PHARMACY AND POISONS RULES

ARRANGEMENT OF RULES

Rule

SCHEDULES

SCHEDULE IX – PERMIT AUTHORISING FARMERS AND OTHER PERSONS TO BE IN POSSESSION OF SUBSTANCES SPECIFIED IN GROUP II OF SCHEDULE IV TO THE RULES

PHARMACY AND POISONS RULES

[L.N. 186/1957, L.N. 443/1957, L.N. 332/1958, L.N. 426/1958, L.N. 498/1958, L.N. 550/1959, L.N. 114/1960, L.N. 587/1961, L.N. 242/1963, L.N. 631/1963, L.N. 92/1964, L.N. 365/1964, L.N. 115/1968, L.N. 125/1969, L.N. 248/1969, L.N. 41/1971, L.N. 120/1984, Corr. No. 52/1984, L.N. 51/1985, L.N. 61/2002, L.N. 91/2004, L.N. 191/2010.]

1. Citation

These Rules may be cited as the Pharmacy and Poisons Rules.

2. Interpretation

(1) In these Rules, unless the context otherwise requires—

"animal" includes bird;

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

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

"British Pharmaceutical Codex", **"British Pharmacopoeia"** and **"British Veterinary Codex"** include supplements;

"food" includes drink;

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

"poison" means a poison included in Part I or Part II of the Poisons List as the case may be;

"Poisons List" means the Poisons List for which provision is made in section 25 of the Act;

"sell" includes an agreement to sell and an offer to sell or any other act whatsoever by which willingness to enter into any transaction of sale is expressed, and an offer to sell includes the exposing of goods for sale.

(2) A reference to the percentage of a poison contained in a substance shall, unless otherwise expressly provided, be construed so that a reference to a substance containing 1 per cent of a poison means—

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

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

and so in proportion for any greater or lesser percentage.

(3) For the purposes of these Rules—

(a) a poison shall not be taken to be sold, issued or supplied otherwise than in accordance with a prescription or other order by reason only that the prescription or order specifies a quantity of the poison in terms of the imperial system and the quantity sold, issued or supplied is the equivalent of that amount in the metric system, or by reason only that the prescription or

order specifies a quantity of the poison in terms of the metric system and the quantity sold, issued or supplied is the equivalent of that amount in the imperial system; and

(b) the quantity of a poison in the imperial system which is the equivalent of a particular quantity in the metric system, and the quantity of a poison in the metric system which is the equivalent of a similar quantity in the imperial system, shall be deemed to be that set out as such in the Tables of Equivalents contained in the *British Pharmacopoeia*, the *British Pharmaceutical Codex* or the *British Veterinary Codex*.

3. Importation of drugs and Part I poison

(1) Any person, other than a person issued with an import licence in form 17 set out in Schedule VIII, who imports any drug or Part I poison from any place outside Kenya shall be guilty of an offence.

(2) The Board may issue an import licence authorizing the importation of any drug cosmetics, herbals, medical devices, technologies upon payment of two per cent Freight on Board value or Part I poison to the following persons—

(a) an authorized seller of poisons;

(b) persons licensed under the provisions of sections 27 and 28 of the Act, in accordance with the terms of such licence;

(c) the Government or a local authority and its institutions for public purposes;

(d) a person requiring to import poisons for industrial purposes;

(e) any *bona fide* tourist or visitor having in his possession, on his arrival in Kenya, any drug or poison for the medical treatment or any other lawful use by himself or any other member of his party;

(f) any duly qualified medical practitioner, dentist or veterinary surgeon who satisfies the Board that he is urgently in need of a drug or poison which he is unable to obtain in Kenya;

(g) a hospital at and of which a medical practitioner registered under the Medical Practitioners and Dentists Act (Cap. 253), is resident and in direct control.

(3) A person requiring to import Part I poison under the provisions of paragraph (2)(d) shall indicate in his application for an import licence the purpose for which the poison is required and, if the importer is not the person who will use the poison, the name or names of the person or persons to whom the poison will be sold.

(4) The Board may, without assigning any reason therefor, refuse an application for a licence to import any drug or Part I poison; and any person aggrieved by the decision of the Board may appeal to the Minister whose decision shall be final.

(5) A person issued with an import licence under these Rules shall comply with the rules and regulations of the Central Bank of Kenya which may be in force from time to time.

(6) A person, issued with an import licence under these Rules who imports any drug or Part I poison from any place outside Kenya shall keep a full, accurate and separate record of such importation.

(7) A person referred to in paragraph (2) and a licensed seller of Part II poison shall not import Part II poison without an import licence issued under these Rules.

[Corr. No. 52/1984, L.N. 120/1984, L.N. 191/2010, r. 3.]

3A. Restriction on the importation or manufacture of specified drugs

(1) No person, shall, without the approval of the Registrar, in writing import or manufacture any of the following drugs—

(a) amphetamine;
(b) amoberbital;
(c) amferpramone;
(d) barbital;
(e) dexamphetamnie;
(f) cyclovarbital;
(g) ethinamate;
(h) lysergide, or its salts;
(i) glutethimide;
(j) methamphetamine;
(k) methyphenidate;
(l) meprobamate;
(m) methaqualone, or its salts;
(n) methylphenobarbital;
(o) methylprylon;
(p) psilocin;
(q) psilocybine;
(r) phencyclidine;
(s) phenmetrazine;
(t) phenobarbital;
(u) pentobarbital;
(v) pipradrol;
(w) secobarbital;
(y) medroxyprogeshrone and its salt; and
(z) foreign traditional medicine of any description.

(2) A person who contravenes paragraph (1) shall be guilty of an offence.

[L.N. 125/1969, L.N. 191/2010, r. 2.]

4. Exportation of drugs and poisons

(1) A person, other than a person, issued with an export licence in form 23 set out in Schedule VIII, who exports any drug or poison to a destination outside Kenya shall be guilty of an offence.

(2) The Board may issue an export licence authorizing the exportation of any drug or poison to an authorized seller of poisons or other person licensed to deal in poisons under section 27 or section 28 of the Act.

(3) The Board may, without assigning any reason therefor, reject an application for a licence to export drugs or poisons to any destination outside Kenya; and a person who is aggrieved by the decision of the Board may appeal to the Minister whose decision shall be final.

(4) A person issued with an export licence under these Rules shall comply with the rules and regulations of the Central Bank of Kenya which are in force from time to time.

(5) Every authorized seller of poison and any other person licensed to deal in poisons under section 27 or section 28 of the Act who exports any drugs or poisons to a destination outside Kenya shall—

(a) keep a full and accurate record of those exports; and

(b) if the drug or poison is sent by post, send the export by registered or parcel post; and

(c) comply with the requirement of rule 15 relating to the transportation of poisons.

(6) A person who fails to comply with the provisions of paragraph (5) shall be guilty of an offence.

5. Exemptions

(1) A person who imports a Part I poison for industrial purposes in accordance with the provision of rule 3 may, notwithstanding the provisions of section 26 of the Act—

(a) lawfully possess the Part I poison in the quantity authorised to be imported;

(b) sell the poison so imported to the person named in the application as the purchaser, and the purchaser may, notwithstanding the provisions of section 26 of the Act, lawfully possess the poison.

(2) An authorised seller of poisons shall not be required to comply with the provisions of section 29(2) and section 30 of the Act in the case of—

(a) substances specified in Schedule I if the sale is effected by, or under the supervision of, a registered pharmacist; and

(b) machine-spread plaster;

(c) surgical dressings;

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

(e) corn paints in which the only poison is a poison included in the Poisons List under the heading of “Cannabis”.

(3) Nothing in Part III of the Act or in these Rules shall apply to—

(a) an article in Group I of Schedule II;

(b) a poison specified in the first column of Group II of Schedule II to these Rules if contained in or in the form of any of the articles or substances specified in the second column.

(4) The requirements of subrule (c) of section 34(1) of the Act shall not apply to any substance specified in Schedule III.

6. Poisons to be supplied only upon prescription

(1) Subject to subrule (2), no person shall sell by retail a Part I poison specified in Schedule IV except on and in accordance with a prescription given by a duly qualified medical practitioner, dentist or veterinary surgeon in the form provided by this rule.

(2) Where an authorised seller of poisons has reasonable cause to believe that a person ordering a Part I poison is a duly qualified medical practitioner, dentist or veterinary surgeon and who is by reason of some emergency unable to furnish such a prescription immediately, he may, notwithstanding that no such prescription has been given, if the person undertakes to furnish him with such a prescription within the twenty-four hours next following, deliver the poison ordered in accordance with the directions of the person, so, however, that notwithstanding anything in the directions, the supply shall not be repeated unless the prescription has been given.

(3) A person by whom any such undertaking has been given who fails to deliver to the seller a prescription in accordance with the undertaking, or who, for the purpose of obtaining delivery of a poison under subrule (2), makes a statement which is to his knowledge false, shall be guilty of an offence.

(4) The provisions of this rule shall not apply to—

(a) a sale referred to in section 29(1) of the Act;

(b) the sale by an authorised seller of poisons of a substance specified in Group II of Schedule IV to a farmer or other person concerned with the welfare of animals as a regular part of the exercise of his trade, business or profession who is in possession of a permit issued by a duly qualified veterinary surgeon;

(c) the sale of strychnine, in quantities not exceeding four ounces at any one time to persons authorised by the District Commissioner to obtain this substance for the purposes of poisoning vermin.

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

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

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

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

(d) have written thereon, if given by a dentist, the words “for dental treatment only” or, if given by a veterinary surgeon, the words “for animal treatment only”;

(e) specify the total amount of the medicine to be supplied and, except in the case of a preparation which is to be used for external treatment only, the dose to be taken.

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

(a) the prescription shall not be dispensed more than once unless the prescriber has directed thereon either that it may be dispensed a stated number of times or that it may be dispensed at stated intervals;

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

(c) a prescription which contains a direction that it may be dispensed a stated number of times but no direction as to the intervals at which it may be dispensed shall not be dispensed more often than once in three days, and a prescription which contains a direction that it is to be dispensed at stated intervals but no direction as to the number of times that it may be dispensed shall not be dispensed more often than three times;

(d) at the time of dispensing or, where a poison has been delivered in accordance with subrule (2), on the subsequent receipt of the prescription there shall be noted on the prescription above the signature of the prescriber the name and address of the seller and the date on which the prescription was dispensed;

(e) except in the case of a prescription which may be dispensed again, the prescription shall, for a period of two years, be retained and kept on the premises on which it was dispensed so as to be readily available for inspection.

(7) For the purposes of subrule (4)(b) a permit—

(a) shall be in the form set out in Schedule IX; and

(b) shall be produced on every occasion when supplies are required; and

(c) on every occasion the supplier shall endorse the permit with his name and address and the date.

(8) A person who fails to comply with the provisions of subrule (6) shall be guilty of an offence.

7. Restriction of sales by licensed sellers of Part II poisons

(1) No person may, by virtue of being a licensed seller of Part II poisons, sell or offer for sale a poison otherwise than in accordance with the provisions of his licence.

(2) A licensed seller of Part II poisons shall not sell a poison, other than ammonia, hydrochloric acid, nitric acid, potassium quadroxalate and sulphuric acid, except in a closed container as closed by the manufacturer or other person from whom the poison was obtained.

(3) A person who fails to comply with the provisions of subrule (2) shall be guilty of an offence.

8. Restriction of sales by person licensed to deal in poisons for mining, agricultural or horticultural purposes

(1) No person may, by virtue of being licensed to deal in poisons for mining, agricultural or horticultural purposes, sell or offer for sale a poison otherwise than in accordance with the provisions of his licence.

(2) A person licensed to deal in poisons for mining, agricultural and horticultural purposes shall not sell—

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

(b) a Part I poison unless—

(i) the purchaser thereof is a person engaged in the trade, business or profession of mining, agriculture or horticulture and requires the poison for the purposes of his trade, business or profession; and

(ii) the sale is made by one of the persons named in the application for the licence to sell the poisons; and

(iii) the poison, if it be one of the substances referred to in Schedule V, shall, in addition to any other requirements of the Act and these Rules, be labelled in the manner described in that Schedule; and

(iv) the requirements of section 30 of the Act are complied with.

(3) A person who fails to comply with the provisions of subrule (2) shall be guilty of an offence.

9. Labelling of containers

(1) A container of poison required to be labelled in accordance with section 34 of the Act shall be labelled clearly and distinctly in the English language with the required particulars and in the following manner—

(a) the name of the poison shall be the term by which the poison is specified in the Poisons List:

Provided that—

(i) where the term describes a group of poisons and not the poison specifically, the name of the poison shall be—

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

(B) in any other case, the accepted scientific name or name descriptive of the true nature and origin of the poison, and in such cases the appropriate name of the poison shall be written in English or in Latin;

(ii) in the case of a preparation in the *British Pharmacopoeia* or the *British Pharmaceutical Codex* or the *British Veterinary Codex* or a dilution or admixture of such a preparation, or a surgical dressing for which a standard is described in the *British Pharmaceutical Codex* it shall be sufficient to state the name, synonym or abbreviated name used to describe the preparation or surgical dressing in the *British Pharmacopoeia* or the *British Pharmaceutical Codex* or the *British Veterinary Codex* with the addition of the letters B.P. or B.P.C or B.Vet.C., as the case may be;

(b) the particulars as to the proportion which a poison contained in a preparation bears to the total ingredients shall be expressed as the percentage which the poison bears to the total ingredients:

Provided that—

(i) in the case of a preparation containing a poison specified in the first column of Schedule VI, it shall be sufficient to state on the label the particulars specified in the second column of that Schedule against the description of the poison;

(ii) in the case of a preparation or surgical dressing which is named in accordance with the provisions of proviso (ii) to subrule (1)(a), it shall not be necessary to state on the label the proportion of the poison contained in the preparation, and in the case of any dilution or admixture of such a preparation, it shall be sufficient to state the proportion which the preparation bears to the total ingredients of the dilution or admixture;

(iii) where the poison is in tablets, pills, cachets, capsules, lozenges or similar articles, or in ampoules, it shall be sufficient to state on the container thereof the number of the articles, and the amount of the poison or the amount of the preparation contained in each tablet, pill, catchet, capsule, lozenge or other similar article;

(c) the word “Poison” or the alternative indication of character specified in rule 10, as the case may be, shall—

(i) in the case of a poison not specified in Schedule I or in Group B of Part II of the Poisons List, either be printed in red letters on a contrasting background or in letters of some other colour set against a red background;

(ii) in all cases be easily legible and either on a separate label or surrounded by a line within which there must be no other words.

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

(3) Directions for the use of a poison shall be given in the English language, in addition to any other language.

(4) Where poison is contained in an ampoule, cachet or other similar article the box or receptacle containing the ampoules, catchets or other articles only need be labelled in pursuance of the provisions of section 34 of the Act and these Rules.

(5) Where the container of a poison or the container of an ampoule, cachet or other similar article is labelled in accordance with the provisions of the Act and these Rules, an outer cover or wrapper to that container used only for the purpose of delivery or transport need not be similarly labelled if it complies with the provisions of rule 15.

(6) A person who sells a poison not labelled in accordance with the provisions of these Rules shall be guilty of an offence.

10. Indication of character of poison

(1) A poison specified in Schedule V shall be labelled with the words and in the manner specified in that behalf in Schedule V.

(2) The words specified in Schedule V shall not be modified in meaning by the addition of other words or marks and shall—

(a) in the case of a poison not specified in Schedule I or in Group B of Part II of the Poisons List, be printed in red letters on a contrasting background or in some other colour on a red background;

(b) in all cases be easily legible on a separate label or surrounded by a line within which there must be no other words.

11. Directions as to use

(1) No person shall sell liquid poison in bottles of more than 120 fluid ounces capacity unless the bottle is labelled with the words “NOT TO BE TAKEN”.

(2) No person shall sell embrocation, liniment, lotion, liquid or antiseptic, or other liquid medicine for external application, which contains poison, unless the container is labelled with the name of the article and the words “FOR EXTERNAL USE ONLY”.

(3) No person shall sell hydrocyanic acid or cyanide unless the container is labelled with the words “WARNING. This container holds a poisonous substance and should be opened and used by persons having expert knowledge of the precautions to be taken in its use.”

(4) A person who fails to comply with any provision of this rule shall be guilty of an offence.

12. Containers for poisons

(1) No person shall keep, sell or consign for transport a poison unless—

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

(b) in the case of a liquid contained in a bottle of capacity of not more than 120 fluid ounces, not being a medicine made up ready for the internal treatment of human ailments, the outer surface of the bottle is fluted vertically with ribs or grooves recognisable by touch.

(2) The provisions of subrule (1)(b) shall not apply to the sale or the keeping of poisons for the purposes of education, research or analysis by a person or institution concerned with scientific education, research or chemical analysis.

13. Safe custody of poisons

(1) No person engaged in a trade, business or profession shall knowingly have in his possession or under his control a poison, unless the following conditions are complied with at all times when the poison is not in actual use—

(a) the poison shall be kept under lock and key—

(i) in a separate room or compartment specially reserved for keeping poisons and partitioned off from the rest of the premises; or

(ii) in a cupboard, box or other receptacle specially reserved for keeping poisons, clearly marked with the words "Poisons Only", and kept in a place apart from anything containing food or drink;

(b) the poison shall be kept in a place ordinarily accessible only to persons lawfully having access thereto;

(c) the key of the room, compartment, cupboard, box or other receptacle in which poisons are kept shall be retained under the control of the person in charge of the poison.

(2) The provisions of subrule (1) of this rule shall not apply to the possession of—

(a) a substance specified in Schedule I;

(b) a substance specified in Group B of Part II of the Poisons List;

(c) medicines prescribed for the personal use of the person having possession or control thereof.

(3) A person in possession of a container or other receptacle which has been used for containing a poison and which is no longer required for that purpose shall by destruction or other means render that container or receptacle innocuous.

(4) Poisons for the treatment of human ailments shall be kept entirely separate from other poisons.

(5) A person who fails to comply with any provisions of this rule shall be guilty of an offence.

13A. Pharmaceutical representative's permit

(1) A representative of a person engaged in the sale and supply of pharmaceuticals containing a poison may, in the course of business, give free samples of such products to persons who may lawfully possess Part I poisons if he—

(a) is in possession of a permit issued by the Board in that behalf; and

(b) enters the following particulars, at the time of issue, in a book used regularly for the purpose—

(i) the date on which the poison was issued;

(ii) the name and quantity of the poison given; and

(iii) the name and address and signature of the person to whom the poison was given.

(2) Every application for a permit under paragraph (1) of this rule shall be made to the Board in form 18 in Schedule VIII and shall be accompanied by a fee of twenty-five shillings in respect of the issue of the permit.

(3) Every permit under paragraph (1) of this rule—

(a) shall be in form 19 in Schedule VIII to these Rules;

(b) shall expire on the 31st December of the year of issue or on the earlier termination of the employment by the person concerned of the person in respect of whom the permit is issued.

[L.N. 41/1971.]

14. Special provisions with respect to hospitals

(1) All poisons not in actual use in any hospital, infirmary, dispensary, clinic, nursing home or other similar institution at which human ailments are treated shall be kept under the control of the person in charge of the institution or some fit and proper person specially detailed for that purpose and shall only be issued for use as required.

(2) In any such institution, at which medicines are dispensed in a dispensing or pharmaceutical department in charge of a person appointed for that purpose, no medicine containing a poison shall, except in a case of emergency, be supplied from that department for use in the wards, operating theatres or other sections of the institution except upon a written order signed by a duly qualified medical or dental practitioner or by a sister or nurse in charge of a ward, theatre or other section of the institution; and the person supplying the medicine shall label the container with the words describing its contents and, in the case of medicines containing poisons other than poisons specified in Schedule I to these Rules or in Group B of Part II of the Poisons List, in addition thereto, an indication that the poison is to be stored in a cupboard reserved solely for the storage of poisons.

(3) Any poison, other than a poison specified in Schedule I or in Group B of Part II of the Poisons List, issued for use in any ward, theatre or other section of the institution shall, at all times when not actually in use, be stored in a cupboard reserved solely for the storage of poisons.

(4) The person in charge of the institution shall, not less than once in every three months, carry out, or arrange and be responsible for the carrying out by a medical practitioner, a pharmacist or some other person appointed for the purpose by the person in charge, of an inspection of—

(i) all stores, cupboards and other places where poisons are kept in the institution;

(ii) the methods by which poisons are issued, dispensed and used in the institution; and

(iii) all books and other records whatsoever kept in the institution for the purpose of recording the purchase, issue and use of poisons.

(5) The person carrying out the inspection shall submit copies of his report in form 20 in Schedule VIII to these Rules—

(i) to the person in charge of the institution, if that person has not himself carried out the inspection; and

(ii) to the registrar.

(6) A person who fails to comply with any provision of this rule shall be guilty of an offence.

[L.N. 41/1971.]

15. Transport of poisons

(1) No person shall consign for transport a poison specified in Schedule VII unless the outside of the package is labelled conspicuously with the name or description of the poison and a notice indicating that it is to be kept separate from food and from empty food containers.

(2) No person shall knowingly transport a poison specified in Schedule VII in a vehicle in which food is being transported unless the food is carried in a part of the vehicle effectively separated from that containing the poison, or is otherwise adequately protected from the risk of contamination.

(3) A person who fails to comply with any provision of this rule shall be guilty of an offence.

16. Manufacture of drugs

(1) No person shall manufacture for sale any drug which is or may be used for the treatment of any human or animal ailment unless he is in possession of a licence for that purpose issued by the Board.

(2) Every application for a licence under paragraph (1) of this rule shall be made to the Board in Form 21 in Schedule VIII to these Rules and shall be accompanied by a fee of one hundred shillings in respect of the issue of the licence, which shall be refundable if the licence is not granted.

(3) Upon an application for a licence under this rule, the Board may, in its absolute discretion, refuse to grant the licence, or may grant the licence either unconditionally or subject to conditions as it may think fit.

(4) A licence under this rule shall be in Form 22 in Schedule VIII to these Rules.

(5) In an establishment in which drugs are manufactured, whether for sale or otherwise, for the purpose of the treatment of any human or animal ailment, such manufacture shall be carried out by, or under the supervision of—

(a) a registered pharmacist; or

(b) a person having a Fellowship or Associateship of the Royal Institute of Chemistry or an equivalent qualification recognized by the Board.

(6) The Board may, by notice in the *Gazette*, exempt any establishment or class of establishment from any or all of the provisions of this rule.

(7) A person who contravenes any of the provisions of this rule, or who fails to comply with any condition of a licence issued thereunder, shall be guilty of an offence.

[L.N. 41/1971.]

17. Restriction on sale of mepacrine and bisulphate tablets

(1) A person who sells mepacrine tablets containing less than 95.0 per cent or more than 105.0 per cent of 100 milligrams of Mepacrine Hydrochloride as described in the *British Pharmacopoeia* shall be guilty of an offence and liable to a fine not exceeding five hundred shillings or to imprisonment for a term not exceeding one month or to both, and in addition to any penalty imposed under these Rules the Court may order any article in respect of which the offence has been committed or which has been used for the commission of the offence to be forfeited.

(2) A person who sells quinine bisulphate tablets containing less than 95.0 per cent or more than 105.0 per cent of 5 grains of Quinine Bisulphate as described in the British Pharmacopoeia and containing any colouring matter shall be guilty of an offence and liable to a fine not exceeding five hundred shillings and to imprisonment for a term not exceeding one month or to both such fine and such imprisonment , and in addition to any penalty imposed under these Rules the court may order any article in respect of which such offence has been committed or which has been used for the commission of the offence to be forfeited.

18. The Poisons Book

(1) The Poisons Book shall be in the form set out in Schedule VIII.

(2) In the case of a person licensed under the provisions of section 27 of the Act as a wholesale dealer in poisons or an authorised seller of poisons having a wholesale section distinct and separate from any retail shop in which complete and detailed records of the receipts and disposals of all poisons are regularly maintained, the Board may, upon such conditions as it may deem fit to impose, relieve that person of the necessity to record sales by way of wholesale in the Poisons Book.

19. Fees

The following fees shall be paid in connection with matters arising under the Act—

	Annual Amount (*KSh.*)
(a) For a certificate of registration as a pharmacist/ Pharmaceutical Technologist	5,000
(b) For the restoration of name to the register	5,000
(c) Professional Practice	5,000
(d) For the registration of premises	10,000
(e) For a wholesale dealer's license per annum	30,000
(f) For a license to deal in mining, agricultural and horticultural Poisons per annum	5,000
(g) For a license to sell Part II poisons per annum	5,000
(h) For a license to manufacture drugs per product	5,000
(i) Advertisement per product	5,000
(j) Pharmaceutical representative permit	5,000
(k) For application approval for import permit 2% value Freight on Board.	
(l) Good Manufacturing Practice Audit per site—	
(i) Foreign manufacturing site	USD 4,000
(ii) Local manufacturing site	USD 1,000

(m) Training and Assessment/Evaluation fees for pharmacists and pharmaceutical technologists

	Kenyan Citizen	Foreigners
Stage/Level I	9,500/=	22,000/=
Stage/Level II	7,000/=	20,000/=

	KSh.
(n) New application, inspection and course approval fees for pharmacy training institutions	
(i) Degree programmes	400,000
(ii) Diploma programme	210,000
(o) Renewal of Annual course approval fees (sect 8)	KSh.
(i) Degree programmes	60,000
(ii) Diploma programme	30,000
(p) Indexing of students in the pharmacy training institutions in Kenya	KSh.
(i) Degree programmes	1,000
(ii) Diploma programme	1,000

20. Forms

The forms to be used under the Act and these Rules shall be those set out in Schedule VIII.

21. Preservation of books

All books and other prescribed records for the purposes of Part III of the Act shall be preserved on the premises on which the sales recorded therein were made for a period of two years from the date on which the last entry was made therein.

SCHEDULE I

[Rule 5.]

SUBSTANCES EXEMPTED FROM THE PROVISIONS OF SECTION 29(2) AND SECTION 30(1)(A) AND (B) OF THE ACT

GROUP I

A substance containing any of the poisons specified in the first column below if the poison content is less than the percentage specified in the second column.

Poison	*Percentage of poison content below which substance is exempted*
1. Alkaloids, including their salts simple or complex:—	
2. Aconite, alkaloids of	0.02 percent.
3. Apomorphine	0.20 percent.
4. Atropine	0.15 percent.
5. Belladonna, alkaloids of	0.15 percent, calculated as hyoscyamine.
6. Brucine	0.20 percent.
7. Coca, alkaloids of	0.10 percent.
8. Cocaine	0.10 percent.
9. Codeine	1.50 percent.
10. Colchicum, alkaloids of	0.50 percent, calculated as colchicine.
11. Coniine	0.10 percent.
12. Cotarnine	0.20 percent.
13. Ecgonine and its esters	0.10 percent.
14. Emetine	1.00 percent.
15. Ethylmorphine	0.20 percent.
16. Gelsemium, alkaloids of	0.10 percent.
17. Homatropine	0.15 percent.
18. Hyoscine	0.15 percent.
19. Hyoscyamine	0.15 percent.
20. Jaborandi, alkaloids of	0.50 percent.
21. Lobelia, alkaloids of	0.50 percent.
22. Morphine	0.20 percent, calculated as anhydrous morphine.
23. Morpholinylethylmorphine	1.50 percent.
24. Papaverine	1.00 percent.
25. Pomegranate, alkaloids of	0.50 percent.
26. Sabadilla, alkaloids of	1.00 percent.
27. Solanaceous alkaloids, not otherwise included in this Schedule	0.15 percent, calculated as hyoscyamine.
28. Stavesacre, alkaloids of	0.20 percent.
29. Strychnine	0.20 percent.
30. Thebaine	1.00 percent.
31. Veratrum, alkaloids of	1.00 percent.
32. Adrenalin, its salts, in preparations for external use only	0.10 percent.

SCHEDULE I—*continued*

Poison	*Percentage of poison content below which substance is exempted*
33. Amino-alcohols, esterified with benzoic acids, phenylacetic acid, phenylpropionic acid, cinnamic acid or the derivatives of these acids	10.00 per cent of esterified amino-alcohols.
34. Antimonial poisons	Equivalent of 1.00 per cent of antimony trioxide.
35. Arsenical poisons	Equivalent of 0.01 per cent of arsenic trioxide and dentifrices containing less than 0.50 per cent of acetarsol.
36. Butyl chloral hydrate	10.00 percent.
37. Cantharidin	0.01 percent.
38. Cantharidates	Equivalent of 0.01 per cent of cantharidin.
39. Chloral formamide	10.00 percent.
40. Chloral hydrate	10.00 percent.
41. Digitalis, glycosides and other active principles of	One unit of activity (as defined in the *British Pharmacopala*) in two grams of the substance.
42. Dinitrocresols (DNC), their compounds with a metal or a base	Equivalent of 5.00 per cent of dinitrocresols.
43. Hydrocyanic acid	0.15 per cent weight in weight of hydrocyanic acid (HCN).
44. Insulin	Not exceeding 80 units in 1 mil.
45. Cyanides	Equivalent of 0.10 per cent weight in weight of hydrocyanic acid (HCN).
46. Mercuric chloride	1.00 percent.
47. Mercuric iodide	2.00 percent.
48. Nitrates of mercury	Equivalent of 3.00 per cent weight in weight of mercury (Hg).
49. Potassio-mercuric iodides	Equivalent of 1.00 per cent of mercuric iodide.
50. Organic compounds of mercury	Equivalent of 0.20 per cent weight in weight of mercury (Hg).
51. Nux vomica	0.20 per cent of strychnine.
52. Opium	0.20 per cent of morphine calculated as anhydrous morphine.
53. Para-amino-benzoic acid, esters of; their salts	1.00 percent.
54. Para-aminobenzenesulphonamide; its salts; derivatives of para-aminobenzenesulphonamide having any of the hydrogen atoms of the para-substituted group or of the sulphonamide group substituted by another radical; their salts; when incorporated in a base for external application only	50 percent.

SCHEDULE I—*continued*

GROUP II

Antibiotics, the following—

Bacitracin

Gramicidin

Neomycin

Polymyxins

when incorporated in a base for treatment of the skin.

Chloramphenicol

when incorporated in a special base for the treatment of the feet of animals.

Anti-histamine substances, the following; their salts; their molecular compounds—

Antazoline.

Bromodiphenhydramine.

Buclizine.

Carbinoxamine.

Chlorcyclizine.

Chlorpheniramine.

Cinnarizine.

Clemizole.

Cyclizine.

Cyproheptadine.

3-Di- n-butylaminomethyl-4, 5, 6-trihydroxypthalide.

Diphenhydramine.

Diphenylpyraline.

Doxylamine.

Isothipendyl.

Mebhydrolin.

Meclozine.

Phenindamine.

Pheniramine.

Phenyltoloxamine.

Promethazine.

Pyrrobutamine.

Thenalidine.

Tolpropamine.

Triprolidine.

Substances being tetra-substituted N derivatives of ethylenediamine or propylenediamine.

SCHEDULE II

[Rule 5.]

[L.N. 92/1964, L.N. 125/1969.]

ARTICLES EXEMPTED FROM PART III OF THE ACT AND THESE RULES

GROUP I

Adhesives, anti-fouling compositions; builders' materials; ceramics; distempers; electrical valves; enamels; explosives; fillers; fireworks; fluorescent lamps; glazes; glue; inks; lacquer solvents; loading materials; matches; medicated soaps; motor fuels and lubricants; paints other than pharmaceutical paints; photographic paper; pigment;: plastics; propellants; rubber; varnishes; tyrothricin, framycetin.

GROUP II

Poison	*Substance or article in which exempted*
1. Acetanilide; alkyl acetanilides	Substances not being preparation for the treatment of human ailments.
2. Brucine	Surgical spirit containing not more than 0.015 per cent of brucine.
3. Emetine	Ipecachuana; extracts and tinctures of ipecachuana; substances containing less than 0.05 per cent of emetine.
4. Ephedra, alkaloids of	Substances containing less than 1 per cent of the alkaloids of ephedra.
5. Formic acid	Substitutes containing not less than 5 per cent weight in weight formic acid (HCOOH).
6. Jaborandi, alkaloids of	Substances containing less than 0.025 per cent of the alkaloids of jaborandi,
7. Lobelia, alkaloids of	Preparations for the relief of asthma in the form of cigarettes, smoking mixtures or fumigants, substances containing less than 0.1 per cent of the alkaloids of lobelia.
8. Nicotine	Tobacco.
9. Pomegranate, alkaloids of	Pomegranate bark.
10. Solanaceous alkaloids	Stramonium contained in preparations for the relief of asthma in the form of cigarettes, smoking mixtures or fumigants.
11. Stavesacre, alkaloids of	Soaps; ointments; lotions for external use.
12. Ammonia	Substances not being solutions of ammonia or preparations containing solutions of ammonia substances containing less than 5 per cent weight in weight of ammonia (NH_3); refrigerators; smelling bottles.
13. Antibiotics as defined in the Poisons List	Preparations or concentrates for animal feeding.
14. Antihistamine substances as defined in the Poisons List	Preparations intended for external application only.
15. Antimony, chlorides of	Polishes.
16. Arsenical poisons	Pyrites ores or sulphuric acid conltaining arsenical poisons as naturaimpurities.
17. Barium, salts of	Witherite other than finely ground witherite.

SCHEDULE II—*continued*

	Poison	*Substance or article in which exempted*
18.	Beta-aminopropylbenzene; its salts; its N-alkyl derivatives; their salts; beta-aminoisopropyl-benzene; its salts; its Nalkyl derivatives; their salts	Appliances for inhalation in which the poison is absorbed in inert solid material.
18A.	Carbarsone	Poultry feeding stuffs containing not more than 0.0375 per cent Carbarsone.
19.	Chloroform	Substances containing less than 10 per cent of chloroform.
20.	Creosote obtained from wood	Substances containing less than 50 per cent of creosote obtained from wood.
21.	Formaldehyde	Substances containing less than 5 per cent weight in weight of formaldehyde (HCHO); photographic glazing or hardening solutions.
22.	Hormones as defined in the Poisons List	Cosmetic preparations for external applicaton and plant hormones.
23.	Hydrochloric acid	Substances containing less than 9 per cent weight in weight of hydrochloric acid (HCL).
24.	Lead acetate	Substances containing less than 4 per cent of lead acetate.
25.	Lead, compounds of	Machine-spread plasters.
26.	Mercuric chloride	Batteries.
27.	Mercuric chloride; mercuric iodide; organic compounds of mercury	Dressings on seeds or bulbs.
28.	Mercury, nitrates of	Ointments containing less than the equivalent of 3 per cent weight in weight of mercury (Hg).
29.	Nitric acid	Substances containing less than 9 per cent weight in weight of nitric acid (HNO_3).
30.	Nitrobenzene	Substances containing less than 0.1 per cent of nitrobenzene; soaps containing less than 1 per cent of nitrobenzene; polishes.
31.	Oxalic acid; metallic oxalates	Laundry blue; polishes.
32.	Oxycinchoninic acid; derivatives of their salts; their esters	Preparations for external applications only containing not more than the equivalent of 3 per cent oxycinchoninic acid.
33.	Paranitrobenzylcyanide	Photographic solutions containing less than the equivalent of 0.1 per cent of HCN.
34.	Paranitrophenol	Preparations for use in agriculture and horticulture containing not more than 0.5 per cent of paranitrophenol as a preservative.
34A.	Phenylcinchoninic acid	Preparations for external application only containing not more than the equivalent of 10.1 % of phenylcinchoninic acid.
35.	Phenols	Carvacrol; creosote obtained from coal tar; essential oils in which phenols occur naturally; medicines containing less than 1 per cent of phenols; nasal sprays, mouth washes, pastilles, lozenges, capsules,

SCHEDULE II—*continued*

Poison	*Substance or article in which exempted*
	pessaries, ointments or suppositories containing less than 2.5 per cent of phenols; paratertiaryamyl phenol; smelling bottles; soaps for washing; solid substances other than pastilles, lozenges, capsules, pessaries, ointments and suppositories, containing less than 60 per cent of phenols; tar (coal or wood), crude or refined; tertiary butylcresol; thymol; animal dips containing less than 6 per cent of phenols.
36. Phenylene diamines; toluene diamines; other alkylatedbenzene diamines; their salts	Substances other than preparations for the dyeing of hair.
37. Phenylmercuric salts	Toilet, cosmetic and therapeutic preparations containing not more than 0.01 per cent of phenylmercuric salts as a preservative, and textiles containing not more than 0.01 %, as a bacteriostat and fungicide.
38. Picric acid	Substances containing less than 5 per cent of picric acid.
39. Potassium hydroxide	Substances containing less than 12 per cent of potassium hydroxide; accumulators; batteries.
40. Procaine	Combined with antibiotics when contained in preparations or concentrates for animal feeding.
41. Sodium ethyl mercurithiosalicylate	Therapeutic substances containing less than 0.1 per cent of sodium ethyl mercurithiosalicylate as a preservative.
42. Sodium fluoride	Substances containing less than 3 per cent of sodium fluoride as a preservative.
43. Sodium hydroxide	Substances containing less than 12 per cent of sodium hydroxide.
44. Sodium silicofluoride	Substances containing less than 3 per cent of sodium silicofluoride as a preservative.
44A. Sulphone	Substance containing a mixture of dapsone and pyrimethamine, recommended for use as an antimalarial.
45. Sulphuric acid	Substances containing less than 9 per cent weight in weight of sulphuric acid (H_2SO_4); accumulators, batteries; fire extinguishers.

SCHEDULE III

[Rule 5.]

[L.N. 248/1969.]

SUBSTANCES EXEMPT FROM CERTAIN LABELLING REQUIREMENTS

1. Antibiotics.
2. Hormones; natural and synthetic; any preparations, admixture, extract or other substance containing any proportion of any substance having the action of any hormone.
3. Isoniazid; its salts, derivatives of isoniazid; their salts.
4. Para-amino-salicylic acid; its salts; any preparation of para-amino-salicylic acid; its salts.
5. Sulphones; their salts; their derivatives.
6. Thiacetazone; its salts; its derivatives.
7. Drugs as defined in the Pharmacy and Poisons (Control of Drugs) Rules, 1969, which are not specifically named in the Schedule to the Poisons List Confirmation Order.

SCHEDULE IV

GROUP I

Substances required to be sold by retail only upon a prescription given by a duly qualified medical practitioner, dentist or veterinary surgeon.

1. Acetanilide; alkyl acetanilides.
2A. Acetohexamide.
3. Acetylcarbromal.
4. Allylisopropylacetylurea.
5. Amidopyrine; amidopyrine sulphonates; their salts.
6. Amitryptyline; its salts.
7. Antibiotics.
8. Antimony, organic compounds of, for injection.
9. Arsenic, organic compounds of, for injection.
10. Azacyclonal; its salts.
11. Barbituric acid; its salts, derivatives of barbituric acid; their salts; compounds of barbituric acid, its salts, its derivatives, their salts, with any other substance.
12. Benactyzine; its salts.
13. Benztropine and its homologues; their salts.
14. Benzhexol; its salts.
15. Bromvaletone.
16. Busulphan; its salts.
17. B-Aminopropylbenzene and B-aminoisopropylbenzene and any compound structurally derived from either of those substances by substitution in the side chain or by ring closure therein (or by both such substitution and such closure), except ephedrine N-methylephedrine, N-diethylamioethylephedrine, phenylpropanolamine and prenylamine; any salt of any substance falling within this item.
18. Captodiame; its salts.
19. Carbromal.

SCHEDULE IV—*continued*

20. Carisoprodol.
21. Chlordiazepoxide; its salts.
22. Chlormethiazole; its salts.
23. Chlorothiazide and other derivatives of benzo-1, 2, 4-thiadiazine-7-sulphonamide 1, 1-dioxide, whether hydrogenated or not.
24. Chlorphenoxamine.
25. Chlorphentermine.
26. Chlorpropamide; its salts.
27. Chlorprothixene, and other derivatives of 9-methylenethiaxanthen; and their salts.
28. Chlorthalidone, and other derivatives of O-Chlorobenzene sulphonamide.
29. Chlorexolone.
30. Curare; alkaloids of; curare bases and salts.
31. Cyclarbamate.
32. Cycrimine; its salts.
33. Demecarium bromide.
34. Desipramine; its salts.
35. 4; 4-diamidino-diazoamino-benzene; its salts.
36. Diazepam, and other compounds containing the chemical structure of 1:4 benzodiazepine substituted to any degree; their salts.
37. Dinitrocresols (DNC); their compounds with a metal or a base, except preparations for use in agriculture or horticulture.
38. Dinitronapthols; dinitrophenols; dinitrothymols.
39. Disulfiram.
40. Dithienylallylamines; dithienylalkylallylamines; their salts except diethylthiambutene, dimethylthiambutene and ethylmethylthiambutene.
41. Ectylurea.
42. Emylcamate.
43. Ergot; alkaloids of; homologues of; their salts.
44. Ethchlorvynol.
45. Ethinamate.
46. Ethionamide.
47. Ethoheptazine; its salts.
48. Gallamine; its salts; its quaternary compounds.
49. Haloperidol, and other 4 substituted derivatives of N-(3-p. fluorobenzoylpropyl) piperidine.
50. Hexapropymate.
51. Hormones, adrenal cortical, natural and synthetic; any preparations, admixture, extract or other substance containing any proportion of any substance having the action of any adrenal cortical Hormone.
52. Hormones, sex, natural and synthetic and analogous substance, except when in the form of avian implants.
53. Hydrazines, benzyl phenethyl or phenoxyethyl; their a-methyl derivatives; acyl derivatives of any of the foregoing substances comprised in this item; salts of any compounds comprised in this item.
54. 4-Hydroxymethyl-2, 2-diisopropyl-1, 3-dioxolan.
55. Hydroxy N-N-dimethyl tryptamines, esters or ethers of these; salts of any of the foregoing (Psilocin and Psilocybe).
56. Hydroxyzine; its salts.
57. Imipramine; its salts.
58. Indomethacin; its salts.

SCHEDULE IV—*continued*

59. Isoniazid; its salts, derivatives of isoniazid; their salts.
60. Mannomustine; its salts.
61. Mephenesin; its esters.
62. Meprobamate.
63. Mercaptopurine; its salts, derivatives and their salts.
64. Metaxolone.
65. Metformin; its salts.
66. Methaqualone; its salts.
67. Methixene; its salts.
68. Methocarbamol.
69. Methoxsalen.
70. Methylpentynol; its esters and other derivatives.
71. Mustine and any other N-substituted derivatives of di-(2 Chloroethyl) amine; their salts.
72. Nortryptyline; its salts.
73. Orphenadrine; its salts.
74. Oxethazaine.
75. Oxyphenbutazone.
76. Para-aminobenzenesulphonamide; its salts; derivatives of para-aminobenzenesulphonamide having any of the hydrogen atoms of the para-amino group or of the sulphonamide group substituted by another radical; their salts, except when contained in ointments or surgical dressings or in preparations for the prevention and treatment of diseases in poultry.
77. Para-amino-salicylic acid; its salts; any preparation of para-aminosalicylic acid, its salts.
78. Paramethadione.
79. Pargyline; its salts.
80. Pemoline; its salts.
81. Phenacemide.
82. Phenaglycodol.
83. Phenanthridinium and its derivatives.
84. Phenbutrazate.
85. Phenetidylphenacetin.
86. Phenformin; its salts.
87. Phenothiazine, derivatives of; their salts; except dimethoxanate, its salts and promethazine, its salts and its molecular compounds.
88. Phenylbutazone; its salts.
89. Phenylcinchoninic acid; salicylcinchoninic acid; their salts, their esters.
90. Phenylhydantoin; its alkyl and aryl derivatives; their salts.
91. Pituitary gland, the active principles of; except when contained in preparation intended for external application only or, except in the case of lysinevasopressin or oxytocin, in inhalants.
92. Polymethylenebistrimethylammonium salts.
93. Procyclidine; its salts.
94. Promoxolan.
95. Propylhexedrine; its salts; except when contained in inhalers.
96. Prothionamide.
97. Prothipendyl.
98. Quinapyramine and analogous substances; their salts.
99. Quinethazone.

SCHEDULE IV—*continued*

100. Rauwolfia, alkaloids of; derivatives of; their salts.
101. Strychnine except in preparations included in Part II of the Poisons List.
102. Styramate.
103. Sulphinpyrazone.
104. Sulphonal; alkyl sulphonals.
105. Sulphones; their derivatives; their salts.
106. Suprarenal gland medulla, the active principles of; their salts; except when contained in preparations intended for external application only or in inhalants, rectal preparations or preparations intended for use in the eye.
107. Syrosingopine.
108. Tetrabenazine; its salts.
109. Thalidomide; its salts.
110. Thiacetazone; its salts; its derivatives.
111. Thyroid gland, the active principles of; their salts.
112. Tolbutamide.
113. Tretamine; its salts.
114. Triazi quone.
115. Tribromethyl alcohol.
116. Trimipramine.
117. Troxidone.
118. Zoxazolamine; its salts.

GROUP II

SUBSTANCES TO WHICH RULE 6(3)(B) APPLIES

1. Antibiotics.
2. Arsenic, organic compounds of, for injection.
3. 4:4?-diamidino-diazoaminobenzene; its salts.
4. Phenanthridinium and its derivatives.
5. Para-aminobenzenesulphonamide; its salts; derivatives of para-aminobenzene-sulphonamide having any of the hydrogen atoms of the para-substituted group or any of the sulphonamide group substituted by another radical; their salts.
6. Quinapyramine; its salts.

SCHEDULE V

[Rules 8 and 10.]

INDICATION OF CHARACTER OF POISON

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

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

Beta-aminopropylbenzene; its salts; its N-alkyl derivatives; their salts.

SCHEDULE V*continued*

Beta-aminoisopropylbenzene; its salts; its N-alkyl derivatives; their salts.

Insulin.

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

Pituitary gland, the active principles of.

Thyroid gland, the active principles of; their salts.

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

Medicines (other than medicines mentioned in paragraph 1 of this Schedule) made up ready for the internal treatment of human ailments except in the case of a substance included in the First Schedule.

3. To be labelled with the words “Poison. For animal treatment only”—

Medicines made up ready for the treatment of animals.

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

Preparations for the dyeing of hair containing phenylene diamines, toluene diamines or other alkylated-benzene diamines or their salts.

5. To be labelled with the words “Caution. This substance is caustic”—

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

6. To be labelled with the words “Caution. This substance is poisonous. The inhalation of its vapour, mist, spray or dust may have harmful consequences. It may also be dangerous to let it come into contact with the skin or clothing”—

Dinitrocresols (DNC), their compounds with a metal or a base, except preparations for the treatment of human ailments and except winter washes containing not more than the equivalent of five per cent of dinitrocresols.

Dinosam, its compounds with a metal or a base.

Dinoseb, its compounds with a metal or a base.

Fluoroacetamide; Fluoroacetanilide.

Phosphorus compounds, the following—

Diethyl thiophosphate of ethyl-mercapto-ethanol, dimefox, ethyl-para-nitrophenyl-benzene thiophosphonate, hexaethyl tetraphosphate (HETP), 4-methyl hydroxy-coumarin-diethyl thiophosphate, mipafox, parintrophenyl-diethyl phosphate, parathion, schradan, tetraethyl pyrophosphate (TEPP), triphosphoric pentadimethylamide, di-isopropyl fluorophenate, demeton, mazidox, methyl demeton, sulphotepp, amiton, demeton-O, demeton-S, demeton-O-methyl, demeton-S-methyl, diethyl 4-methyl-7-coumarinyl phosphorothionate, diethyl p-nitrophenyl phosphate, ethyl p-nitrophenyl phenyl-phosphonothionate.

7. To be labelled with the words “Caution. This preparation should be administered only under medical supervision. The vapour is dangerous”—medicines made up ready for the internal or external treatment of human ailments and containing di-isopropyl fluorophosphonate.

8. To be labelled with the words “Caution. This may cause drowsiness”—

Anti-histamine substances, the following; their salts; their molecular compounds—

Antazoline.

SCHEDULE V*continued*

Bromodiphenhydramine.
Chlorcyclizine.
(p-Chlorophenylpyrid-2-ylmethyl) 2-dimethylaminoethyl ether 1-(4-p-Chlorophenyl-3-phenyl-but-2-enyl)-pyrrolidine.
Chlorpheniramine.
Cyclizine.
Di-n-butylaminomethyl-4:5:6-trihydroxyphthalide.
1 -Dimethylamino-3-phenyl-3-(2-pyridyl)-propane.
Diphenhydramine.
Substances being tetra-substituted N derivatives of ethylenediamine or propylenediamine.

SCHEDULE VI

[Rule 9(1)(b).]

STATEMENT OF PARTICULARS PERMITTED IN CERTAIN CASES AS TO PROPORTION OF POISON

Name of poison	*Particulars*
1. Alkaloids.	
2. Aconite, alkaloids of	The proportion of any one alkaloid of aconite that the preparation would be calculated to contain on the assumption that all the alkaloids of aconite in the preparation were that alkaloid.
3. Belladonna, alkaloids of	The same as above, with the substitution for the reference to aconite of a reference to belladonna, calabar bean or such other of the said poisons as the case may require.
4. Calabar bean, alkaloids of	
5. Coca, alkaloids of	
6. Ephedra, alkaloids of	
7. Ergot, alkaloids of	
8. Gelsemium, alkaloids of	
9. Jaborandi, alkaloids of	
10. Lobelia, alkaloids of	
11. Pomegranate, alkaloids of	

SCHEDULE VI—*continued*

Name of poison	*Particulars*
12. Quebracho, alkaloids of, other than the alkaloids of red quebracho	
13. Sabadilla, alkaloids of	
14. Solanaceous alkaloids not otherwise included in the Poisons List	
15. Stavesacre, alkaloids of	
16. Veratrum, alkaloids of	
17. Yohimba, alkaloids of	
18. Colchicum, alkaloids of	
19. Antimonial poisons	The proportion of antimony trioxide (Sb2 O3) or antimony pentoxide (Sb2 O5) that the preparation would be calculated to contain on the assumption that the antimony (Sb) in the poison had been wholly converted into antimony trioxide or antimony pentoxide as the case may be.
20. Arsenical poisons	The proportion of arsenic trioxide (As2 O3) or arsenic pentoxide (As2 O5) that the preparation would be calculated to contain on the assumption that the arsenic (As) in the poison had been wholly converted into arsenic trioxide or arsenic pentoxide as the case may be.
21. Barium, salts of	The proportion of one particular barium salt which the preparation would be calculated to contain on the assumption that the barium (Ba) in the poison had been wholly converted into that salt.
22. Digitalis, glycosides of; other active principles of digitalis	The number of units of activity as defined in the *British Pharmacopoeia* contained in a specified quantity of the preparation.
23. Hydrocyanic acid; cyanides, double cyanides of mercury and zinc	The proportion of hydrocyanic acid (HCN) that the preparation would be calculated to contain on the assumption that the cyanides in the poison had been wholly converted into hydrocyanic acid.
24. Insulin	The number of units of activity as defined in the *British Pharmacopoeia* contained in a specified quantity of the preparation.
25. Lead, compounds of, with acids from fixed oils	The proportion of lead oxide (Pbo) that the preparation would be calculated to contain on the assumption that the lead in the poison had been wholly converted into lead ozide.
26. Mercury, organic compounds of	The proportion of organically combined mercury (Hg) contained in the preparation.
27. Nux vomica	The proportion of strychnine contained in the preparation.
28. Opium	The proportion of morphine contained in the preparation.
29. Phenols	The proportion of phenols (added together) contained in the preparation.

SCHEDULE VI—*continued*

Name of poison	*Particulars*
30. Compounds of a phenols with a metal	The proportion of phenols (added together) that the preparation would be calculated to contain on the assumption that the compounds of phenols with a metal had been wholly converted into the corresponding phenols.
31. Pituitary gland, the active principles of	Either— (a) the number of units of activity as defined in the *British Pharmacopoeia* contained in a specified quantity of the preparation; or (b) the proportion of pituitary gland, or of anterior or of posterior lobe of the gland, as the case may be, contained in the preparation; or (c) the amount of pituitary gland or of anterior or of posterior lobe of the gland, as the case may be, from which a specified quantity of the preparation was obtained, together with an indication whether the amount relates to fresh or to dried gland substance.
32. Potassium hydroxide	The proportion of potassium monoxide (K2O) which the preparation would be calculated to contain on the assumption that the potassium hydroxide in the preparation had been wholly converted into potassium monoxide.
33. Sodium hydroxide	The proportion of sodium monoxide (Na20) which the preparation would be calculated to contain on the assumption that the sodium hydroxide in the preparation had been wholly converted into sodium monoxide.
34. Strophanthus, glycosides of	The amount of Standard Tincture of Strophanthus as defined in the *British Pharmacopoeia* which possesses the same activity as a specified quantity of the preparation when assayed by the method described in the said *Pharmacopoeia*.
35. Suprarenal gland, the active principles of their salts	Either— (a) the proportion of suprarenal gland or of the cortex or of the medulla of the gland, as the case may be, contained in the preparation; or (b) the amount of suprarenal gland or of the cortex or of the medulla of the gland, as the case may be, from which a specified quantity of the preparation was obtained, together with an indication whether the amount relates to fresh or to dried gland substance.
36. Thyroid gland, the active principles of their salts	Either— (a) the proportion of thyroid gland contained in the preparation; or

SCHEDULE VI—*continued*

Name of poison	*Particulars*
	(b) the amount of thyroid gland from which a specified quantity of the preparation was obtained, together with an indication whether the amount relates to fresh or to dried gland.

SCHEDULE VII

[Rule 15.]

POISONS REQUIRED TO BE SPECIALLY LABELLED FOR TRANSPORT

1. Arsenical poisons.
2. Barium, salts of.
3. Dinitrocresols (DNC), their compounds with a metal or a base when contained in preparations for use in agriculture or horticulture, except winter washes containing not more than the equivalent of 5 per cent of dinitrocresols.
4. Dinitrophenols when contained in preparations for use in agriculture or horticulture.
5. Dinosam, its compounds with a metal or a base, when contained in preparations for use in agriculture or horticulture.
6. Dinoseb, its compounds with a metal or base, when contained in preparations for use in agriculture or horticulture.

6A. Endosulfan.

7. Fluoroacetamide; Fluoroacetanilide.
8. Hydrocyanic acid; cyanides.
9. Nicotine.
10. Phosphorus compounds, the following—

    Diethyl thiophosphate of ethyl-mercapto-ethanol, dimefox, ethyl-paranitro-phenyl-benzene thiophosphonate, hexaethyl tetraphosphate (HETP), 4-methyl-hydroxy-coumarin-diethyl thiophosphate, mipafox, paranitrophenyl-diethyl phosphate, parathion, schradan, tetraethyl pyrophosphate (TEPP), triphosphoric pentadimethylamide, di-isopropyl-fluorophenate, demeton, mazidox, methyl demeton, sulphotepp, amiton, demeton-O, demeton-S, demeton-O-methyl, demeton-S-methyl, diethyl 4-methyl-7-coumarinyl phosphorothionate, diethyl p-nitrophenyl phosphate, ethyl p-nitrophenyl phenylphosphonothionate, ethion, mecarbam, phenkapton.

11. Strychnine.
12. Thallium, salts of.

SCHEDULE VIII

[L.N. 365/1964, L.N. 41/1971, r. 20, L.N. 61/2002, s. 2, L.N. 91/2004, s. 2.]

FORMS

1. Application for registration as a pharmacist (section 7).
2. Register of pharmacists (section 6).
3. Certificate of registration as a pharmacist (section 9).
4. Application for registration of premises (section 23).
5. Register of premises (section 23).
6. Application for wholesale dealer's licence (section 27).

SCHEDULE VIII—*continued*

7. Wholesale dealer's licence (section 27).
8. Register of wholesale dealer's licences (section 27).
9. Application for licence to deal in poisons for mining, agricultural and horticultural purposes (section 28).
10. Licence to deal in poisons for mining, agricultural and horticultural purposes (section 28).
11. Register of dealers in mining, agricultural and horticultural poisons (section 28).
12. Certificate for purchase of poison (section 29).
13. Application for licence to sell Part II poisons (section 32).
14. Licence to sell Part II poisons (section 32).
15. Register of licences issued to sellers of Part II poisons (section 32).
16. Poisons Book (section 30).
17. Permit to import Part I poisons (rule 3).
18. Application for pharmaceutical representative's permit (rule 13A).
19. Pharmaceutical representative's permit (rule 13A).
20. Institution inspection report (rule 14).
21. Application for licence to manufacture drugs for sale (rule 16).
22. Licence to manufacture drugs for sale (rule 16).
23. Application for licence for the exportation of drugs and poisons.
24. Annual professional practise licence as a pharmacist (section 9A).
25. Roll of Pharmaceutical Technologists (section 6(2)).
26. Application for enrolment as a pharmaceutical technologist (section 7(2)).
27. Application for annual practice licence for a pharmacist (section 9A).
28. Certificate of enrolment as a pharmaceutical technologist (section 9(2)).
29. Application for licence as a pharmaceutical technologist (section 20(1A)).
30. Application for registration of premises for a pharmaceutical technologist (section 20(1A)).
31. Certificate for registration of premises for a pharmaceutical technologist (section 20(1A)).
32. Annual licence to practice as a pharmaceutical technologist (section 20(2A).
33. Certificate of registration of premises for pharmacist (section 23(c)).

FORM 1

APPLICATION FOR REGISTRATION AS A PHARMACIST

The Registrar, Pharmacy and Poisons Board,
Afya House, P.O. Box 30016, Nairobi.

I, ... of .....
hereby make application for registration as a pharmacist.

I hereby declare that to the best of my knowledge and belief I am not aware of any circumstances which would disqualify me for registration.

My qualifications are .....
.....

I enclose the following certificates/diplomas—
.....
.....

Date ....
Signature

SCHEDULE VIII—*continued*

FORM 2

REGISTER OF PHARMACISTS

REGISTRATION		Name of Applicant	Address	Qualification	Date of Qualification	Registration Fee
No.	Date					

FORM 3

CERTIFICATE OF REGISTRATION AS A PHARMACIST

.....

is hereby registered as a pharmacist in accordance with the provisions of Part II of the Pharmacy and Poisons Act.

Given at Nairobi on the ....., 20.....

.....

Registrar, Pharmacy and Poisons Board

FORM 4

APPLICATION FOR REGISTRATION OF PREMISES

The Registrar, Pharmacy and Poisons Board,

Afya House, P.O. Box 30016, Nairobi.

In accordance with the provisions of section 23 of the Pharmacy and Poisons Act, I/We .....

.....

wishing to carry on the business of a pharmacist, do hereby apply for registration of premises situated at .....

in the town of .....

The business, in so far as concerns the retail sale of drugs, will be under the control of .....

..... a pharmacist registered in accordance with Part II of the Act.

Date ....

Signature of Applicant

N.B.—Any change of pharmacist under whose control the business is carried on must be notified to the Registrar within seven days.

Fee: Sh. 100.

SCHEDULE VIII—*continued*

FORM 5

REGISTER OF PREMISES

REGISTRATION		Name(s) of owner(s) of the business	Address of premises where business of a pharmacist is carried on (give name of minor settlement/town)	Name of pharmacist under whose control the business of a pharmacist is carried on
No.	Date			

FORM 6

APPLICATION FOR WHOLESALE DEALER'S LICENCE

The Registrar, Pharmacy and Poisons Board,
Medical Headquarters, P.O. Box 30016, Nairobi.

I/We ..... of .....
wishing to carry on business as a wholesale dealer in poisons at .....
..... in the town of .....
hereby apply for the issue/renewal of a wholesale dealer's licence.

The registered pharmacist in control of the distribution of poisons is .....
....., resident in .....

Date ....

Signature of Applicant

N.B.—Any change of registered pharmacist under whose control the distribution of poisons is effected must be notified to the Registrar within seven days.

FORM 7

WHOLESALE DEALER'S LICENCE

Messrs ..... of ....., carrying on business at ..... are hereby authorised to sell poisons by way of wholesale dealing.

Date .....

.....

Registrar, Pharmacy and Poisons Board

Note.—This licence expires on the 31st day of December, 20.....
Fee: Sh. 400.

SCHEDULE VIII—*continued*

FORM 8

REGISTER OF WHOLESALE DEALERS

REGISTRATION		Name(s) of owner(s) of the business	Address of premises where business is carried on	Name of pharmacist in control of the distribution of poisons
No.	Date			

FORM 9

APPLICATION FOR LICENCE TO DEAL IN POISONS FOR MINING, AGRICULTURAL AND HORTICULTURAL PURPOSES

The Registrar, Pharmacy and Poisons Board,
Medical Headquarters, P.O. Box 30016, Nairobi.

I/We ..... of .....
carrying on a regular business in *mining/agricultural/and/or horticultural accessories at .....
..... in the town of ....., hereby apply for the issue/renewal of a licence to deal in the following poisons .....
.....
.....

I/We hereby nominate the following person(s) .....
.....
.....
who may sell in accordance with the provisions of rule 10 of the Pharmacy and Poisons Rules.

Date .....

.....
Signature of Applicant

* Delete as necessary.

Note.—Not more than two persons may be nominated.

FORM 10

LICENCE TO DEAL IN MINING, AGRICULTURAL OR HORTICULTURAL POISONS

Messrs. ..... of .....
carrying on business at ..... are hereby licensed to deal in the following poisons .....
.....
.....
.....

SCHEDULE VIII, Form 10—*continued*

The following person(s) are hereby authorised to sell these poisons in accordance with the provisions of rule 10 of the Pharmacy and Poisons Rules.

.....

.....

Date .....

.....

Registrar, Pharmacy and Poisons Board

N.B.—Any change in persons authorised to sell must be notified to the Registrar within seven days.

Note.—This licence expires on the 31st day of December, 20.....

Fee: Sh. 50.

FORM 11

REGISTER OF DEALERS IN MINING, AGRICULTURAL AND HORTICULTURAL POISONS

REGISTRATION		Name of owner(s) of the business	Address of premises where business is carried on	Name(s) of person(s) authorised to sell poisons
No.	Date			

FORM 12

CERTIFICATE FOR PURCHASE OF POISON

For the purposes of paragraph (b) of subsection (2) of section 29 of the Pharmacy and Poisons Act, I, the undersigned, hereby certify from my knowledge of (a) .... of (b) ....., that he is a person to whom (c) ..... may properly be supplied.

I further certify that (d) ..... is the signature of the said (a) .....

Date ....

Signature and designation of officer giving certificate

(a) Insert full name of intending purchaser.

(b) Insert full postal address.

(c) Insert name of poison.

(d) Intending purchaser to sign his name here.

SCHEDULE VIII—*continued*

FORM 13

APPLICATION FOR LICENCE TO SELL PART II POISONS

To the Civil Secretary,

.....

I/We ..... , being engaged in the business of ..... ,

hereby apply to sell poisons by wholesale/retail in Group A/Group B of Part II of the Poisons List or specified poisons, on the following premises .....

.....

I/We hereby nominate the following person(s) .....

.....

.....

who will sell such poisons in accordance with the provisions of the Pharmacy and Poisons Act and the Pharmacy and Poisons Rules.

Date..... .....

Signature of Applicant

FORM 14

LICENCE TO SELL PART II POISONS

....., of ..... ,

carrying on the business .....

at ..... , is hereby licensed to sell the following

Part II Poisons .....

Insert here either .....

Group A,

Group B, or .....

specified poisons .....

as the case may .....

be and whether .....

by wholesale or .....

retail sale. .....

.....

The following person(s) are hereby authorised to sell these poisons in accordance with the provisions of the Pharmacy and Poisons Rules.

.....

.....

Date ....

Civil Secretary

N.B.—Any change of persons authorised to sell must be notified to the Civil Secretary within seven days.

Note.—This licence expires on 31st December, 20 .....

Fee: Sh. 40.

SCHEDULE VIII—*continued*

FORM 15

REGISTER OF LICENCES ISSUED TO SELLERS OF PART II POISONS

REGISTRATION		Name of licensee	Address of premises where business is carried on	Class of licence	Name(*s*) of person(*s*) authorised to sell poisons
No.	Date				

FORM 16

POISONS BOOK

Date of sale	Name and quantity of poison supplied	PURCHASER'S			Purpose for which stated to be required	Date of certificate (if any)	Name and address of person giving certificate (if any)	Signature of purchaser, or where a signed order is permitted the date of the signed order
		Name	Address	Business trade or occupation				

FORM 17

MINISTRY OF HEALTH AND CENTRAL BANK OF KENYA

FOR EXCHANGE CONTROL USE No.

APPLICATION FOR IMPORT AND/OR FOREIGN EXCHANGE ALLOCATION

IMPORTER'S FULL NAME AND ADDRESS:	NOTE.—Applicant to attach sellers Proforma Invoice. Proforma Invoice No. Date: ..... Reference: .....
IMPORTER'S BANK AND BRANCH:	TOTAL AMOUNT IN FOREIGN CURRENCY: In Figures: In Words: Exchange rates: Kenya Currency equivalent Sh.
SELLER'S FULL NAME AND ADDRESS:	APPLICABLE SCHEDULE:
	COUNTRY OF ORIGIN:

SCHEDULE VIII—*continued*

Date of Shipment:			Terms of Payment (State commission rate if applicable):			
Mode of Transport Port of Loading			F.O.B.	Freight		Insurance
Port of Discharge:						
S.I.T.C. Code	Generic Name	Trade Name	Package Size	Quantity	Reg. No.	Unit price

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

FOR OFFICIAL USE OF MINISTRY OF HEALTH

....

Valid up to Replacement

.....

Extended to

FOR USE OF CENTRAL BANK OF KENYA

.....

Replacement

.....

Exchange Control Authorization Stamp and Signature

SPECIAL INSTRUCTIONS:

Approved subject to clean report of finding by general superintendence company limited as to: quality and quantity inspection and Price comparison:

To be contacted at:

FOR USE BY REMITTING BANK

PAYMENTS MADE

Date	Foreign Currency Remitted	Exchange Rate	Kenya Currency Equivalent	Branch Stamp and Authorized Signature

SCHEDULE VIII—*continued*

FORM 18

APPLICATION FOR PHARMACEUTICAL REPRESENTATIVE'S PERMIT

I/We .....,
of (postal address) .....,
being engaged in the sale and supply of pharmaceutical goods, hereby make application that our representative Mr. .....
..... be permitted to possess pharmaceutical goods containing Part I poisons as scheduled below, for the purpose of giving free samples to persons who may lawfully possess such goods.

SCHEDULE

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

Date ....
(Signature of Applicant)

FORM 19

PHARMACEUTICAL REPRESENTATIVE'S PERMIT

Mr. ..... as representative
of ..... is hereby permitted to possess and supply free samples of pharmaceutical goods containing Part I Poisons, as scheduled below, to persons who are authorized to use them in their trade, business or profession as laid down in the Pharmacy and Poisons Act, subject to maintenance of records as required by rule 13A(1)(b) of the Pharmacy and Poisons Rules.

SCHEDULE

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

Date ....

The Pharmacy and Poisons Board,
P.O. Box 30016,
Nairobi.

Note.—This permit expires on 31st December, 20 ....., or upon the person named ceasing to be employed as a pharmaceutical representative of the firm stated above.

FEE: Sh. 25.

FORM 20

INSTITUTION INSPECTION REPORT

I, the undersigned of (postal address) .....,
have today carried out an inspection of .....
as required by rule 14 of the Pharmacy and Poisons Rules.

SCHEDULE VIII, Form 20—*continued*

The following defects are reported—

1. Storage .....
.....
.....
2. Methods of Handling .....
.....
.....
3. Records .....
.....
.....

I have the following recommendations to make—

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

The previous inspection was carried out on .....

Signature .....

Designation .....

Date .....

To: 1. ..... (person in charge of the Institution).

2. The Registrar, Pharmacy and Poisons Board.

FORM 21

APPLICATION FOR A LICENCE TO MANUFACTURE DRUGS FOR SALE

The Registrar, The Pharmacy and Poisons Board .....
.....

I/We .....,

of (postal address) .....,

having premises situated at .....

and being engaged in the business of .....

hereby apply to manufacture for sale the following drug(s) medicine(s) .....
.....

This/These drug(s)/medicine(s) has/have the following composition .....
.....

The manufacture of the above drug(s)/medicine(s) will be carried out under the direct personal supervision of .....
..... who has

the following qualifications .....

The manufacture of the above drug(s)/medicine(s) will be carried out at .....
.....

Date ....

(Signature of Applicant)

SCHEDULE VIII, Form 21—*continued*

Note.—Any change of the person under whose direct personal supervision the manufacture is carried out, whether temporary or permanent, must be notified to the Registrar immediately.

FORM 22

LICENCE TO MANUFACTURE DRUGS FOR SALE

.....

of (postal address) .....

and having premises situated at .....

is hereby licensed to manufacture for sale the following drug(s)/medicine(s).....

under the direct personal supervision of .....

at .....

Note—The licence expires on 31st December, 20.....

Registration No. .....

Date..... .....

Registrar,
Pharmacy and Poisons Board,
P.O. Box 30016,
Nairobi.

Any change of the person under whose direct personal supervision the manufacture is carried on, whether temporary or permanent, must be notified to the Registrar immediately.

FORM 23

MINISTRY OF HEALTH

APPLICATION FOR LICENCE FOR THE EXPORTATION OF DRUGS AND POISONS

<table>
<tr><td colspan="7">EXPORTER'S NAME AND ADDRESS CODE No.</td></tr>
<tr><td colspan="3">CONSIGNEE'S NAME AND ADDRESS:</td><td colspan="4">INVOICE No.:</td></tr>
<tr><td colspan="3" rowspan="2"></td><td colspan="4">CD3 No.</td></tr>
<tr><td colspan="4">Country of origin: Destination of goods:</td></tr>
<tr><td colspan="3">DATE OF SHIPMENT:</td><td colspan="4" rowspan="2">Terms of delivery and payments:

F.O.B. Value:</td></tr>
<tr><td colspan="3">Mode of transport: Port of loading
Port of discharge:</td></tr>
<tr><td>Generic Name</td><td>Trade Name</td><td>Pack Size</td><td>Unit Price</td><td>Quantity</td><td>Batch No.</td><td>Country of Manufacture</td></tr>
<tr><td></td><td></td><td></td><td></td><td></td><td></td><td></td></tr>
</table>

SCHEDULE VIII, Form 23—*continued*

I declare that the particulars which I have given are true and accurate to the best of my knowledge and belief.

Date ..... Signed .....

Applicant

This document will be effective as an Export Licence only when it has been validated by the Chief Pharmacist.

FOR OFFICIAL USE ONLY: EXPORT LICENCE: NUMBER .....

Export of goods described above is approved, subject to .....

.....

Date ....

for Chief Pharmacist

This licence is not transferable.

FORM TO BE FULLY COMPLETED IN TRIPLICATE (PREFERABLY TYPEWRITTEN) BY APPLICANT:

PHARMACY AND POISONS ACT (CAP. 244) RULE .....

FORM 24 **(L.N. 61/2002, s. 2.)**

ANNUAL PROFESSIONAL PRACTICE LICENCE AS A PHARMACIST

Serial No. .....

Prof./Dr. .....

(Full names in block letters)

is hereby licensed by the Pharmacy and Poisons Board to render pharmaceutical services in Kenya.

Dated the ..... day of ..... 20 .....

.....

Registrar, Pharmacy and Poisons Board

Licence No. .....

This Licence expires on 31st December, 20 .....

Fee: KSh. 2,500

SCHEDULE VIII—*continued*

FORM 25

ROLL OF PHARMACEUTICAL TECHNOLOGISTS

ENROLMENT		NAME OF APPLICANT	ID No.	ADDRESS	QUALIFICATION	DATE OF QUALIFICATION	TRAINING INSTITUTION

SCHEDULE VIII—*continued*

FORM 26

APPLICATION FOR ENROLMENT AS A PHARMACEUTICAL TECHNOLOGIST

The Registrar,
Pharmacy and Poisons Board,
P.O. Box 27663-00506,
Nairobi

I .....

of P.O. Box .....

.....

ID No. ..... do hereby apply to be enrolled as a Pharmaceutical Technologist in accordance with the Pharmacy and Poisons Act.

Qualification .....

Institution .....

.....

Date of Qualification .....

Period of Internship: From ..... to .....

(Attach proof of Internship*)

.....

Signature of Applicant

* Applicants are advised to attach genuine evidence from recognized institution of attachment. Any false information given may lead to prosecution.

FORM 27

APPLICATION FOR ANNUAL PRACTICE LICENCE FOR A PHARMACIST

The Registrar,
Pharmacy and Poisons Board,
P.O. Box 27663-00506,
Nairobi

I .....

of P.O. Box .....

.....

Registration No. ..... do hereby apply for a Practice licence as a pharmacist.

.....

Date

.....

Signature of Applicant

SCHEDULE VIII—*continued*

FORM 28

SERIAL No. .....

CERTIFICATE OF ENROLMENT AS A PHARMACEUTICAL TECHNOLOGIST

The Registrar,
Pharmacy and Poisons Board,
P.O. Box 27663-00506,
Nairobi

.....

(Name and Address)

ID/No. .....

Having duly satisfied the Pharmacy and Poisons Board is hereby enrolled as a Pharmaceutical Technologist in accordance with the Pharmacy and Poisons Act.

Given on the ..... day of ..... in the year 20 .....

Enrolment No. .....

.....

(Registrar, Pharmacy and Poisons Board)

Fee: KSh. 500

FORM 29

APPLICATION FOR LICENCE AS PHARMACEUTICAL TECHNOLOGIST

The Registrar,
Pharmacy and Poisons Board,
P.O. Box 27663-00506,
Nairobi

Dear Sir/Madam

I, ...

of P.O. Box .....

ID/No. ..... do hereby apply for a licence as a pharmaceutical technologist.

Enrolment No. ..... Date of enrolment .....

Name of premises .....

Plot No. ..... Road .....

Town .....

.....

Signature of Applicant

.....

Date

SCHEDULE VIII—*continued*

FORM 30

APPLICATION FOR REGISTRATION OF PREMISES FOR A PHARMACEUTICAL TECHNOLOGIST

The Registrar,
Pharmacy and Poisons Board,
P.O. Box 27663–00506,
Nairobi

I/We .....

.....

wishing to carry on the business of a Pharmaceutical Technologist, do hereby apply for registration of premises situated at .....

.....

in the township of .....

The business in so far as concerns the retail sale of drugs will be under the control of .....

.....

..... a Pharmaceutical Technologist enrolled in accordance with Part II of the Act.

Date ....

Signature of the Applicant

Note.—Any change of premises of a Pharmaceutical Technologist under whose control the business is carried on must be notified to the Registrar within seven days.

FORM 31

MINISTRY OF HEALTH

PHARMACY AND POISONS BOARD

PREMISES REGISTRATION CERTIFICATE FOR PHARMACEUTICAL TECHNOLOGIST'S PRACTICE

SERIAL No. .....

Name of Premises .....

Registration No. of premises .....

Location of premises .....

Town ..... Street .....

Plot No. .....

Name of pharmaceutical technologist .....

ID No. ..... Enrolment No. .....

Has met the necessary conditions for the business of a pharmaceutical technologist to be carried therein.

.....

(Registrar, Pharmacy and Poisons Board)

.....

Date

SCHEDULE VIII , Form 31—*continued*

Note: (a) This registration expires on 31st December, 20 .....

(b) No change of premises is permitted without the authority of the Board.

(c) This registration shall become void upon expiration of 30 days from any change of ownership of the business.

Fee: KSh. 5,000

FORM 32

SERIAL NO. .....

MINISTRY OF HEALTH

PHARMACY AND POISONS BOARD

ANNUAL LICENCE TO PRACTICE AS A PHARMACEUTICAL TECHNOLOGIST

.....

(Name and Address)

is hereby licensed to practice as a pharmaceutical technologist in accordance with the Pharmacy and Poisons Act.

Name of Premises .....

Plot No. ..... Road .....

Town .....

Given at Nairobi on the ..... day of ..... of the year 20 .....

.....

(Registrar, Pharmacy and Poisons Board)

.....

Date

This licence expires on the 31st December, 20 .....

Fee: KSh. 2,500

FORM 33

SERIAL No.....

CERTIFICATE FOR REGISTRATION OF PREMISES

Messrs. .....

of .....

Plot No. ..... is registered to carry on business of a pharmacist as provided for by section 23.

Date .....

.....

Registrar, Pharmacy and Poisons Board.

SCHEDULE VIII—*continued*

Note: (a) This registration expires on 31st December, 20 .....

(b) No change of premises is permitted without the authority of the Board.

(c) This registration shall become void upon expiration of 30 days from any change of ownership of the business.

Fee: KSh. 5,000

SCHEDULE IX

[Rule 6.]

PERMIT AUTHORISING FARMERS AND OTHER PERSONS TO BE IN POSSESSION OF SUBSTANCES SPECIFIED IN GROUP II OF SCHEDULE IV TO THE RULES

For the purposes of rule 6 of the Pharmacy and Poisons Rules, I, the undersigned, of .....

.....

hereby authorise .....

of ..... to purchase and possess the following substances in Group II of Schedule IV to the Rules—

.....

.....

.....

.....

.....

.....

.....

1. If any quantity is specified against any or all of the items listed above the permit holder may not purchase or possess more than that quantity at any time.
2. This permit is valid for a period of six months from date of issue.
3. This permit must be produced to the authorised seller of poisons on each occasion when supplies are purchased.

Date ....

Signature
