

## FOOD AND DRUGS ACT

### ARRANGEMENT OF SECTIONS

#### *Food, drugs, cosmetics and devices*

#### SECTION

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FOOD AND DRUGS  
ACT

**An Act to make provision for the regulation of the manufacture, sale and advertisement of food, drugs, cosmetics and devices and repeal the existing State laws on those matters.**

*[10th February,  
1976]*

[Commencement.]

*Food, drugs, cosmetics and devices*

**1. Prohibition of sale of certain food, drugs, cosmetic and devices**

(1) No person shall sell, import, manufacture or store any article of food which-  
[1999 No. 21.]

- (a) has in it or upon it any poisonous or harmful substance not being a food additive or contaminant of a type and within the level permitted by regulations made under this Act;
- (b) is unfit for human consumption; or
- (c) consists in whole or in part of any filthy, disgusting, rotten or diseased substance.

(2) No person shall sell, import, manufacture or store any article of food or any drug which is adulterated.

[1999 No. 21.]

(3) No person shall sell, import, manufacture or store any article of food or any drug or cosmetic which was manufactured, prepared, preserved, packaged, or stored under insanitary conditions.

[1999 No. 21.]

(4) No person shall sell, import, manufacture or store any cosmetic which-

- (a) has in it or upon it any substance which may cause injury to the health of the user thereof when the cosmetic is used-
  - (i) according to the directions on the label or otherwise accompanying the cosmetic; or
  - (ii) for such purposes and by such methods of use as are customary or usual therefor; or
- (b) consists wholly or in part of any filthy or decomposed substance or of any foreign matter.

[1999 No. 21.]

(5) No person shall sell, import, manufacture or store any device which, when used according to the directions on the label or otherwise accompanying the device, or under such conditions as are customary or usual therefor, may cause injury to the user thereof.

[1999 No. 21.]

## **2. Prohibition of sale or advertisement of food, etc., as treatment, etc., for certain diseases**

Except as otherwise provided by regulations, no person shall-

- (a) advertise to the general public any food, drug, cosmetic or device as treatment, preventative or cure for any of the diseases, disorders or abnormal physical states specified in the First Schedule to this Act; or  
[First Schedule.]
- (b) sell any food, drug, cosmetic or device that is represented on the label or is advertised to the general public as a treatment, preventative or cure for any of the diseases, disorders or abnormal physical states specified in the said First Schedule to this Act.

[First Schedule.]

## **3. Prohibition of importation, exportation, distribution, etc., of specified drugs**

Except as authorised by regulations, no person shall import into Nigeria or export therefrom, manufacture, sell, distribute or cause to be distributed (whether as samples or otherwise) any of the drugs specified in the Second Schedule to this Act.

## **4. Power of Minister to obtain particulars in respect of certain substances**

- (1) Where any person carries on any business in the course of which any substance-
  - (a) is used in the preparation of any food, drug or cosmetic; or
  - (b) is imported, produced or sold for use in the preparation of any food, drug or cosmetic,

the Minister or any person authorised by the Minister in that behalf may serve on that person a notice requiring him to furnish to the Minister within such time as may be specified in the notice such particulars as may be so specified in respect of every substance which, being a substance or a substance of a class specified in the notice, is in the course of that business used as mentioned in paragraph (a) of this subsection or imported, produced or sold for use as mentioned in paragraph (b) of this subsection.

(2) Without prejudice to the generality of subsection (1) of this section, a notice under that subsection may in particular require the furnishing of the following particulars of any substance to which it applies, that is to say-

- (a) particulars of the composition of the substance and the chemical formula of every ingredient thereof;
- (b) particulars of the manner in which the substance is used or intended to be used in the preparation of any food, drug or cosmetic;
- (c) particulars of-
  - (i) any investigations (and over what period of time) carried out by or on behalf and to the knowledge of the person carrying on the business for the purpose of determining whether or not the substance or any product produced when the substance is used as mentioned in paragraph (b) of this subsection is injurious to or otherwise affects health, and the result of any such investigations;
  - (ii) any investigations or inquiries (and over what period or time) carried out by or on behalf and to the knowledge of the person carrying on the business for the purpose of determining the cumulative effect on the health of any person consuming in ordinary quantities that substance or any product produced when the substance is used as mentioned in paragraph (b) of this subsection.

(3) No person shall disclose any information supplied to the Minister in pursuance of a notice under subsection (1) of this section except-

- (a) with the written consent of the person who supplied the information; or
- (b) in accordance with the directions of the Minister; or
- (c) for the purposes of any proceedings under this Act.

## **5. Prohibition of various misleading practices**

No person shall-

- (a) label, package, treat, process, sell or advertise any food, drug, cosmetic or device in a manner that is false or misleading or is likely to create a wrong impression as to its quality, character, value, composition, merit or safety;
- (b) where a standard has been prescribed for any food, drug, cosmetic or device; label, package, sell or advertise any substance or article in such a manner that the substance or article is likely to be mistaken for that food, drug, cosmetic or device unless the substance or article complies with the prescribed standard;

- (c) where a standard for a drug is contained in any of the publications specified in the Third Schedule to this Act, label, package, sell or advertise any substance which is not of the published standard in a manner likely to cause it to be mistaken for a drug of the published standard;  
[Third Schedule.]
- (d) in the case of drug for which no standard has been prescribed under the regulations or in any of the publications specified in the Third Schedule to this Act-
  - (i) sell that drug in any manner which is likely to deceive or mislead a purchaser into thinking that the drug conforms to a standard prescribed as aforesaid; or
  - (ii) sell that drug as complying with some other standard unless the drug complies with the professed standard under which it is sold.

#### **6. Manufacture, etc., of food, etc., under unsanitary conditions**

No person shall under unsanitary conditions manufacture, prepare, preserve, package or store for the purpose of selling, any food, drug or cosmetic.

#### **7. Minister's certificate for manufacture, etc., of drugs specified in the Fourth or Fifth Schedule**

(1) No person shall manufacture for sale any drug specified in the Fourth Schedule to this Act without first obtaining, in accordance with the regulations, a certificate of the Minister to the effect that premises in which the drug intended to be manufacture and the process and conditions by and under which the manufacture is to be carried on are in the opinion of the Minister suitable for ensuring that the drug will be safe for use.

[Fourth Schedule.]

(2) No person shall sell any drug specified in the Fifth Schedule to this Act without first obtaining in accordance with the regulations a certificate of the Minister that the batch from which the drug was taken is safe for use.

[Fifth Schedule.]

(3) Except as provided in the regulations no person shall distribute or cause to be used as samples any of the drugs listed in the Fourth or Fifth Schedule to this Act.

[Fourth or Fifth Schedule.]

#### **8. Declaration by manufacturer**

(1) The Minister or any person authorised by him in that behalf may order the manufacture of any article of food, drug, cosmetic or device to furnish a declaration in the prescribed form that the article in question was manufactured in accordance with the provisions of this Act and the regulations, and it shall be the duty of the manufacturer to comply with the requirements of the order.

(2) Except as provided by the regulations, no article of food, drug, cosmetic or device shall be imported or otherwise brought into Nigeria unless-

- (a) it is accompanied by a certificate from the manufacturer to the effect that it was manufactured in accordance with any existing standard or code of practice pertaining to such product or, where such standard or code of practice does not

exist for the particular product, in accordance with any international standard laid down, in the case of food, under the directive of the Codex Alimentarius Commission; and

- (b) a certificate issued by or on behalf of the Government of the country where it was manufactured to the effect that its sale in that country would not constitute a contravention of the law of that country.

(3) Notwithstanding anything contained in the foregoing provisions of this section, the Minister may by regulations provide that any such article as is mentioned in subsection (2) of this section shall be imported into Nigeria unless it was manufactured in accordance with any standards that may be specified in the regulations.

#### **9. Designation of inspecting officers and analysts**

(1) The Minister may, on the recommendation of the Food and Drugs Advisory Council, and subject to the provisions of this section, designate-

- (a) as a food and drug analyst; or
- (b) as a drug analyst; or
- (c) as a food and drug inspection officer,

any person (whether or not a member of the public service of the Federation) who possesses such qualifications and fulfil such other requirements, if any, as are prescribed under this section and shall furnish to every person so designated a certificate of designation.

(2) A person shall not be designated a food and drug analyst unless he is a graduate in chemistry and-

- (a) has at his disposal such laboratory facilities as are, in the opinion of the Minister, adequate to enable him to discharge his functions properly under this Act;
- (b) produces evidence of post-graduate training or qualification in the analysis of food and drugs; and
- (c) has had not less than five years' continuous post-graduate relevant experience in the analysis of food and drugs in an approved laboratory.

(3) A person shall not be designated a drug analyst unless he holds a professional qualification in pharmacy acceptable to the Minister and-

- (a) has at his disposal such laboratory facilities as are, in the opinion of the Minister, adequate to enable him to discharge his functions properly under this Act;
- (b) produces evidence of post-qualification training or experience in the analysis of drugs; and
- (c) has had not less than five years' continuous post-qualification experience in the analysis of drugs in an approved laboratory.

#### **10. Powers of inspecting officers**

(1) An inspecting officer may, in the course of his duty, at any reasonable time and on production of his certificate of designation if so required-

- (a) enter (if need be by force) any premises in which he reasonably believes that any article to which this Act or the regulations apply is manufactured, prepared, preserved, packaged, stored or sold;
- (b) examine any article in the premises which appears to him to be an article to which this Act or the regulations apply or anything in the premises which he reasonably believes is used or is capable of being used for the manufacture, preparation, preservation, packaging, storage or sale of any such article;
- (c) take a sample or specimen of any article to which this Act or the regulations apply or which he has power to examine under paragraph (b) of this subsection;
- (d) open and examine, while on the premises, any container or package which he reasonably believes may contain anything to which this Act or the regulations apply or which may help him in his investigation;
- (e) examine any books, documents or other records found on the premises which he reasonably believes may contain any information relevant to the enforcement of this Act or the regulations and make copies thereof or extracts therefrom; and
- (f) seize and detain for such time as may be necessary for the purposes of this Act any article by any means of or in relation to which he reasonably believes any provision of this Act or the regulations has been contravened.

(2) In subsection (1) of this section, the expression "**article to which this Act or the regulations apply**" means-

- (a) any food, drug, cosmetic or device;
- (b) anything used for the manufacture, preservation, packaging or storing of any food, drug, cosmetic or device; and
- (c) any labelling or advertising material relating to or for use in connection with any food, drug, cosmetic or device,

but does not include live animals.

(3) The owner or person in charge of any premises entered by an inspecting officer pursuant to this section, and every person found therein, shall give all reasonable assistance in their power to the inspecting officer and shall make available to the inspecting officer all such information as the officer may reasonably require for the purposes of this Act.

(4) Any article seized under this Act shall be kept or stored in such a place as the inspecting officer may direct and shall be returned to the owner or the person from whom it was seized if the article, upon analysis or examination, is found to conform with the requirements of this Act and the regulations.

(5) Any article seized by an inspecting officer pursuant to this Act or the regulations may be submitted to an analyst for analysis or examination and the analyst, upon making

such analysis or examination, shall issue a certificate or report in the prescribed form setting forth the result of such analysis or examination and, without prejudice to the generality of section 19 (2) of this Act, the inspecting officer shall on demand deliver a copy of such certificate or report to the owner of the article if the article is to be the subject of proceedings under this Act.

(6) In this section, "**animals**" includes birds and fishes.

#### **11. False statement**

No person shall wilfully make or insert any false statement in any certificate or other document required by this Act or the regulations to be issued, made or furnished.

#### **12. Obstruction of inspecting officers**

No person shall-

- (a) obstruct or resist an inspecting officer in the execution of his duty under this Act or the regulations; or
- (b) make in any manner whatsoever to an inspecting officer acting in the course of his duties any statement which that person knows or has reasonable cause to believe to be false or misleading; or
- (c) without the authority of an inspecting officer remove, alter or interfere in any way with any HOLD LABEL tag or article seized under this Act.

[1999 No. 21.]

#### **13. Power of inspecting officers on importation of food, etc.**

(1) An inspecting officer shall have the right to examine any customs entries of any food, drug or cosmetic imported for use in Nigeria and for the purposes of analysis or examination thereof to take samples of any such food, drug, or cosmetic while still in any customs shed or government warehouse in Nigeria.

(2) Where sample are taken by an inspecting officer pursuant to subsection (1) of this section, the food, drug or cosmetic from which they are taken shall not be released to the importer except on production of an analyst's certificate or report to the effect that the food, drug or cosmetic complies with the requirements of this Act and the regulations.

(3) Where, in the course of his duties under this Act or the regulations an inspecting officer takes a sample of any food, drug or cosmetic for the purposes of analysis, he shall, in the presence of the owner or importer or any person in apparent control of the food, drug or cosmetic, seal the article in triplicate, one copy of which shall be sent to the analyst, the second retained by him and the third delivered to the owner, importer, or person in apparent control, as the case may be.

#### *Forfeiture*

#### **14. Forfeiture**

(1) Where the owner of an article seized under this Act consents to the forfeiture thereof the article shall thereupon be forfeited to the Minister.



(2) Where a person has been convicted of an offence under this Act or the regulations the court may order that the article by means of or in respect of which the offence was committed, and anything of a similar nature belonging to or in the possession of the person convicted or found with that article, be forfeited to the Minister.

(3) Without prejudice to the provisions of subsection (1) of this section, where any article has been seized under this Act, a judge of a Federal High Court may, upon application by an inspecting officer and after the giving of notice to such persons as the judge or magistrate may direct, order that the article and anything of a similar nature found therewith be forfeited to the Minister, if after hearing all the parties concerned he is of the opinion that the article is one by means of or in relation to which any of the provisions of this Act or the regulations is being or has been contravened.

(4) Where, whether in pursuant of an order under this section or otherwise, any article or thing is forfeited to the Minister under this section it shall vest in the Minister free from encumbrances, and the Minister may retain it or cause it to be destroyed or otherwise disposed of as the Minister thinks fit.

#### *Advisory Council*

### **15. Food and Drugs Advisory Council**

(1) The Minister may set up a Council to be known as the Food and Drugs Advisory Council, to assist and advise him in the preparation and review of regulations for carrying out the purposes and provisions of this Act and with respect to any other matters connected with this Act.

[1999 No. 21.]

(2) The said Council shall consist of such persons as the Minister may appoint, being persons who appear to the Minister to be suitable for appointment-

- (a) by reasons of their knowledge or experience of the matters to which this Act relates; or
- (b) as representing the interests of producers or distributors of food, drugs, cosmetics or devices; or
- (c) as representing the interests of consumers or users thereof.

#### *Regulations*

### **16. Regulations**

(1) The Minister may make regulations for carrying out the purposes and provisions of this Act and with respect to any other matter connected with this Act.

[1999 No. 21.]

(2) Without prejudice to the generality of subsection (1) of this section the Minister may make regulations-

- (a) for determining what constitutes the adulteration of any food or drug or class of food or drug and for determining what constitutes foreign matter in relation to any cosmetic or class of cosmetics;

- (b) prescribing the type and level of food additive or contaminant that may be present in any food offered for sale;
- (c) with respect to-
- (i) the labelling and packaging and the offering or exposing in any manner for sale of any food, drug, cosmetic or device;
  - (ii) the specifications and fill of packages of food, drugs, cosmetics and devices; and
  - (iii) the sale of any food, drug, cosmetic or device;
- (d) with respect to the use of any substance as an ingredient in any food, drug, cosmetic or device, for the purpose of preventing consumers or purchasers thereof from being misled or deceived as to the quantity, character, value, composition, merit or safety of that substance when so used or of preventing injury to the health of consumers or users thereof;
- (e) prescribing standards of composition, potency, purity or quality, or of any other property, for any article of food, drug, cosmetic or device;
- (f) with respect to the importation of any food, drug, cosmetic or device for the purpose of ensuring compliance with the provisions of this Act and the regulations;
- (g) with respect to the method of preparing, manufacturing, preserving, packaging, storing or testing of any food, drug, cosmetic or device, in the interests of, or for the prevention of injury to the health of, consumers or users thereof;
- (h) requiring persons who sell food, drugs, cosmetics or devices to maintain and keep such books or records as may be prescribed;
- (i) with respect to the form of certificate to be issued by the Minister for the purposes of section 7 (1) of this Act, and the manner of application therefor including the fees payable therefor, and with respect to the premises or processes or conditions of manufacture, including the qualifications of technical staff, which purposes of that subsection;
- (j) for requiring manufacturers of any drugs specified in the Forth Schedule to this Act to submit test portions of any batch of any such drug and with respect to the form of certificate to be issued by the Minister for the purposes of section 7 (2) of this Act including the fees payable therefor;
- (k) with respect to the manner of taking samples and the reporting of results of analysis or examination of samples;
- (l) exempting any food, drug, cosmetic or device from all or any of the provisions of this Act or of the regulations whether unconditional or subject to prescribed conditions;
- (m) prescribed forms for the purpose of this Act and the regulations;
- (n) providing for the analysis of food, drugs, or cosmetics otherwise than for the purpose of this Act and prescribing a tariff of fees to be paid for such analysis;
- (o) with respect to the distribution of samples of any drug;
- (p) with respect to any advertisement of drugs and cosmetics;
- (q) amending the First to Fifth Schedule to this Act in the interests of, or for the prevention of injury to the health of consumers or users; and
- (r) prescribing anything authorised or required by this Act to be prescribed.

*Penalties and legal proceedings*

**17. Penalties**

(1) Any person who contravenes any of the provisions of this Act or the regulations made under it or fails to comply with any requirement imposed on him by a notice under subsection (1) of section 4 of this Act shall be guilty of an offence and liable on conviction to a fine of not less than fifty thousand naira or imprisonment for a term not exceeding two years or to both such fine and imprisonment.

[1999 No. 21.]

(2) Where an offence under this section committed by a body corporate is proved to have been committed with the consent or connivance of, or to be attributable to any neglect on the part of any director, manager, secretary or other similar officer of the body corporate, or any person purporting to act in any such capacity, he, as well as the body corporate, shall be deemed to be guilty of the offence and shall be liable to be proceeded against and punished accordingly.

### **18. Trial of offences**

The Federal High Court shall have exclusive jurisdiction to try offences under this Act.

[1999 No. 62.]

### **19. Defence in proceedings for sale of food, etc.**

(1) Subject to subsection (2) of this section, it shall be a defence in any proceedings for an offence consisting of the sale of any article in contravention of this Act or the regulations to prove-

- (a) that the accused sold the article in the same package and in the same condition as it was in when he bought it; and
- (b) that the accused could not with reasonable diligence have ascertained that the sale of the article would be in contravention of this Act or the regulations.

(2) A person charged with an offence under this Act shall not be entitled to avail himself of the provisions of subsection (1) of this section unless he has given notice of this intention to do so at least ten days before the date of the trial and has at the same time disclosed to the prosecution the name of the person from whom he bought the article in question and the date of the purchase thereof.

### **20. Certificates and presumptions**

(1) In any proceedings under this Act or the regulations the production of a certificate purporting to be signed by an analyst shall be *prima facie* evidence of the matters stated therein; but the party against whom the certificate is produced may require the attendance of the analyst for the purpose of cross-examination.

(2) A certificate such as is mentioned in subsection (1) of this section shall not be received in evidence unless the party producing it has not less than three days before the trial supplied a copy thereof to the party against whom it is intended to be produced and has notified him in writing of the intention to produce it.

(3) Where any employee or agent commits an offence under this Act, his employer or principal shall be deemed to have committed the offence and be liable to be proceeded against and punished accordingly, whether or not the employee or agent has been prosecuted for the same offence; and for the purpose of this subsection, any person selling or ostensibly employed to sell on behalf of another person shall be presumed to be employed by him.

(4) Where, in any proceedings against any person for manufacturing any adulterated food or drug contrary to the provision of this Act or the regulations, it is established that the food or drug is one which, under the regulations, becomes adulterated if any prescribed substance is added thereto, and that that substance was found in his possession or on his premises, it shall be presumed, unless the contrary is proved, that the food or drug manufactured by him is adulterated.

### *General*

## **21. Interpretation**

In this Act, unless the context otherwise requires-

**"analyst"** means any person designated as a food and drug analyst or as a drug analyst under section 9 of this Act;

**"cosmetic"** means any substance or mixture of substances manufactured, sold or advertised for use in cleansing, improving or altering the complexion, skin, hair or teeth, and includes deodorants;

**"device"** means any instrument, apparatus or contrivance (including components, parts and accessories thereof) manufactured, sold or advertised for use in the diagnosis, treatment, mitigation or prevention of any disease, disorder, abnormal physical state or the symptoms thereof, in man or in animals;

**"drug"** includes any substance or mixture of substances manufactured, sold or advertised for use in-

- (a) the diagnosis, treatment, mitigation or prevention of any disease, disorder, abnormal physical state, or the symptoms thereof, in man or in animals;
- (b) restoring, correcting or modifying organic functions in man or in animals;
- (c) disinfection, or the control of vermin, insects or pests; or
- (d) contraception;

**"food"** includes any article manufactured, processed, packaged, sold or advertised for use as food or drink for human consumption, chewing gum and any ingredient which may be mixed with food for any purpose whatsoever and excludes-

- (a) live animals, birds or fish;

(b) articles or substances used as drugs;  
[1999 No. 21.]

**"insanitary conditions"** means such conditions or circumstances as might contaminate any food, drug or cosmetic with dirt or filth or render it injurious to health;

**"inspecting officer"** means any person designated as a food and drug inspecting officer under section 9 of this Act by the National Agency for Food and Drug Administration and Control;  
[1999 No. 21.]

**"label"** in relation to any food, drug, cosmetic, device or package includes any legend, word or mark attached to, included in, belonging to or accompanying that food, drug, cosmetic, device or package;

**"Minister"** means the Minister of Health;

**"package"** includes anything in which any food, water, drug, cosmetic or device is wholly or partly contained, wrapped, placed or packed;  
[1999 No. 21.]

**"prescribed"** means prescribed by the regulations;

**"regulations"** means any regulation made under this Act;

**"selling"** includes offering for sale, exposing for sale and having in possession for sale or distribution.

## 21. Short title

This Act may be cited as the Food and Drugs Act.

## SCHEDULES

### FIRST SCHEDULE

*Diseases, etc., referred to in section 2*

Acquired Immune Deficiency Syndrome  
[1999 No. 21.]

Alcoholism  
Appendicitis  
Arteriosclerosis  
Asthma  
Blood disorders  
Cancer  
Cataract

Diabetes  
Cholera  
Diphtheria  
Disorders of menstrual flow  
Disorders of prostate gland  
Dysentery  
Encephalitis  
Enteric fever  
Epilepsy  
Erysipelas  
Filariasis  
Gallstones, kidney stones, and bladder stones  
Gangrene  
Any genital or urinary diseases not mentioned elsewhere in this Schedule  
Glaucoma  
Goitre  
Hay fever  
Heart disease  
Hernia  
High blood pressure  
Infective hepatitis  
Influenza  
Jaundice  
Kidney disease  
Leprosy  
Locomotor ataxis  
Loss of youth  
Measles  
Meningitis  
Mental conditions  
Mumps  
Nervousness  
Nutritional disorders  
Obesity  
Onchocerciasis  
Paralysis  
Plague  
Pleurisy

Pneumonia  
 Poliomyelitis  
 Rabies  
 Rheumatic fever  
 Schistosomiasis  
 Sexual impotence, loss of virility or sterility  
 Sleeping sickness  
 Smallpox  
 Snake bite  
 Syphilis  
 Tetanus  
 Trachoma  
 Tuberculosis  
 Tumours  
 Typhoid fever  
 Undulant fever  
 Ulcers of the gastro-intestinal tract  
 Venereal diseases  
 Yaws  
 Yellow fever.

## SECOND SCHEDULE

*Drugs referred to in section 3*

|                | PART A  |
|----------------|---|
| DET            | (N, N-diethyltryptamine)  |
| DMHP           | (3-(1, 2-dimethylhepty)-1-hydroxy-7, 8, 9, 10 tetrahydro-6, 6, 9-trimethyl 6H-dibenzo (b, 9) pyran) |
| DMT            | (N,N-dimethyltryptamine)  |
| LYSERGIDE/LSD, | (+)-N,N-diethyllysergamide (d-lysergic acid diethylamide)   |
| LSD-25         | (3,4, 5-trimethoxyphenethylamine)   |
| Mescaline      | (3-hexyl-1-hydroxy-7, 8, 9, 10-tetrahydro-6, 6, 9-trimethyl-6H-dibenzo (b,d) pyran)                 |
| parahexyl      | (3-(2-dimethylaminethyl)-4-hydroxyindole)   |
| psilocine,     | (3-(2-dimethylaminoethyl) indol-4-yl dihydrogen phosphate)  |
| psilotsin      | (2-amino-1-(2, 5-dimethoxy-4-methyl) phenyl-propane)  |
| Psilocybine    |   |
| STP,DO         |   |
| M              |   |

PART A-continued

tetrahydrocannabinols, all isomers 1-hydroxy-3-pentyl-6a, 7, 10, 10a,-tetrahydro-6, 6, 9 tri-  
methyl-6-H-dibenzo (b,d) pyran

|                 |  |
|-----------------|--|
| AMPHETAMINE     | (+)-2-amino-1-phenylpropane)                         |
| DEXAMPHETAMINE  | (+)-2-amino-1-phenylpropane)                         |
| METHAMPHETAMINE | (+)-2-methylamino -1-phenyl propane                  |
| METHYLPHENIDATE | (2-phenyl-2-(2-piperidyl) acetic acid, methyl ester) |
| PHENMETRAZINE   | (3-methyl-2-phenylmorpholine)                        |
| PHENCYCLIDINE   | (1-(1-phenylcyclohexyl) piperidine)                  |

Any synthetic compound or salt (however structurally derived) of Amphetamine or of any substance of the like nature as Amphetamine.

PART B

|               |  |
|---------------|--|
| AMOBARBITAL   | (5-ethyl-5-(3-methylbutyl)barbituric acid) |
| CYCLOBARBITAL | (5-C)                                      |
| GLUTETHIMIDE  |  |
| PENTOBARBITAL |  |
| SECOBARBITAL  |  |
| AMFEPRAMONE   |  |
| BARBITAL      |  |

ethchlorvynol

ETHINAMATE  
MEPROBAMATE  
METHAQUALONE  
METHYLPHENOBARBITAL  
AL  
METHYPRYLON  
PHENOBARBITAL  
PIPRADROL  
SPA  
COBALT PREPARATIONS  
CYCLIZINE  
SULPHAMETHOXYDIAZINE  
SULPHADIMETHOXINE  
AMINOPYRINE AND  
DIPYRONE  
ACECARBROMAL



PART B-continued

CARBROMA  
L  
BROMOSOV  
AL

METALLIC TIN AND ITS COM-  
POUNDS  
HEXACHLOROPHANE (2-(P-Iaminophenyl)-ethylglytarimide)

ELIPTEN  
MECLOZINE 2. 2-thiobis (4, 6-dichlorophenol)

BITHIONOL  
IMIPRAMINE  
EFOCAINE  
NEONOVUM  
XENAZOIC ACID OR  
XENALAMINE

DITHLAZANINE  
IODIDE  
CHLOROPHENTERMI  
NE

CYCLAMATES, CYCLOHEXYLEMINE AND THEIR

DERIVATIVES  
COMPHETAMINES  
CHLORAMPHENICOL AND ITS  
PREPARATIONS  
CAFFEINE  
EPHEDRIN AND ITS SALT

THIRD SCHEDULE

*Publications referred to in section 5*

Pharmacopoeia Internationalis

Any other pharmacopoeia published under the authority of a recognised medical or pharmaceutical council of any country.

FOURTH SCHEDULE

*Drugs referred to in section 7 (1)*

Liver extract in all forms  
Insulin in all forms  
Anterior pituitary extracts

Radioactive isotopes  
Living vaccines for oral or parenteral use  
Drugs prepared from micro-organisms or viruses, for parenteral use  
Sera and drugs analogous thereto, for parenteral use  
Antibiotics for parenteral use.

#### FIFTH SCHEDULE

*Drugs referred to in section 7 (2)*

Arsphenamine  
Dipchlorophenarsine hydrochloride  
Neoarsphenamine  
Oxophenarshine hydrochloride  
Sensitivity discs and tablets  
Sulpharsphenamine.

#### FOOD AND DRUGS ACT

#### **SUBSIDIARY LEGISLATION**

*No Subsidiary Legislation*