UNITED NATIONS

E/NL. 1986/41
24 October 1988

ENGLISH ONLY *

LAWS AND REGULATIONS

PROMULGATED TO GIVE EFFECT TO THE PROVISIONS OF THE INTERNATIONAL TREATIES ON NARCOTIC DRUGS AND PSYCHOTROPIC SUBSTANCES

In accordance with the relevant articles of the international treaties on narcotic drugs and psychotropic substances, the Secretary-General has the honour to communicate the following legislative texts.

MALTA

Communicated by the Government of Malta

NOTE BY THE SECRETARIAT

(a) Some editing of texts may be done by the Secretariat in the interest of clarity. In this connection, words in square brackets [ ] have been added or changed by the Secretariat.

(b) Only passages directly relevant to the control of narcotic drugs or psychotropic substances have been reproduced in this document. Non-relevant parts of laws and regulations have been deleted by the Secretariat; such deletions are indicated by [...].

MEDICAL AND KINDRED PROFESSIONS ORDINANCE

(CAP. 51)

Drugs (Control) Regulations, 1985

* Note by the Secretariat: The present document is a direct reproduction of the text received by the Secretariat.

V.88-30295

MEDICAL AND KINDRED PROFESSIONS ORDINANCE

(CAP. 51)

Drugs (Control) Regulations, 1985

IN exercise of the powers conferred by section 35B of the Medical and Kindred Professions Ordinance, the Minister of Health, on the advice of the Council of Health, has made the following regulations:—

Citation.

1. These regulations may be cited as the Drugs (Control) Regulations, 1985.

Interpretation.

2. In these regulations, unless the context otherwise requires, —

Act LI of 1975

"identity card" means a valid identity card issued in accordance with the provisions of the Identity Card Act, 1975 and of any regulations made thereunder;

"register" means a bound book with consecutively numbered pages, but does not include any form of loose leaf register or card index;

"restricted drug" means any one of such drugs or chemical substances as are listed in the First Schedule to these regulations;

"specified drug" means any one of such drugs or chemical substances as are specified in the Second Schedule to these regulations;

"Superintendent" means the Superintendent of Public Health.

Importation, etc., of restricted drugs to be specially authorised.

3. (1) No person may import, manufacture, export, purchase, sell, use or be in possession of any restricted drug without a special authorisation in writing by the Superintendent.

(2) The Superintendent shall not grant a special authorisation to use any restricted drug except in special cases for scientific or very limited medical purposes inside medical or scientific establishments under Government control or specifically approved by the Superintendent, who shall have the power to impose in any authorisation any conditions and requirements he may deem fit to impose with a view to safeguard against abuse.

(3) The Superintendent may at any time, without giving any reason whatsoever, withdraw any authorisation granted by him under paragraph (1) of this regulation.

(4) Any person who has been authorised in accordance with paragraph (1) of this regulation shall keep such registers as shall be necessary in accordance with the Third Schedule to these regulations in relation to the manufacture, acquisition and/or disposal of restricted drugs, in which registers there shall be entered the details relative to such drugs. Each entry shall be signed, by the person authorised under paragraph (1), within twenty-four hours of any transaction or process, and any such register shall be preserved and kept available for inspection by the Superintendent or his representative for at least two years after the date of the last entry recorded therein.

(5) Any entry in the register shall be made in ink or other indelible material and shall be entered on the same day in which the transaction, administration or process is effected, or, when this is not possible, on the next following day.

(6) Any register used or in use with regard to one particular premises shall not be used also with regard to any other premises or for any purpose other than of this regulation.

(7) No entry in the register referred to in the last preceding paragraphs of this regulation shall be cancelled, obliterated or altered or shall be entered with some untrue particulars:

Provided that if any mistake is committed in any entry, such mistake shall be corrected by means of a note in the margin or at the foot of the page, which note shall contain the correction required and the date of the note.

(8) No person may dispose of any restricted drug except to persons specially authorised to possess or use such drugs.

Transactions regarding specified drugs to be registered.

4. (1) Any person who is licensed or authorised to manufacture, import, sell, supply, distribute, or administer any specified drug, whether in the line of his trade or profession or as a medicament on the presentation of a prescription, shall keep such registers as shall be necessary to show any transaction regarding the manufacture, importation, purchase, sale, supply, distribution or administration of any such drug.

(2) (i) No person shall import, stock or sell any specified drug by wholesale unless he holds a specific licence for the purpose from the Superintendent.

(ii) No licence for the purposes of the foregoing subparagraph shall be issued unless the drug or drugs in respect of which it is issued are under the direct responsibility of an apothecary.

(3) Any specified drug manufactured, imported, exported, purchased, sold, supplied, distributed or administered shall be entered in the register in the form shown with such variations as circumstances may require, and containing such particulars as are shown in the Third Schedule to these regulations and as are applicable to the case.

(4) Any entry in the register shall be made in ink or other indelible material and shall be entered on the same day in which the transaction, administration or process is effected, or, when this is not possible, on the next following day.

(5) Any register used or in use with regard to one particular premises shall not be used also with regard to any other premises or for any purpose other than of this regulation.

(6) No entry in the register referred to in the preceding paragraphs of this regulation shall be cancelled, obliterated or altered or shall be entered with some untrue particulars:

Provided that if any mistake is committed in any entry, such mistake shall be corrected by means of a note in the margin or at the foot of the page, which note shall contain the correction required and the date of the note.

(7) No person may export any specified drug without the prior authorisation from the Superintendent.

(8) In the case of importation or exportation of specified drugs, the Superintendent shall have the power to direct the procedure to be followed and the details to be submitted in the relative application, and may also, before issuing an export authorisation, require that the authorisation of the importing country be obtained.

(9) The Superintendent may prohibit, or otherwise restrict, the importation of any specified drug if he so considers it necessary in the public interest.

Possession of specified drugs.

5. (1) No person, unless duly authorised, may have in his possession any specified drug.

(2) For the purpose of this regulation, a person shall be deemed to be duly authorised if his name is entered in the Medical Register, in the Register of Dental Surgeons, or in the Veterinary Surgeons' Register, or if he is a managing apothecary, or is in possession of a licence issued by the Superintendent under section 74 of the Medical and Kindred Professions Ordinance, or has obtained such drug in virtue of a medical prescription, or otherwise in accordance with the provisions of these regulations:

Provided that, in respect of a person authorised in virtue of a medical prescription, such prescription shall not be deemed to be valid if, at the time of the receipt of the prescription, such person was under the treatment of another medical practitioner and had been receiving such specified drugs in virtue of a prescription from this other medical practitioner and had not informed the prescriber of this fact.

Safe keeping of restricted and specified drugs.

6. Any person authorised to be in possession of specified drugs in accordance with the provisions of regulation 5 of these regulations and any person specially authorised in respect of restricted drugs, as the case may be, shall, when such drugs are not in use, keep same under lock and key and it shall be the duty of any such person to take all steps necessary to ensure security and to prevent theft or other diversification of stock:

Provided that nothing in this regulation shall apply to any person who is in possession of specified drugs in virtue of a medical prescription.

Use and contents of prescription.

7. (1) Every prescription for a specified drug shall be written in ink or in other indelible material on the form set out in the Seventh Schedule to these regulations.

(2) It shall be the duty of a medical practitioner issuing a prescription for a specified drug to fill in a clear and legible hand Part A of the form set out in the Seventh Schedule, and to supply all the details and give all the information in the appropriate space as therein required; the medical practitioner shall further add his signature in full and the date when the prescription was issued:

Provided that in the case of a medical practitioner authorised under section 4 of the Medical and Kindred Professions Ordinance to practise the medical profession but who is not yet registered with the Medical Council, such practitioner shall insert the number given to him by the Superintendent instead of the Medical Council registration number on the said prescription.

(3) Subject to the provisions of paragraph (11) of this regulation, no medical practitioner shall issue a prescription for a specified

drug unless the prescription complies with the provisions of this regulation and such drug is required for the purpose of medical treatment:

Provided that a medical practitioner may, subject to the other provisions of these regulations, issue a prescription for professional use for an amount not in excess of ten phials for injection or of twenty tablets or capsules.

(4) Every medical practitioner who obtains a specified drug for professional use as provided in the proviso to paragraph (3) of this regulation shall keep, in accordance with, and without prejudice to, the provisions of regulation 4 of these regulations, a record in an appropriate register of the name and surname (if applicable), the age and address of the patient to whom the drug has been administered and the date of administration.

(5) No medical practitioner shall issue a prescription for a specified drug to any person unless such person is well known to him or unless the medical practitioner has ascertained the identity of such person through his identity card.

(6) No medical practitioner shall issue a prescription for a specified drug to any person unless the said medical practitioner has taken reasonably sufficient steps to ascertain that such person is not, at the time of issuing the prescription, receiving treatment from another medical practitioner in respect of addiction to any specified drug or otherwise, and that such person has not been supplied with any such drug on a prescription issued by that other medical practitioner.

(7) A medical practitioner shall use a separate form in respect of every drug prescribed by him under this regulation and no drug other than a specified drug or a drug falling under the Dangerous Drugs Ordinance, 1939, may be prescribed on those forms.

(8) A medical practitioner shall in prescribing a specified drug, use only prescription forms from the booklet of forms in serial number, issued to him by the Superintendent upon a request made on the form set out in the Eighth Schedule to these regulations and it shall be the duty of every medical practitioner, whether for the purpose of his private practice or for carrying out his duties in an official capacity as a result of his employment with Government, with a view to meeting the needs of his patients, to make any request for such prescription booklets on the said form; such request form shall be correctly filled in all respects and signed by the practitioner, and shall be either handed in by such practitioner personally to the Chief Pharmacist at the Government Medical Stores at Gwardamangia or sent by post to the Superintendent of Public Health at the Department of Health in Valletta; when sent by post the envelope may be marked "Public Health Notifications" for the purpose of exemption from postage.

(9) It shall be duty of a medical practitioner to report in writing forthwith to the Superintendent any case of theft or loss of such booklet and it shall not be lawful for a medical practitioner to use any prescription form from any booklet of forms issued to another medical practitioner.

(10) For the purpose of this regulation, unless the content otherwise requires, the expression "medical practitioner" includes a dental surgeon, a dentist and a veterinary surgeon.

(11) The provisions of paragraphs (1), (2), (7) and, in so far as applicable, paragraphs (8) and (9) of this regulation shall not apply

to the prescription of specified drugs for administration to ward patients in Government hospitals, which prescription shall be controlled by the hospital internal rules.

Dispening of of prescriptions.

8. (1) No person other than an apothecary shall dispense a prescription for a specified drug.

(2) No apothecary shall dispense a prescription for a specified drug unless —

(a) he is acquainted with the signature of the person by whom it purports to have been issued, he has no reason to suppose that it is not genuine and he has taken reasonable steps to satisfy himself that it is genuine; and

(b) the prescription complies with the provisions of regulation 7 of these regulations.

(3) No specified drug shall be supplied more than once on the same prescription.

(4) An apothecary dispensing a prescription for a specified drug shall fill in a clear and legible hand, in ink or other indelible material, Part C of the form set out in the Seventh Schedule to these regulations, and supply all the details and give all the information as therein required; the apothecary shall further add his signature in full and the date, and shall, after dispensing the prescription, retain it.

(5) For the purpose of paragraph (4) of this regulation it shall be the duty of an apothecary dispensing a prescription for a specified drug to request the identity card of the person in respect of whom the prescription has been issued and it shall be the duty of the person who intends to acquire a specified drug, whether for himself or on behalf of the person to whom the drugs have been prescribed, to present the said identity card as well as his own identity card together with the prescription:

Provided that the provisions of this paragraph and the relative part of the provisions of paragraph (4) of this regulation shall not apply in respect of a person who has not yet been issued with an identity card.

(6) It shall be the duty of every managing apothecary of a licensed dispensary to send to the Superintendent in a sealed envelope on the first day of every month all the prescriptions for specified drugs dispensed from that dispensary during the preceding month; that envelope shall be addressed "Superintendent of Public Health, Department of Health, Valletta" and may be marked "Public Health Notifications" for the purpose of exemption from postage.

Specimen signature of prescription.

9. (1) Saving the other provisions of these regulations, no medical practitioner, dental surgeon, dentist or veterinary surgeon shall prescribe any specified or restricted drug unless the Superintendent is in possession of his specimen signature.

(2) The Superintendent may, for the purpose of the control of drugs, circulate to all managing apothecaries a copy of any such specimen signature in his possession.

Delivery of specified drugs to licensed or authorised persons.

10. (1) Saving the provisions of the following paragraphs of this regulation, no person shall deliver any specified drug to any other person except against the presentation of a duly signed receipt therefor.

(2) No person shall deliver any specified drug to any other person who is not a managing apothecary or who is not otherwise law-

fully authorised to be in possession of such drug, unless such other person produces a written authorisation in that behalf duly signed by the managing apothecary or by the person otherwise lawfully authorised to be in possession of such drug, and unless the person delivering the said drug is satisfied that the authorisation is genuine.

(3) Nothing in the preceding paragraphs of this regulation shall be deemed to refer to the supply of any specified drug against the presentation of a medical prescription in accordance with the provisions of these regulations or to the administration of any such drug by or under the supervision of a medical practitioner, dental surgeon, dentist or veterinary surgeon.

(4) For the purpose of this regulation, "deliver" includes any act or omission whereby a person allows or suffers any other person, not being a person authorised to be in possession of specified drugs under regulations, to take possession of any such drug.

Preservation of prescriptions, registers, records, and other documents.

11. Saving the provisions of paragraph (6) of regulation 8 of these regulations any dispensed prescription, register, invoice or other document relating to the manufacture, importation, purchase, sale, supply, distribution, or other disposal of specified drugs shall be kept at the premises to which they refer and shall be preserved for a period of not less than two years from the date of the prescription, record, invoice or other document, or from the date of the last entry in the register, as the case may be, and during such time they shall be open to inspection by the Superintendent or his representative.

Notification by prescriber and managing apothecary in the case of certain drugs.

12. (1) No medical practitioner may issue a prescription for any preparation consisting of or containing amphetamine, or any isomer of amphetamine, flunitrazepam, mecloqualone, methaqualone, methylphenidate, or phenmetrazine, or any salts or esters of any such drugs, or for any preparation consisting of secobarbital or of a combination of secobarbital and amobarbital, in either case in tablet or capsule form, or in sachet or cachet, or of any salts or esters thereof, unless he has the prior authorisation in writing by the Superintendent.

(2) The Superintendent may, before granting any authorisation as is referred to in paragraph (1) of this regulation, request from the medical practitioner such information as the Superintendent may require, including whether according to the opinion of the medical practitioner the patient is suffering from drug dependency.

(3) When a request for such authorisation is made, the Superintendent shall submit such request to a panel of medical specialists who shall advise the Superintendent as to whether such request is to be authorised.

(4) Before making their recommendations it shall be the duty of the panel to see whether such prescribing is indicated and the panel shall also, where indicated, make such recommendations as may be necessary with a view to weaning of patient from such drug.

(5) Notwithstanding the provisions of paragraphs (3) and (4) of this regulation, the Superintendent may, if he considers it is in the interest of the patient, authorise the prescription of such drug until the report of the panel is received.

(6) It shall be the duty of any medical practitioner who delivers a prescription as is referred to in paragraph (1) of this regulation to inform accordingly the Superintendent within twenty-four hours

of his prescribing such a preparation, giving the details as set out in the Fourth Schedule to these regulations.

(7) It shall be the duty of any apothecary dispensing a prescription as is referred to in paragraph (1) of this regulation to inform the Superintendent accordingly within twenty-four hours of his dispensing such a preparation, giving the details as set out in the Fifth Schedule to these regulations.

(8) All notifications made in compliance with paragraphs (6) and (7) of this regulation shall be marked "Confidential".

(9) No veterinary surgeon, dental surgeon or dentist may prescribe any preparation as is referred to in paragraph (1) of this regulation.

(10) If the President of the Republic has reason to suspect that a medical practitioner, or a dental surgeon or dentist, is supplying or prescribing any specified drug to or for either himself or any other person other than, or more than, is properly required for medical or dental treatment of himself or that other person, as the case may be, or that a veterinary surgeon is supplying or prescribing any specified drug in respect of an animal under his care other than, or more than, is properly required for the treatment of such animal, as the case may be, the said President may refer the matter to the Medical Council, and, if the said Council so recommends, the said President may prohibit the said medical practitioner, dental surgeon or dentist, or veterinary surgeon, from supplying, prescribing, procuring or possessing such drug, for such period as the President may deem fit, and it shall then be unlawful for that medical practitioner, dental surgeon or dentist, or veterinary surgeon to supply, prescribe, procure or possess any such drug during such period:

Provided that the said President may, at any time, modify or withdraw any such prohibition.

(11) Notice of any prohibition, modification or withdrawal of any prohibition as is referred to in paragraph (10) of this regulation shall be given in writing to the person affected thereby.

(12) It shall be lawful for the Superintendent to circulate to —

(a) all medical practitioners the name, and other details as are shown in the Sixth Schedule to these regulations, of any person in respect of whom any preparation as is referred to in paragraph (1) of this regulation has been prescribed, and

(b) all medical practitioners, dental surgeons and dentists the name, and other details as are shown in the Sixth Schedule to these regulations, of any person in respect of whom any specified drug has been prescribed or dispensed.

Warning and cautions on label, etc.

13. (1) The Superintendent may require all manufacturers, importers, exporters, and all other persons who trade or otherwise deal in any restricted drug or any specified drug, to include such cautions or other warnings on the label where practicable, or in the accompanying leaflet of retail packages of such drugs, as are in his opinion necessary for the safety of the persons using such drugs.

(2) It shall not be lawful for any person to advertise in any way to the general public any restricted drug or any specified drug.

14. (1) These regulations shall not apply, in respect of any specified drug, to any drug when it forms part of, or is incorporated in, a preparation ready for use and which requires no further compounding, unless such drug is a main or the only active ingredient of the preparation. Exemptions.

(2) These regulations shall likewise not apply in the case of a preparation containing a specified drug, compounded in such a way that it presents no risk, or only a negligible risk, of abuse, and the substance cannot be recovered by readily applicable means in a quantity liable to abuse, so that the preparation does not give rise to a public health or social problem.

15. (1) The Drugs (Control) Regulations, 1976,[1/] other than regulation 13 thereof, are hereby revoked without prejudice to any liability incurred and to any proceedings taken or that may be taken thereunder. Revocation and saving. L.N. 31 of 1976

(2) Any dispensed prescription, register, record, invoice or other document which, immediately before the coming into force of these regulations, are preserved under the regulations hereby revoked, shall continue to be preserved and to be open for inspection and shall for these purposes be deemed to have always been preserved under these regulations.

FIRST SCHEDULE

(Regulation 2)

International or other nonproprietary name or other trivial name	*Chemical name*
DET	*N, N* — diethyltryptamine
DMHP	3-(1, 2-dimethylheptyl)-1-hydroxy-7, 8, 9, 10-tetrahydro-6, 6, 9-trimethyl-6H-dibenzo [b, d] pyran
DMT	*N, N*-dimethyltryptamine
DOB, dimethoxy bromoamphetamin	2, 5-dimethoxy-4-bromoamphetamine
(+) LYSERGIDE, LSD, LSD-25	(+) *N, N*-diethllysergamide (d-lysergic acid diethylamide)
MDA, methylene dioxyamphetamine	3, 4, methylinedioxyamphetamine
MESCALINE, MESCAL BUTTON, PEYOTE, PEYOTL	3, 4, 5-trimethoxyphenethylamine
PARAHEXYL	3-hexyl-l-hydroxy-7, 8, 9, 10-tetrahydro-6, 6, 9-trimethyl-6H-dibenzo [b, d] pyran
PCE	N-ethyl-l-phenylcyclohexylamine
PHP, PCPY	l-(1-phenyclyclohexyl) pyrrolidine
PSILOCINE, PSILOTSIN	3-(2-dimethylaminoethyl)-4-hydroxyindole
PSILOCYBINE	3-(2-dimethylaminoethyl) indol-4-yl dihydrogen phosphate
STP, DOM	2-amino-1-(2, 5-dimethoxy-4-methyl) phenylpropane
TETRAHYDROCANNABINOLS, ALL ISOMERS	1-hydroxy-3-pentyl-6a, 7, 10, 10a-tetrahydro-6, 6, 9-trimethyl-6-H-dibenzo [b, d] pyran
TCP	1-[1-(2-thienyl) cyclohexyl] piperidine

And any derivatives, salts, or esters of the above.

1/ Note by the Secretariat: E/NL.1977/54

SECOND SCHEDULE

(Regulation 2)

International or other nonproprietary name or other trivial name	*Chemical name*
AMFEPRAMONE, DIETHYLPROPION	2-(diethylamino) propiophenone
AMOBARBITAL	5-ethyl-5-(3-methylbutyl) barbituric acid
AMPHETAMINE	(±)-2-amino-1-phenylpropane
BARBITAL	5, 5-diethylbarbituric acid
BENZPHETAMINE	
CHLORAL HYDRATE	2, 2, 2-trichloroethane-1, 1-diol
CHLORDIAZEPOXIDE	7-chloro-2-methylamino-5-phenyl-3H-1, 4-benzo-diazepine 4-oxide
CHLORPHENTERMINE	p-chloro-α-dimethylphenethlylamine
CYCLOBARBITAL	5-(1-cyclohexen-1-yl)-5-ethylbarbituric acid
DEXAMPHETAMINE	(+)-2-amino-1-phenylpropane
DIAZEPAM and other compounds containing the chemical structure of DIHYDRO-1:4 BENZODIAZEPINE or of DIHYDRO-1:5 BENZODIAZEPINE substituted to any degree	7-chloro-2, 3-dihydro-1-methyl-5-phenyl-1H-1, 4-benzodiazepine-2-one
ETHCHLORVYNOL	ethyl-2-chlorovinylethinyl-carbinol
ETHINAMATE	1-ethynylcyclohexanol carbamate
FENCAMFAMIN	2-Ethylamino-3-phenylnorbornane
FENFLURAMINE	N-ethyl-α-methyl-m(trifluoromethyl) phenethylamine
GLUTETHIMIDE	2-ethyl-2-phenylglutarimide
MAZINDOL	5-p-chlorophenyl-2, 3-dihydro-5H-imidazo [2, 1-a] isoindol-5-ol
MECLOQUALONE	3-(O-chlorophenyl)-2-methyl-4 (3H)-quinazolinone
MEPHENTERMINE	N-α-trimethylphenethylamine
MEPROBAMATE	2-methyl-2-propyl-1, 3-propanediol dicarbamate
METHAMPHETAMINE	(+)-2-methylamino-1-phenylpropane
METHAQUALONE	2-methyl-3-o-tolyl-4(3H)-quinazolinone
METHYLPHENIDATE	2-phenyl-2-(2-piperidyl) acetic acid, methyl ester
METHYLPHENOBARBITAL	5-ethyl-1-methyl-5-phenyl-barbituric acid
METHYPRYLON	3, 3-diethyl-5-methyl-2, 4-piperidinedione
PARALDEHYDE	acetaldehyde trimer
PENTAZOCINE	1, 2, 3, 4, 5, 6, Hexahydro-6, 11 dimethyl-3 -(3 methylbutyl-2 enyl)-2, 6 Methano-3-benzazocin-8 ol
PENTOBARBITAL	5-ethyl-5-(1-methylbutyl) barbituric acid
PHENTERMINE	dimethylphenethylamine
PIPRADROL	1, 1-diphenyl-1-(2-piperidyl) methanol
PHENCYCLIDINE	1-(1-phenylcyclohexyl) piperidine
PHENDIMETRAZINE	(+)-3, 4, -Dimethyl-2-phenylmorpholine
PHENMETRAZINE	3-methyl-2-phenylmorpholine
PHENOBARBITAL	5-ethyl-5-phenylbarbituric acid
SECOBARBITAL	5-allyl-5-(1-methylbutyl) barbituric acid
SPA	(-)-1-dimethylamine-1, 2, diphenylethane

And any salts or esters of the above.

THIRD SCHEDULE

(Regulations 3 and 4)

FORM OF REGISTERS

I (a) Restricted Drugs, imported, purchased, or otherwise obtained

Name of substance imported, purchased or otherwise obtained	Date on which supply is received	Name of person or firm from whom obtained	Address of person or firm from whom obtained	Amount obtained	Form in which obtained	Purpose of use	Date of Authorisation from S.P.H.	Signature of Authorised Person

(b) Restricted Drugs sold, supplied, administered or otherwise used

Name of substance	Date	Name of person or firm to whom sold, supplied or administered or purpose of use whichever applicable	Address of person to whom sold, supplied or administered or premises where used whichever applicable	Amount sold or used	Form	Prescription No. or authorisation from S.P.H.	Quantity remaining in stock	Signature of Authorised Person

II (a) Specified Drugs imported, purchased or otherwise obtained

Name of substance imported, purchased or otherwise obtained	Date on which supply is received	Name of person or firm from whom obtained	Address of person or firm from whom obtained	Amount obtained	Form in which obtained

(b) Specified Drugs exported, sold, supplied or administered

Name of substance exported, sold, supplied or administered	Date on which exported, sold, supplied or administered	Name of person or firm to whom exported, sold, supplied or administered	Address of person or firm to whom exported, sold, supplied or administered	Amount exported, sold, supplied or administered	Form in which exported, sold, supplied or administered	When sale is on prescriptions prescription number, the name of the prescriber and the official serial number printed on the prescription form

III (a) Manufacture of Restricted Drugs

Name of substance manufactured	Date of manufacture	Amount manufactured	Form	Signature of Authorised Person	Date of authorisation

(b) Manufacture of Specified Drugs

Name of substance manufactured	Date of manufacture	Amount manufactured	Form	Signature of person responsible

FOURTH SCHEDULE

Regulation 12 (4)

CONFIDENTIAL

(Return in terms of regulation 12 — Prescriptions for amphetamine and its isomers, flunitrazepam, mecloqualone, methaqualone, methylphenidate and phenmetrazine and secobarbital or a combination of secobarbital and amobarbital in tablet or capsule form or in sachet or cachet and their respective salts or esters)

IMPORTANT — Prescription for these preparations can only be issued after written approval is obtained from the Superintendent of Public Health for each prescription.

NAME AND ADDRESS OF PATIENT .....

.....

IDENTITY CARD NUMBER .....

AGE .....

NAME OF PREPARTION AND STRENGTH .....

.....

QUANTITY PRESCRIBED .....

DAILY DOSE .....

DATE OF PRESCRIPTION .....

DATE AND REF. NO. OF APPROVAL BY SUPERINTENDENT OF PUBLIC HEALTH .....

SIGNATURE OF MEDICAL PRACTITIONER

NAME (Block Letters) .....

ADDRESS .....

.....

.....

FIFTH SCHEDULE

Regulation 12 (5)

CONFIDENTIAL

(Return in terms of regulation 12 — Prescriptions for amphetamine and its isomers, flunitrazepam, mecloqualone, methaqualone, methylphenidate and phenmetrazine and secobarbital or a combination of secobarbital and amobarbital in tablet or capsule form or in sachet or cachet and their respective salts or esters)

NAME AND ADDRESS OF PATIENT .....

.....

IDENTITY CARD NUMBER .....

NAME OF PREPARTION AND STRENGTH .....

.....

QUANTITY PRESCRIBED .....

DATE OF PRESCRIPTION .....

NAME OF PRESCRIBER .....

SIGNATURE OF APOTHECARY .....

NAME (Block Letters) .....

DH LICENCE NUMBER .....

ADDRESS (Of Pharmacy) .....

.....

DATE OF DISPENSING .....

SIXTH SCHEDULE

(Regulation 12)

Name and address of patient .....

Identity Card Number .....

Date of prescription .....

Name of drug .....

Amount prescribed .....

Name of medical practitioner, dental surgeon or dentist who has prescribed the drug

.....

SEVENTH SCHEDULE

(Regulation 7)

PRESCRIPTION FOR NARCOTIC AND PSYCHOTROPIC DRUGS*
(Only one item may be prescribed on this Form)

A. TO BE FILLED BY PRESCRIBER

PATIENT
Name and Surname .....
Address .....

ITEM (incl. Form & Dose)	Quantity

Directions for Use

Additional Information in respect of Prescriptions for METHADONE

Expected Duration of Treatment .....
Medical Indication .....

PRESCRIBER Med. Council Reg. No. ☐☐☐☐☐☐

Name and Surname .....
Address .....

Signature ..... Date .....

B. FOR OFFICIAL USE ONLY

To be filled by Government Dispenser

PRESCRIPTION TYPE
Pink Form ☐ Schedule III ☐ Others ☐
Direct ☐ Postal ☐

LOCATION DISPENSED ☐☐☐☐

To be filled by Coder

Item No. ☐☐☐☐☐☐ Prescription Origin ☐☐

C. TO BE FILLED BY DISPENSING APOTHECARY

PATIENT
Surname ☐☐☐ Name ☐☐☐
(First three letters)
I.D. ☐☐☐☐☐☐☐☐
Male ☐ Female ☐

ITEM
Quantity supplied
Total Quantity in Units .....
Total Quantity in Weight/Measure .....

PRESCRIPTION ORIGIN
Hospitals ..... NAME
Govt. Clinics ..... DISTRICT
Private ☐

DISPENSER
Signature .....
Pharmacy Board Reg. No. ☐☐☐
Dispensary Address
.....
Dispensary DH Licence No. ☐☐☐☐
Date ☐☐☐☐☐☐

Serial No.

EIGHTH SCHEDULE

(Regulation 7)

REQUEST FOR PRESCRIPTION FORMS FOR NARCOTIC AND PSYCHOTROPIC DRUGS

I request ..... booklet/s by ..... prescription forms.

Name and surname of prescriber .....
(IN BLOCK LETTERS)

Private address .....
(IN BLOCK LETTERS)

Signature ..... Med. Council Reg. No.

FOR OFFICIAL USE

Supplied ..... booklet/s by ..... prescription forms.

Serial No. ..... to .....

(a) If withdrawn personally by prescriber.

....
Signature of Prescriber Date

(b) If forwarded by mail.

Forwarded by ..... on .....

A.R. Card to be attached.