial in respect of any drug unless he or she is in possession of a certificate issued under subsection (1).

41. Local research and production

- (1) The National Drug Authority shall encourage research by persons carrying on research and development in herbal and other medicines and where appropriate take such medicines into production as a component of the drug supply.
- (2) Where the drug authority considers it economically advantageous and it is in the interest of the development of a national drug industry, it shall encourage and develop national production of essential drugs.

42. Storage

- (1) Where restricted or classified drugs are kept on any premises, they shall be kept in accordance with the Seventh Schedule, but that Schedule shall not apply to drugs supplied to an individual for the treatment of himself or herself or another individual residing with him or her or an animal in his or her possession or control.
- (2) If an act is done on any premises in contravention of the above subsection then—
 - (a) in a case where the act constitutes a breach of a duty imposed by or under the terms of his or her employment upon a person employed on the premises, that person shall be deemed to have committed an offence;

- (b) in any other case, the occupier of the premises shall be deemed to have committed an offence.
- (3) Nothing contained in subsection (2) shall prevent any person who wilfully removes or alters the label on any container, or does any other act, as opposed to an omission, in respect of a restricted drug, from being treated as having committed an offence under subsection (1).

Part V – Control of transport, import and export of drugs

43. Transportation of drugs

The Minister may, on the advice of the drug authority, make regulations for the control of the transportation of any drug or class of drugs.

44. Importation of pharmaceuticals

- (1) No person or body shall import any drugs into Uganda without having a licence in relation to their import from the drug authority.
- (2) The licence shall be valid for one year and shall state the range of preparations to be imported during that period.

45. Exportation of drugs

- (1) No person or body shall export any drug or preparation without having a licence in relation to that export from the drug authority.
- (2) The licence shall be valid for one year and shall specify the drug to be exported.
- (3) A person who exports any classified drugs shall keep a record in the prescribed form of all exports.

46. Import and export licences

- (1) The authority may grant a permit for the import or the export of a classified drug if—
 - (a) an application for the permit is made in the prescribed form and the applicant pays the prescribed fee; and
 - (b) the authority is satisfied that the applicant is a person to whom the permit can properly be granted.
- (2) No permit shall be granted for the import or export of any narcotic drugs or psychotropic substances under international control, other than for medical, dental or veterinary use.
- (3) A permit granted under this section may be granted generally for the import or export of classified drugs or limited to specified drugs.

Part VI – Further restrictions on narcotics

47. ***

[section 47 repealed by section 93 of [Act 3 of 2016](#)]

48. ***

[section 48 repealed by section 93 of [Act 3 of 2016](#)]

49. ***

[section 49 repealed by section 93 of [Act 3 of 2016](#)]

Part VII – Powers of entry and investigation**50. Powers of entry**

- (1) An inspector or assistant inspector of drugs may enter—
 - (a) at all reasonable times, any premises in respect of which a certificate issued under this Act is in force or on which any person is required to carry out any functions imposed under this Act;
 - (b) at any time, any premises on or in relation to which he or she has reasonable cause to suspect that an offence under this Act has been or is being committed;
 - (c) at any reasonable time, any premises on which a business relating to the manufacture or supply of narcotic drugs is carried on;
 - (d) at any time, any vehicle or vessel which he or she reasonably suspects is being or is about to be used in the commission of an offence under this Act.
- (2) Any police officer not below the rank of assistant superintendent may enter, at any reasonable time, any premises or detain and enter any vehicle or vessel on or in relation to which he or she has reasonable cause to suspect that an offence under this Act has been or is being committed.

51. Powers of investigation

- (1) A drug inspector, assistant inspector of drugs or police officer of the rank of assistant superintendent empowered under this Act to enter any premises, vehicle or any other means of transport may—
 - (a) inspect the premises, vehicle or vessel and any articles found in the premises, vehicle or vessel;
 - (b) require any person on or in the premises, vehicle or vessel to furnish any information in his or her possession as to the activities carried on or in the premises and the person by whom they are carried on or the purposes for which the vehicle or vessel is being used;
 - (c) take away any drug or records and other documents found on or in the premises, vehicle or vessel.
- (2) Where a drug is taken away pursuant to this section, reasonable payment thereof shall be tendered by the inspecting officer, but—
 - (a) no payment need be tendered in respect of a drug if the inspecting officer reasonably suspects that the drug is unfit for its purpose by reason of deterioration, impurity, adulteration or other defect; but if the drug is later found on analysis not to be so unfit, reasonable payment shall be tendered by the inspecting officer in respect of the drug which is not returned to its owner in good condition;
 - (b) no payment shall be made in respect of a drug if the inspecting officer anticipates that proceedings for an offence under this Act will be brought in respect of the drug; but if the proceedings are not commenced within six months, reasonable payment shall be tendered by the inspecting officer in respect of the drug which is not returned to its owner in good condition.

52. Authority to be shown

An inspecting officer exercising any powers conferred by this Act shall produce on demand a duly authenticated document showing that he or she is entitled to exercise those powers.

53. Obstruction

No person shall obstruct an inspecting officer exercising powers under this Part of this Act or fail to comply with a requirement made by him or her in exercise of those powers.

Part VIII – The secretariat and financial provisions**54. Secretariat**

- (1) The drug authority shall have a secretariat which shall be responsible for the day-to-day operations of the drug authority.
- (2) The secretariat shall be headed by the secretary to the drug authority who shall be appointed by the drug authority on terms and conditions that the drug authority may determine.
- (3) In addition to any other functions that may be conferred upon him or her by the drug authority, the secretary shall—
 - (a) have custody of the seal of the drug authority;
 - (b) be responsible for taking the minutes of the drug authority and the commission and for keeping the records of the transactions of the drug authority.
- (4) There shall be other officers and employees of the drug authority as the drug authority may determine.
- (5) An employee of the drug authority shall not, in his or her personal capacity, be liable to any civil or criminal proceedings in respect of any act done or omission made in good faith in the performance of his or her duties under this Act.

55. Funds of the drug authority

- (1) The funds of the drug authority shall consist of—
 - (a) grants from the Government;
 - (b) grants and loans from any body, organisation or person;
 - (c) interest on savings made by the drug authority;
 - (d) money that may accrue to the drug authority in the discharge of its functions; and
 - (e) money from any other source as may be approved by the Minister.
- (2) The drug authority shall possess a bank account in a bank approved by it.

56. Estimates

- (1) The drug authority shall, within three months before the commencement of each financial year, prepare and submit to the Minister, estimates and expenditure for the drug authority for the next ensuing year; and any time before the end of a financial year, the drug authority may prepare and submit to the Minister for approval any estimates supplementary to the estimates of a current year.
- (2) The Minister shall notify the drug authority of his or her decision on the estimates submitted to him or her within one month of the submission of the estimates.

- (3) No expenditure shall be made out of the funds of the drug authority unless that expenditure is part of the expenditure approved by the Minister under the estimates for the financial year in which the expenditure is to be incurred or in supplementary estimates for that year.

57. Accounts

- (1) The drug authority shall keep proper books of account of all its income and expenditure and proper records in relation to those accounts.
- (2) Subject to any direction given by the Minister responsible for finance, the drug authority shall cause to be prepared in respect of each financial year, a statement of account which shall include—
 - (a) a balance sheet, a statement of income and expenditure and a statement of surplus and deficit; and
 - (b) any other information in respect of the financial affairs of the drug authority as the Minister responsible for finance may require.

58. Audits

- (1) The accounts of the drug authority shall, in respect of each financial year, be audited by the Auditor General or an auditor appointed by him or her.
- (2) The drug authority shall ensure that within four months after the end of the financial year a statement of account is submitted to the Auditor General for auditing.
- (3) The Auditor General and any auditor appointed by him or her shall have access to all books of account, vouchers and other financial records of the drug authority and be entitled to have any information and explanation required by him or her in relation to those records.
- (4) The Auditor General shall, within two months after receipt of statements of accounts under this section, audit the accounts and deliver to the drug authority and the Minister a copy of the audited accounts and his or her report on those accounts.

Part IX – Miscellaneous provisions

59. Rational use of drugs

- (1) The drug authority shall, in the interest of public health and the economical use of resources, and in consultation with the bodies concerned, promote the rational use of drugs both in the private and public sector.
- (2) In the implementation of subsection (1), the drug authority may adopt methods and materials which have proved effective in other countries and shall, among other methods, do the following—
 - (a) develop basic and postgraduate training in the health sector;
 - (b) promote public awareness and knowledge of the proper use of drugs; and
 - (c) disseminate information on the purposes and progress of the national drug policy.

60. Offences and penalties

- (1) A person contravening a provision of this Act commits an offence and, where no punishment is provided, is liable—
 - (a) to a fine not exceeding one million shillings;
 - (b) *[paragraph (b) repealed by section 93 of Act 3 of 2016]*
 - (c) *[paragraph (c) repealed by section 93 of Act 3 of 2016]*

- (d) to imprisonment not exceeding one year; or
 - (e) to any two of the above punishments,
- and for any subsequent offence under this Act, a person is liable to a fine not exceeding two million shillings or to a term of imprisonment not exceeding five years or to both.
- (2) A person who commits an offence under this Act and no other punishment is provided is liable—
 - (a) where the offence relates to class A drugs, to a fine not exceeding two million shillings or to a term of imprisonment not exceeding five years or to both;
 - (b) where the offence relates to narcotic drugs or psychotropic substances under international control and is a second or more subsequent offence, to a term of life imprisonment;
 - (c) where the offence relates to manufacturing, smoking or having possession of any narcotic drug or psychotropic substance under international control and is a second or more subsequent offence, to a term not exceeding ten years.
 - (3) Where no case is proved in respect of any drug or article taken from an accused person, the court shall order reasonable payment to the owner in respect of the drug or article which is not returned to him or her in good condition.
 - (4) No proceedings shall be instituted for an offence under [section 35](#) without the consent of the Director of Public Prosecutions.

61. Vicarious criminal responsibility

- (1) Any act or omission which if done by an individual would be an offence under this Act or any regulations made under it shall, if done by a body corporate, be deemed to be an offence committed by every director, secretary and manager of the body corporate, unless the director, secretary or manager proves that the offence was committed without his or her consent or connivance and that he or she exercised all such diligence to prevent the commission of the offence as he or she ought to have exercised, having regard to the nature of his or her functions in that capacity and to all the circumstances of the case.
- (2) If an offence under this Act or any regulations made under it is committed by a partner in a firm, every person who at the time of the commission of the offence was a partner in that firm, or was purporting to act in that office, shall be deemed to have committed the like offence unless he or she proves that the offence was committed without his or her consent or connivance and that he or she exercised all such diligence to prevent the commission of the offence as he or she ought to have exercised, having regard to the nature of his or her functions in that capacity and to all the circumstances of the case.

62. Evidence

- (1) In any proceedings under this Act—
 - (a) any licence, permit or certificate purporting to have been issued under this Act; or
 - (b) any document purporting to state the results of an analysis carried out on behalf of the authority for the purposes of this Act,shall be *prima facie* evidence of the facts stated in it.
- (2) Where, in any proceedings under this Act, a person is charged with—
 - (a) the unlawful possession, sale or supply of any restricted drug and the drug is in a container; or

- (b) any other offence where the contents of a container are in issue in the proceedings, and the container appears to the court to be intact and in its original state of packing by its manufacturer, the contents of the container shall be deemed, unless the contrary is proved, to be of the description specified on the label of the container.

63. Drugs bureau

- (1) There shall be established a drugs bureau under the office of the inspector of drugs.
- (2) The drugs bureau shall—
 - (a) keep and maintain a register in which shall be entered details of the composition of all drugs registered under [section 35](#);
 - (b) keep and maintain a list of all toxic substances, their composition, toxicity and antidotes;
 - (c) supply such information to medical practitioners, dentists or veterinary surgeons in respect of drugs as may be in its possession in emergency cases of poisoning.
- (3) In order to discharge its functions under this section, the drugs bureau may require any person to give any information in his or her possession or control regarding any drug, and that person shall furnish the information within such period as may be specified by the drugs bureau.
- (4) Subject to subsection (2)(c), any information furnished to the drugs bureau under subsection (3) shall be kept confidential and shall not be published without the consent of the person furnishing the information.

64. Regulations

- (1) The Minister may, on the advice of the drug authority, by statutory instrument, make regulations generally for better carrying into effect the provisions of this Act—
 - (a) including the period within which all drugs imported—
 - (i) should be labelled and prescribed by their International Nonproprietary Names (INN) or generic names; and
 - (ii) but not appearing on the national list of essential drugs or the national formulary may be off the market;
 - (b) prescribing the procedure to be followed at meetings, inquiries and other proceedings of the authority and its committees;
 - (c) prescribing conditions to be inserted in licences or permits granted under this Act, and otherwise prescribing things to be done in relation to such licences or permits;
 - (d) laying down conditions in respect of supplies and issues of drugs by hospitals and the storage of drugs by hospitals and the records to be kept;
 - (e) use of drugs in first-aid boxes notwithstanding any other enactment;
 - (f) prohibiting, regulating or restricting the manufacture, sale or advertising of drugs, pharmaceutical preparations and therapeutic substances;
 - (g) regulating, restricting or prohibiting the importation, sale or advertising of surgical instruments and appliances;
 - (h) regulating and restricting the use of classified drugs for agricultural, horticultural, mining and industrial purposes, and the measures to be taken to protect the persons using such classified drugs, including the types and standards of protective clothing which shall be worn;

- (i) requiring the registration and treatment of persons addicted to drugs;
- (j) the registration and operation of authorised persons;
- (k) prescribing anything which under this Act may be prescribed.

65. Amendment of Schedules

The Minister may, after consulting the authority, by statutory order, amend the First, Second, Third, Fourth, Fifth, Sixth, Seventh and Eighth Schedules to this Act.

First Schedule (Sections 12, 29)

Class A drugs or narcotics

The drugs included in this class may only be imported, or exported, manufactured or used, under authority. They may be sold by retail only on the prescription of a duly qualified medical practitioner, dentist or veterinary surgeon, but only for medical, dental or veterinary purposes and may be supplied only by a registered pharmacist or licensed pharmacy.

Acetorphine

Acetyl-alpha-methylfentanyl

Acetyldihydrocodeine

Acetylmethadol

Alfentanil

Allyprodine

Alphacetylmethadol

Alphameprodine

Alphamethadol

Alpha-methylfentanyl

Alphaprodine

Anileridine

Benzethidine

Betacetylmethadol

Betameprodine

Betamethadol

Betaprodine

Bezitramide

Cannabis (Indian hemp) and Cannabis resin (resin of Indian hemp)

Clonitazene

Coca leaf

Cocaine

Codeine

Codoxime

Concentrate of poppy straw (the material arising when poppy straw has entered into a process for the concentration of its alkaloid when such material is made available in trade)

Desomorphine

Dextromoramide

Dextropropoxyphene

Diampromide

Diethylthiambutene

Difenoxin

Dihydrocodeine

Dimepheptanol

Dimethylthiambutene

Dioxaphetyl butyrate

Diphenoxylate

Dipipanone

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

Ethylmethylthiambutene

Ethylmorphine

Etonitazene

Etorphine

Etoxadine

Fentanyl

Furethidine

Heroin

Hydrocodone

Hydromorphanol

Hydroxypethidine

Isomethadone

Ketobemidone

Levomethorphan

Levomoramide

Levophenacymorphan

Levorphanol

Metazocine

Methadone

Methadone-intermediate

Methyldesorphine

Methyldihydromorphine

3-methylfentanyl

Metopon

Moraine-intermediate

Morpheridine

Morphine

Morphine Methobromide and the pentavalent nitrogen morphine derivatives, including, in particular, the morphine-N-oxide derivatives, one of which is Codeine-N-Oxide

Morphine-N-Oxide

MPPP

Myrophine

Nicodicodine

Nicodine

Nicomorphine

Noracymethadol

Norcodeine

Norlevorphanol

Normethadome

Normorphime

Norpipanone

Opium

Oxycodone

Oxymorphone

Para-fluorofentanyl

PEPAP

Pethidine

Pethidine-intermediate-A

Pethidine-intermediate-B

Pethidine-intermediate-C

Phenadoxone

Phenampromide

Phenazocine

Phenomorphan

Phenoperidine
Pholcodine
Piminodine
Piritramide
Proheptazine
Properidine
Propiram
Racemethorphan
Racemoramide
Racemorphan
Sufentanil
Thebacon
Thebaine
Thiofentanyl
Tilidine
Trimeperidine

The isomers, unless specifically excepted, of the drugs in this Schedule whenever the existence of such isomers is possible within the specific chemical designation.

The salts of the drugs listed in this Schedule, including the salts of the isomers as provided above whenever the existence of such salts is possible.

The esters and ethers, unless appearing in another Schedule of the drugs in this Schedule whenever the existence of such esters or ethers is possible. The salts of the drugs listed in this Schedule, including the salts of esters, ethers and isomers as provided above whenever the existence of such salts is possible.

Second Schedule (Sections 12, 29)

Class B drugs or controlled drugs

Group I

The following drugs may be supplied by retail only on the prescription of a duly qualified medical practitioner, dentist or veterinary surgeon, but only for medical, dental or animal treatment respectively.

Acetahexamide
Acetanilide; alkyl acetanilides (except as provided in Group II)
Acetazolamide
Acetylcarbromal
Acocanthera, glycosides of
Adenium, glycosides of
Alcuronium chloride
Allylisopropyl-acetylurea

Amidopyrine, its salts; amidopyrine sulphonates, their salts.

b-Aminopropylbenzene and b-aminoisopropylbenzene; their salts, synthetic compound structurally derived from either of those substances by substitution in the aliphatic part or by ring closure therein (or by both such substitution and closure except ephedrine, its optical isomers, N-substituted derivations oxethazaine, phenylpropanolamine, phenylamine and tropicamide), any salt of any substance falling within this item

p-Aminobenzenesulphonamide; its salts, derivatives of p-aminobenzenesulphonamide having any of the hydrogen atoms of the p-amino group or the sulphonamide group substituted by another radical; their salts (except as provided for in Group II of this class)

Aminometradine

p-aminosalicylic acid; its salts, any preparation of p-aminosalicylic acid; their salts

Aminorex, its salts

Aminosometradine

Amitriptyline, its salts

Androgenic, oestrogenic and progestational substances, the following—

Benzoestral

Derivatives of stilbene, dibenzyl or naphthalene with oestrogenic activity; their salts; their esters

Steroid compounds with androgenic, oestrogenic or progestational activity; their esters (except as provided in Group II of this class)

Antibiotics, the following (except as provided in Group II of this class) any substance the chemical and biological properties of which are identical with or similar to those antimicrobial substances but which is produced by means other than living organisms

Amphomycin and its salts; its esters and salts of such esters

Amphotericins and their salts

Bacitracin

Capreomycin and its salts, its esters and salts of such esters

Cephaloridine and its salts; its esters and salts of such esters

Chloramphenicol

Chlortetracycline

Cycloserine and its salts

Demethylchlortetracycline and its salts

Erythromycin; its salts; its esters and salts of such esters

Framycetin and its salts

Fusidic acid and its salts, its esters and salts of such esters

Gentamycin and its salts, its esters and salts of such esters

Gramicidin

Griseofulvin and its salts

Kanamycin and its salts

Lincomycins, their salts, their esters and salts of such esters

Neomycin and its salts
Novobiocin and its salts
Oleandomycin and its salts; its esters and their salts
Oxytetracycline and its salts
Paramomycin and its salts, its esters and salts of such esters
Penicillin and its salts, derivatives and their salts
Polymyxins and their salts
Rifamycins, their salts, their esters and salts of such esters
Ristocetins and their salts
Spiramycins and its salts
Streptomycin, its salts, its derivatives and their salts
Sulphomycin
Tetracyclins and their salts
Tyrothricin
Vancomycin and its salts
Viomycin and its salts
Virgiamycin and its salts
Azacyclonol acid, its salts
Barbituric acid, its salts; derivatives of barbituric acid; their salts;
compounds of barbituric acid; their salts; their derivatives, their salts;
with any other substance (except as provided in Group II of this class)
Benactyzine; its salts
Benzhexol; its salts
Benztropine, its homologues; their salts
Bromvaltene
Busulphan, its salts
Captodiamine; its salts
Caramiphen; its salts
Carbromal
Carisoprodal
Chlordiazepoxide; its salts
Chlormethiazole
Chlorophentermine; its salts
Chlorothiazide and other derivatives of Benzol 2:4-thiadiazine-7-sulphonamide 1:1-dioxide, whether hydrogenated or not

Chlorphenoxmine; its salts
Chlorpropamide; its salts
Chlorprothixene, and other derivatives of 9-methylenethiazane-ther, their salts
Chlorthalidone, and other derivatives of o-chlorbenzene sulphonamide
Clorexolone, its salts
Corticotrophinis, natural and synthetic
Cycalrbamate
Cycrimine, its salts
Debrisoquine sulphate
Demacarium bromide
Desipramine; its salts
Diazepam, and other compounds containing the chemical structure of dihydro-1-4-benzodiazepine substituted to any degree; their salts
Dinitrocresols (DNOC) their compounds with a metal or base, except when in preparations especially formulated and approved for use in agriculture or horticulture
Dinitronaphthols dinitrophenols dinitrothymols
Disulfiram
Dithienylalkylamines; dithienylakylallylamines; their salts except diethylthaimbutene
Dyflos
Ecothiopate iodide
Ectylurea
Embutramide
Emylcamate
Ergot, alkaloids of, whether hydrogenated or not; their homologues; any salt of any substance falling within this item (except as provided in Group II of this class)
Ethacrycin acid; its salts
Ethchlorynol
Ethinamate
Ethionamide
Ethoheptazine; its salts
Fenfluramine; its salts
Flufenamic acid; its salts; its esters, their salts
Gallamine; its salts; its quaternary compounds
Glutethimide; its salts
Guaiphenesin
Haloperidol; and other 4-substituted derivatives of N-(3-p-fluorobenzolpropyl) piperidine

Hexapropymate

Hydrazines, benzyl, phenethyl and phenoxythyl; their a-methyl derivates; acyl derivatives of any of the foregoing substances comprised in this item; salts of any compounds comprised in this item

Hydroxy-N, N-dimethyltryptamines; their esters or ethers; any salt of any substance falling within this item

Hydroxyurea Hydroxyzine; its salts

Imipramine; its salts

Indomethacin; its salts

Iprindole; its salts

Isoniazid; its salts; derivatives of isoniazid; their salts

Mafenamic acid; its salts; its esters, their salts

Mannomustine; its salts

Mebezonium; its salts

Mebutamate

Meclofenokate; its salts

Mephenesin; its esters

Meprobamate

Meralluride

Mercaptopurine, its salts, derivatives of mercaptopurine, their salts

Mescaline, and other derivatives of phenethylamine formed by substitution in the aromatic ring, their salts

Metaxalone; its salts

Metformin; its salts

Methaqualone; its salts

Methixene; its salts

Methocarbamol

Methoxysalen

Methyldopa

Methylpentynol; its esters and other derivatives

Methyprylone

Metoclopramide; its salts

Metronidazole

Mitopodozide; its salts

Mustine and any other N-substituted derivatives of di-(2-chloroethyl) amine; their salts

Nortryptiline; its salts

Orphenadrine

Oxethazine

Oxyphenbutazone

Oxytocins, natural and synthetic

Paramethadione

Pargyline; its salts

Pemoline, its salts

Pentazocine; its salts

Phenacemide

Phenaglycodal

Phenanthridinium; its salts; its derivatives; their salts; their compounds with other substances

Phenbutrazate

Phencylidine; its salts

Phenetidylphenacetin

Phenformin; its salts

Phenothiazine; its salts; derivatives of; their salts; except dimethoxanate, its salts and promethazine, its salts and its molecular compounds, except also as shall be provided for in Group II of this class and class C

Phenylbutazone; its salts

2-Phenylcinchoninic acid, 2-salicylcinchoninic acid; their salts; their esters

5-Phenylhydantoin; its alkyl and aryl derivatives; their salts

Pituitary gland, the active principles of whether natural or synthetic except as provided for in Group II of this class

Polymethylnebistrimethylammonium salts

Procarbazine; its salts

Procyclidine; its salts

Promoxalan

Propylhexedrine; its salts

Prothionamide

Prothipendyl; its salts

Quinapyramine; derivatives of quinapyramine; their salts; compounds of quinapyramine with other substances

Quinethazone

Rauwolfia; alkaloids of; their salts, derivatives of the alkaloids of rauwolfia; their salts

Spirocholactone

Strychnine (except in preparations included in Group II of this class and class C)

Styramate

Sulphinpyrazone

Sulphonamide, alkyl sulphonamides

Sulphones; their salts, their derivatives

Suprarenal gland, the active principles of; whether natural or synthetic, their esters and salts of such esters, provided for in Group II of this class

Suramin

Syrosingopine; its salts

Tetrabenazine; its salts

Thalidomide; its salts

Thiacetazone; its salts; its derivatives

Thiocarlide; its salts

Thyroid gland; the active principles of; their salts

Tolbutamide

Tretamine; its salts

Triaziquone; its salts

Tribromethyl alcohol

2:2:2-Trichloroethyl alcohol; esters of; their salts

Trimipramine; its salts

Troxidone

Tybamate

Vasopressins, natural and synthetic

Verapamil

Zoxazolamine; its salts

Group II

The following drugs and preparations containing such drugs may be supplied by retail only by a registered pharmacist or licensed pharmacy.

Acetanilide, alkyl, acetanilides, when contained in preparations not intended for the treatment of human ailments

Acetyldihydrocodeine; its salts, when contained in approved preparations containing not more than 100 milligrammes of the drug per dosage unit or 2.5 percent in individual preparations

Alcuronium chloride

Aldrin (except as provided in class C)

Alkali fluorides (other than those specified in class C)

Alkaloids and related substances; the following; their salts, simple or complex; their quaternary compounds

Aconite, alkaloids of (except as provided in class C)

Atropine

Belladonna, alkaloids of (except as provided in class C)

Brucine (except as provided in class C)

Calabar bean, alkaloids of

Cocaine, but only in approved preparations containing less than 0.10 percent w/v of cocaine

Codeine when contained in approved preparations containing not more than 2.5 percent of codeine

Colchicum, alkaloids of

Coniine (except as provided in class C)

Cotarnine

Curare, alkaloids of; curare bases, their salts and quaternary compounds

Ephedrine, its optical isomers except as provided in class C

Ergotamine tartrate but only when contained in oral preparations for the relief of migraine

Gelsemium, alkaloids of

Homatropine

Hyoscine (except as provided in class C)

Hyoscyamine (except as provided in class C)

Jaberandi, alkaloids of

Lobelia, alkaloids of (except as provided in class C)

Morphine, but only in approved preparations containing less than 0.20 percent w/v and as provided in class C

Nicotine (except as provided in class C)

Papaverine

Pomegranate, alkaloids of

Quebracho, alkaloids of

Sabadilla, alkaloids of

Salanaceous alkaloids not otherwise included in this list

Stavesacre, alkaloids of

Stramonium, alkaloids of (except as provided in class C)

Strychnine when contained in preparations containing not more than 1 percent of strychnine and as provided in class C

Thebaine

Veratrum, alkaloids of

Yohimba, alkaloids of

Allylisopropyl acetylurea

Allylprodine; its salts

Alphameprodine; its salts

Alphaprodine, its salts

Aluminium phosphide

Amino-alcohols esterified with benzoic, phenylacetic acid, phenylpropionic acid, cinnamic acid or the derivatives of these acids, their salts

p-Aminobenzenesulphonamide having any of the hydrogen atoms of the p-amino group or of the sulphonamide group substituted by another radical; their salts, but only when in the form of approved preparations containing

not more than 50 percent of such drugs and intended for external use only; or in specifically formulated, labelled and approved preparations for the prevention and treatment of diseases in poultry, or in animal feeding stuff, containing not more than .5 percent of sulphonamides

Amyl nitrite

Anileridine, its salts

Androgenic, oestrogenic and progestational substances as defined in Group I of this Schedule when contained in approved cosmetic preparations

Antibiotics as listed in Group I of this Schedule, when contained in preparations or concentrates for animal feeding stuffs, as provided in class C

Antihistamine substances; the following: their salts; their molecular compounds (except as provided in class C)

Antazoline

Bromodiphenhydramine

Bucizine

Carbinoxamine

Chlopheniramine

Chlorcyclizine

Cinnarizine

Cyproheptadine

3-Di-n-butylaminomethyl -4:5:6 trihydroxyphthalide

Diphenhydramine

Diphenylpyraline

Doxylamine

Isothipendyl

Mebhydrolin

Meclozine

Phenindamine

Pheniramine

Phenyltoloxamine

Promethazine

Pyrrobutamine

Thenalidine

Tolpropamine

Tripolidine

Substances being tetra substituted N derivatives of ethylenediamine or prophyliendiamine

Antimonial substances; antimonates, antimonites, chlorides of; oxides of; sulphides of; organic compounds of antimony

Apomorphine, its salts

Arsenical substances; arsenates, arsenites, halides of arsenic, oxides of arsenic, sulphides of arsenic, organic compounds of arsenic (except as provided in class C)

Barium, salts of; (other than the salts of barium specified in class C)

Butylchloral hydrate

Calcium phosphide

Cantharidin, cantharidates

Carbachol

Carperidine; its salts

Chloral; its addition and its condensation products, their molecular compounds (except as provided in class C)

Chloroform (except as provided in class C)

Clorprenaline; its salts

Creosote obtained from wood (except as provided in class C)

Dehydroemetine; its salts

Dextromethorphan; its salts

Dextrorphan; its salts

Diacetylnalorphine; its salts

Digitalis, glycodies and other active principles of

Dihydrocodeinse when contained in approved preparation containing not more than 2.5 1/5 of

Dihydrocodeine Dimitrocresols (DNOC); their compounds with a metal or base when in preparations specially formulated and approved for agricultural or horticultural purposes

Dinosam; its compounds with a metal or base

Dinoseb; its compounds with a metal or base

Diphenoxylate; its salts, when contained in approved formulation containing not more than 2.5 milligrammes of diphenoxylate base per dosage unit.

Endosulfan (except as provided in class C)

Endothal; its salts (except as provided in class C)

Endrin (except as provided in class C)

Ethylmorphine when contained in preparations containing not more than 2.5 of ethylmorphine

Ethylnoradrenaline; its salts

Etonitazene; its salts

Etoxidine; its salts

Fenranyl; its salts

Fluanisone

Fluoroacetamide; fluoroacetanilide

Furethidine; its salts

Glyceryl trinitrate

Guanidines, the following—

Polymethylene diguanidines di-p-anizyl-p-phenethlauanidine

Heparin and preparations thereof

Hydrocyanic acid, cyanides; double cyanides of mercury and zinc

Hydroxycinchoninic acid; derivatives of, their salts, their esters

4-Hydroxy-3-nitrophenylarsonic acid

Insulin

Iprindole, its salts

Isetharine, its salts

Isoaminile, its salts

Isoprenaline, its salts

Isoprophenamine, its salts

Isopropylarterenol, its salts

N-Isopropylethylnoradrenaline, its salts

Isopropylnoradrenaline, its salts

Isoproterenol, its salts

Laudexium, its salts

Lead; salts of; compounds of lead with acids from fixed oils

Levisoprendhine; its salts

Mannityl hexanitrate

Mercury, oxides of; nitrate of mercury, mercuric ammonium chlorides, potassio-mercuric iodides, organic compounds of mercury; oxycyanides, mercuric thiocyanates, mercuric chloride, mercuric iodide, mercurous chloride (except as provided in class C)

Methoxiphenadrin; its salts

Methoxyphenamine, its salts

Methylaminoheptane; its salts

Methylbromide

Monofluoroacetic acid; its salts

Nalorphine; its salts

Nialamide; its salts

Nitrobenzene (except as provided in class C)

m-Nitrophenol; o-Nitrophenol; p-Nitrophenol

Norcodeine; its salts when contained in approved preparations containing not more than 2.5 percent of Norcodeine

Nux Vomica (except as provided in class C)

Opium, but only in preparation for external use and containing not more than 0.2½ of morphine, calculated as anhydrous morphine

Orciprenaline; its salts

Organofentin compounds, that is compound of fentin

Orthocaine; its salts

Ouabain

Oxalic acid, salts of (except as provided in class C)

Oxycinchoninic acid, its derivatives; their salts; their esters

Pamaquine

Phenamidine; its salts

Phenobarbitone when contained in approved preparations containing Theophylline, ephedrine and not more than 6 milligrammes of phenobarbitone per dosage unit

Phenols (any members 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 60.5 percent weight in weight of phenols; compounds of phenol with a metal; except in substances containing less than the equivalent of 60 percent weight in weight of phenols

Phenothiazine; its salts; when in approved preparations specially formulated and labelled for animal treatment (and as provided in class C)

Phenylene diamines, toluene diamines, other alkylated benzene diamines, their salts (except as provided in class C)

Phenylpropanolamine; its salts

Pholcodine, salts of, when contained in preparations containing not more than $2\frac{1}{2}$ 1/5 of pholcodine (and as provided in class C)

Phosphorous, yellow (except as provided in class C)

Phosphorus compounds, the following (except as provided in class C)

Amiton

Azniphos-ethyl

Azniphos-methyl

Cholorfenrinphos

Demeton-O

Demeton-O-methyl

Demeton-S

Demeton-S-methyl

Diazonon

Dichlorvos

Diethyl 4- methyl-7-courmarinyl phosphorothionate

Diethyl-p-nitrophenyl phosphate

Dimefox

Disulfotan

Ethion

Ethyl-p-nitrophenyl phenylphosphonothimate
Maridox
Mecardbam
Merimphos
Mipafox
Oxydematon-methyl
Paralithion
Phenhaptan
Phosphamidin
Plorate
Schradan
Sulfotep
Tepp (Hepp)
Thionazin
Triphosphoric pantadimethylamide
Vamidothion
Quinapyramine
Savin, oil of
Selenium; its compound (except as provided in class C)
Sodium chlorate
Strophanthus, glycodides of strophanthus
Suprarenal gland, the active principles of; their salts, but only when contained in preparations for external application or in inhalants, rectal preparations or preparations intended for use in the eye
Schradan
Tetrachloroethylene
Thallium, salts of
Toxaphene (except as provided in class C)
Trichloroethyl phosphate
Trimeperidine; its salts
Warfarin; its salts (except as provided in class C)
Zinc phosphide (except as provided in class C)
All radioactive materials
Hypodermic needles and hypodermic syringes used for injection
Preparations of human blood
Substances commonly known as vaccines, sera, toxin, antitoxins and Antigens

Substances known as stethoscopes

Any other substance or preparation not otherwise specified if in a form intended for, or suitable for, injection

Surgical ligatures; any absorbent or protective material capable of being absorbed by the body tissues offered or intended for use in surgical operations

Third Schedule (Section 12)

Class C licensed drugs

Note—The following drugs, except such as are in a form suitable for administration by injection, are the drugs included in this Schedule. They may be sold by retail only by a person or company operating a licensed pharmacy or a licensed drug seller, but in the case of the latter, only in accordance with the terms of his or her licence.

Group I

Any proprietary preparation which does not contain any class A or B drugs

Aconite, alkaloids of, in preparations containing less than 0.02 percent of the alkaloids of aconite

Antibiotics, when contained in preparations or concentrates for animal feeding stuffs

Antihistamines as listed in class B, Group II, but only when contained in preparations for external application only, other than for the eye or nose, and in preparations containing not more than 1 percent of an antihistamine substances intended for application only to the eye or nose

Arsenic, in preparations containing less than the equivalent of 0.01 percent of arsenic trioxide, and dentifrices containing less than .5 percent of acetarsol

Barium sulphide when contained in depilatories

Belladonna, alkaloids of, in preparations containing less than 0.15 percent of the alkaloids of belladonna, calculated as hyscyamine

Brucine, when contained in surgical spirit containing not more than 0.2 percent of brucine

Chloroform, in preparations containing not more than 5 percent of chloroform

Codeine, when contained in preparations in a proportion of less than 1.5 percent; and also when contained in compound tablets of Codeine B.P., or tablets of a similar composition each containing not more than 10 milligrammes of codeine; and being in sealed containers holding not more than twenty-five such tablets

Coniine, in preparations containing less than 0.1 percent of coniine

Creosote, preparations containing not more than 10 percent of creosote obtained from wood

Emetine, preparations containing less than 1 percent of emetine

Ephedrine salts when contained in preparations containing less than 1 percent of the alkaloids of ephedra; tablets of ephedrine hydrochloride containing not more than 30 milligrammes of ephedrine hydrochloride per tablet, and being in sealed containers holding not more than fifty such tablets

Ethylmorphine, in preparations containing less than 0.2 percent of ethylmorphine

Hyoscine, when in preparations in the form of tablets and intended for use in travel sickness

Hyoscyamine, in preparations containing less than 0.15 percent of hyoscyamine

Lobelia, alkaloids of, in preparations for the relief of asthma or in substances containing less than 0.50 percent of the alkaloids of lobelia

Mercuric ammonium chloride when contained in an ointment containing not more than 5 percent of mercuric ammonium chloride or when contained in approved cosmetic preparations

Mercuric oxide when contained in yellow oxide of mercury ointment

Morphine in approved preparations containing less than 0.2 percent of anhydrous morphine

Nux Vomica, alkaloids of; in preparations containing less than 0.2 percent of the alkaloids of Nux Vomica calculated as strychnine

Ohenothiazine, when in approved preparations, specially formulated and labelled for animal treatment

Phenylene diamines, touene diamines, other alkylated benzene diamine; their salts, when contained in preparations intended for use as hair dyes

Pholcodine in approved preparations containing not more than 1 percent of pholcodine

Solenium; its compounds when contained in hair lotions and similar preparations

Stramonium, alkaloids of, in preparations for the relief of asthma or substances containing less than 0.15 percent of alkaloids calculated as hyoscyamine

Strychnine, in preparations containing less than 0.2 percent of strychnine

Sulphadimidine, when contained in approved preparations formulated and labelled for the treatment of poultry diseases containing not more than 16 percent w/v of sulphadimidine

Group II

Aldrin, when contained in preparations for agricultural or horticultural purposes, and sold in the original container as supplied by the manufacturer

Ammonia

Barium carbonate, when in preparations intended for the destruction of rats and mice

Barium silico fluoride

Chloralose, when contained in preparations for the destruction of rats and mice

Chlordane, when contained in preparations for agricultural or horticultural purposes

Dieldrin, when contained in preparations for agricultural or horticultural purposes and sold in the original container as supplied by the manufacturer

Endosulfan, when contained in preparations for agricultural or horticultural purposes

Endothal, when contained in preparations for agricultural or horticultural purposes

Endrin, when contained in preparations for agricultural or horticultural purposes

Formaldehyde

Formic acid

Hydrochloric acid

Hydrofluoric acid

Metallic oxalates other than potassium quadroxalate in photographic solutions

Nicotine, its salts; when contained in preparations intended for agricultural or horticultural purposes

Nitric acid

Nitrobenzene; when contained in agricultural or horticultural insecticides; in substances for the treatment of bee disease and in ointments for animal treatment

Paraquat; salts of

Phenols as defined in class B, Group II of the Second Schedule to this Act in substances containing less than 60 percent weight of phenols; compounds of phenol with a metal in substances containing less than the equivalent of 60 percent weight of phenols

Phosphoric acid

Phosphorus, yellow, when contained in rat poisons

Phosphorus compounds as listed in class B, Group II, but only when contained in preparations specially formulated, packed and labelled for agricultural or horticultural purposes and sold in the original container as supplied by the manufacturer

Potassium fluoride

Potassium hydroxide

Potassium quadroxalate

Sodium fluoride

Sodium hydroxide

Sodium nitrate

Sodium solico fluoride

Sulphuric acid

Toxaphene, but only in preparations intended for use in agricultural or horticultural purposes, and sold in the original container as supplied by the manufacturer

Warfarin, its salts, when contained in preparations for the destruction of rats and mice

Zinc phosphide when in preparations intended for the destruction of rats and mice

Fourth Schedule (Section 12)

Exempted drugs and articles

The following drugs and articles are known as exempted drugs and articles.

Adhesives

Ammonia, substances containing less than 5 percent of ammonia, refrigerators.

Antifouling compositions

Antimony, chlorides of, when contained in polishes

Batteries and accumulators

Builders' materials

Ceramics

Chemicals not included in class A, B or C when packed and labelled for culinary and cooking purposes

Creasote, obtained from coal tar

Dentifrices

Distempers

Dressings on seeds or bulbs

Electrical valves

Enamels
Explosives
Fireworks
Fitters, fire extinguishers
Fluorescent lamps
Formaldehyde, when in photographic glazing or hardening solution
Glazes
Glue
Inks
Lacquer solvents
Laundry materials; blue, bleaches and starch
Loading materials
Matches
Medicated soap
Motor fuels and lubricants
Nitrobenzene, when contained in polishes
Oxalic acid and metallic ozalates when contained in polishes and cleaning powders
Paints (other than pharmaceutical paints)
Phenylmercuric salts, when used in a concentration not exceeding 0.01 per cent in toilet and cosmetic preparations as a preservative or in textiles or antiseptic dressings as a bacteriostat or fungicide
Photographic paper
Pigments
Plastics
Propellants
Rubber
Tar (coal or wood)
Tobacco
Varnishes

Fifth Schedule (Section 33)

Diseases as to which publication of descriptive matter is restricted or prohibited

1. Syphilis, gonorrhoea, soft chancre and any form of genitourinary disease or other diseases connected with the human reproductive functions.
2. Any of the following—
 - Amenorrhoea
 - Arteriosclerosis

Bladder stones
Blindness Brights' disease
Cancer
Cataract
Deafness
Diabetes
Diphtheria
Dropsy
Epilepsy or fits
Erysipelas
Gallstones
Glaucoma
Goitre
Heart disease
Hernia or rupture
Kidney stones
Leprosy
Locomotorataxy
Lupus
Nephritis or Brights' disease
Paralysis
Pleurisy
Pneumonia
Poliomyelitis
Scarlet fever
Schistosomiasis
Septicaemia
Smallpox
Tetanus or lockjaw
Trachoma
Tuberculosis or consumption
Any structural organic ailment of the auditory system

Sixth Schedule (Section 38)

Preparations that may be manufactured by, or under the supervision of, a duly qualified medical practitioner

1. Preparations containing extracts of pituitary, suprarenal, thyroid, liver, pancreas or parathyroid glands, or stomach
2. Preparations containing the active principles of any of the aforesaid glands or the salts of the active principles of any of those glands

Seventh Schedule (Section 42)

Requirements as to the storage of classified drugs

1. All class B and class C (Group II) drugs and preparations except when in use shall be kept—
 - (a) under secure lock and key—
 - (i) in a separate room or compartment specially reserved for keeping these drugs and partitioned off from the rest of the premises; or
 - (ii) in a suitable cupboard, box or other receptacle specifically reserved for keeping drugs, and kept in a place apart from anything containing food or drink; and
 - (b) the drugs shall be kept in a place ordinarily accessible only to the person in charge of the drugs, or to some person under his or her immediate supervision and control; and

the key of the room, compartment, cupboard, box or other receptacle in which these drugs are kept shall be retained under the control of the person in charge of the drugs.
2. All class A drugs and preparations shall, except when in use, be stored in a separate store or cupboard apart from all other drugs, in accordance with the requirements of paragraph 1 above, except that if stored in a cupboard or similar receptacle the cupboard or other receptacle shall be so fixed in position as to be immovable
3. No class A, class B (Group I) or class C (Group II) drugs shall be kept in a part of any premises to which members of the general public normally have access.
4. All drugs and preparations for external use shall be kept separate from drugs and preparations intended for internal use

Eighth Schedule (Section 43)

Consignment and transportation of classified drugs.

1. No person shall consign for transport any drug specified in this Schedule, unless the outside of the package is labelled conspicuously with the name or description of the drug and a notice indicating that it is to be kept separate from food and from empty food containers
2. No person shall, knowingly, transport any drug specified in this Schedule 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 drug, or is otherwise adequately protected from the risk of contamination.

Aldrin

Aluminium phosphide

Arsenical preparations

Barium, salts of

Dieldrin

Dinitrocresols (DNOC), their compounds with a metal or base when contained in preparations for use in agriculture or horticulture

Dinosam, its compounds with a metal or base when contained in preparations for use in agriculture or horticulture

Dinoset, its compounds with a metal or base when contained in preparations for use in agriculture or horticulture

Endosulfan

Endrin

Endothal, its salts

Ethylene dibromide; ethylene dichloride

Fluoroacetamide, fluoroacetanilide

Hydrocyanic acid, cyanides

Mercury, its halides when contained in preparations for use in agriculture or horticulture

Methyl bromide

Monofluoroacetic acid; its salts

Nicotine; its salts

Organo tin compounds, the following compounds of gentin

Phosphorous compounds, the following—

Amiton, azinphos-ethyl, azinphos-methyl; chlorgenrinphos; demeton-O-methyl; demeton-O; demeton-S; demeton-S-methyl, diazinon, dichlorvos, diethyl 4-methyl-7-coumarinyl phosphorothianate, diethyl-p-nitrophenyl phosphate; dimefox; disulfotam ointment; disulfoton; ethion; ethyl-pitrophenyl, phenylphosphonothionate; mazidox, mecarbam, mevinphos, mipafox, oxydematon-methyl; parathon, phenkapton, photate, phosphamidon schradon, sulfetep, TEPP, HEPP, thionazin, triphosphoric pentadimethylamide; vamidothion

Selenium; its compounds when contained in preparations for use in agriculture or horticulture

Strychnine

Thallium, salts of