1973-06
Revoked
Subsidiary
1974/138

DRUGS (MISUSE) REGULATIONS

Regulations made under s.34.

DRUGS (MISUSE) REGULATIONS

Revoked by LN. 2005/106 as from 14.7.2005

(LN. 1974/138)

1.1.1975

Amending enactments	Relevant current provisions	Commencement date
1988/085	rr. 9(3), 10(1)(c) and (2), 12(2), 13(2), 16(f) and (g) and Sch.5	1.2.1988

ARRANGEMENT OF REGULATIONS.

23. *Omitted.*

Schedule 1.

Schedule 2.

Schedule 3.

Schedule 4.

Schedule 5.

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DRUGS (MISUSE) REGULATIONS

Title.

1. These Regulations may be cited as the Drugs (Misuse) Regulations.

Interpretation.

2. (1) In these Regulations, unless the context otherwise requires–

"health prescription" means a prescription issued by a doctor under the Group Practice Medical Scheme;

"master" has the same meaning as in the Merchant Shipping Ordinance;

"matron or acting matron" includes any male nurse occupying a similar position;

"the Merchant Shipping Ordinance" includes those parts of the Merchant Shipping Acts which apply to Gibraltar;

"prescription" means a prescription issued by a medical practitioner for the medical treatment of a single individual, by a dentist for the dental treatment of a single individual or by a veterinary practitioner for the purposes of animal treatment;

"register" means a bound book and does not include any form of loose leaf register or card index;

"registered dispensary" means a dispensary registered under the Medical and Health Ordinance, 1997;

"retail dealer" means a person lawfully conducting a registered dispensary;

"sister or acting sister" includes any male nurse occupying a similar position;

"wholesale dealer" means a person who carries on the business of selling drugs to persons who buy to sell again.

(2) Nothing in these Regulations shall be construed as derogating from any power or immunity of the Crown, its servants or agents.

Metric and imperial systems.

3. (1) For the purposes of these Regulations–

(a) a controlled drug shall not be regarded as supplied otherwise than on a prescription or other order by reason only that the prescription or order specifies a quantity of the controlled drug in terms of the imperial system and the quantity supplied is the equivalent of that amount in the metric system;

(b) where any person may lawfully be in possession of a quantity of a controlled drug determined by or under these Regulations in terms of the imperial system he shall be deemed not to be in possession of a quantity of that controlled drug in excess of the first-mentioned quantity by reason only that he is in possession of a quantity of that drug which is the equivalent of the first-mentioned quantity in the metric system.

(2) For the purposes of this regulation the quantity of a controlled drug in the metric system which is the equivalent of a particular quantity in the imperial system shall be taken to be the appropriate quantity ascertained in accordance with the provisions of the United Kingdom's Weights and Measures (Equivalents for Dealings with Drugs) Regulations, 1970.

S.I.1970-1897

Exceptions.

4.(1) Sections 5(1) and 7(1) of the Ordinance (which prohibit the importation, exportation and possession of controlled drugs) shall not have effect in relation to the controlled drugs specified in Schedule 1.

(2) Sections 6(1) (which prohibits the production and supply of controlled drugs) and 7(1) of the Ordinance shall not have effect in relation to poppy-straw.

Licences to produce, etc., controlled drugs.

5. Where any person is authorized by a licence of the Governor issued under this regulation and for the time being in force to produce, supply, offer to supply or have in his possession any controlled drug, it shall not by virtue of section 6(1) or 7(1) of the Ordinance be unlawful for that person to produce, supply, or have in his possession that drug in accordance with the terms of the licence and in compliance with any conditions attached to the licence.

General authority to possess.

6. Any of the following persons may, notwithstanding the General provisions of section 7(1) of the Ordinance have any controlled drug authority in his possession, that is to say–

(a) a police officer when acting in the course of his duty;

(b) a person engaged in the business of a carrier when acting in the course of that business;

(c) a person engaged in the business of the Post Office when acting in the course of that business;

(d) a revenue officer when acting in the course of his duty;

(e) a person engaged in the work of any laboratory to which the drug has been sent for forensic examination when acting in the course of his duty as a person so engaged;

(f) a person engaged in conveying the drug to a person authorized by these Regulations to have it in his possession.

Administration of drugs in Schedules 1 and 2.

7. (1) Any person may administer to another any controlled drug specified in Schedule 1.

(2) A practitioner may administer to a patient any controlled drug specified in Schedule 2.

(3) Any person other than a practitioner may administer to a patient, in accordance with the directions of a practitioner, any controlled drug specified in Schedule 2.

Production and supply of drugs in Schedules 1 and 2.

8. (1) Notwithstanding the provisions of section 6(1)(a) of the Ordinance–

(a) a practitioner or pharmacist, acting in his capacity as such, may manufacture or compound any controlled drug specified in Schedule 1 or 2; and

(b) a person lawfully conducting a registered dispensary and acting in his capacity as such may, at the registered dispensary at which he carries on business, manufacture or compound any controlled drug specified in Schedule 1 or 2.

(2) Notwithstanding the provisions of section 6(1)(b) of the Ordinance any of the following persons, that is to say–

(a) a practitioner;

(b) a pharmacist;

(c) a person lawfully conducting a registered dispensary;

(d) the matron or acting matron of a hospital or nursing home which is wholly or mainly maintained by a public authority out of public funds or by a charity or by voluntary subscriptions;

(e) in the case of such a controlled drug supplied to her by a person responsible for the dispensing and supply of medicines at the hospital or nursing home, the sister or acting sister for the time being in charge of a ward, theatre or other department in such a hospital or nursing home;

(f) a person who is in charge of a laboratory the recognised activities of which consist in, or include, the conduct of scientific education or research and which is attached to a hospital or to any other institution approved for the purpose by the Governor;

(g) a public analyst appointed under section 32 of the Food and Drugs Ordinance[5];

(h) a sampling officer within the meaning of section 34 of the Food and Drugs Ordinance[5];

may, when acting in his capacity as such, supply or offer to supply any controlled drug specified in Schedule 1 or 2 to any person who may lawfully have that drug in his possession:

Provided that nothing in this subregulation authorizes–

(i) the matron or acting matron of a hospital or nursing home, having a pharmacist responsible for the dispensing and supply of medicines, to supply or offer to supply any controlled drug;

(ii) a sister or acting sister for the time being in charge of a ward, theatre or other department to supply any controlled drug otherwise than for administration to a patient in that ward, theatre or department in accordance with the directions of a doctor or dentist.

(3) Notwithstanding the provisions of section 6(1)(b) of the Ordinance the owner or master of a ship which does not carry a doctor on board as part of her complement, may supply or offer to supply any controlled drug specified in Schedules 1 and 2–

(i) to any member of the crew;

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[5] **1964-07**

(ii) to any person who may lawfully supply that controlled drug;

(iii) to any police officer for the purpose of destruction.

Possession of drugs in Schedule 2.

9. (1) Notwithstanding the provisions of section 7(1) of the Ordinance a person specified in regulation 8(2) may have in his possession any controlled drug specified in Schedule 2 for the purpose of acting in his capacity as such.

(2) Notwithstanding the provisions of section 7(1) of the Ordinance a person may have in his possession any controlled drug specified in Schedule 2 for administration for medical, dental or veterinary purposes in accordance with the directions of a practitioner:

Provided that this subregulation shall not have effect in the case of a person to whom the controlled drug has been supplied by or on the prescription of a medical practitioner if–

(a) that person was then being supplied with any controlled drug by or on the prescription of another medical practitioner and failed to disclose that fact to the first mentioned medical practitioner before the supply by him or on his prescription; or

(b) that person or any other person on his behalf made a declaration or statement, which was false in any particular, for the purpose of obtaining the supply or prescription.

(3) Notwithstanding the provisions of section 7(1) of the Ordinance a person whose name is for the time being entered in the register kept for the purposes of this subregulation by the Public Health Specialist in Community Medicine may, in compliance with any conditions subject to which his name is so entered, have in his possession any drug specified in Schedule 2.

(4) Notwithstanding the provisions of section 7(1) of the Ordinance–

(a) the owner or master of a ship which does not carry a doctor on board as part of her complement, may have in his possession any drug specified in Schedule 2, so far as necessary for the purpose of compliance with the Merchant Shipping Ordinance[6];

[6] **1935-09**

(b) the master of a foreign ship which is in port in Gibraltar may have in his possession any controlled drug specified in Schedule 2 so far as necessary for the equipment of the ship.

Exemption for midwives in respect of pethidine.

10. (1) Notwithstanding the provisions of sections 6(1)(b) and 7(1) of the Ordinance a registered midwife, may, subject to the provisions of this regulation–

(a) so far as necessary for the practice of her profession or employment as a midwife, have pethidine in her possession;

(b) so far as necessary as aforesaid, administer pethidine; and

(c) surrender to the Public Health Specialist in Community Medicine any stocks of pethidine in her possession which are no longer required by her.

(2) Nothing in subregulation (1) authorizes a midwife to have in her possession pethidine which has been obtained other than on an order in writing of the Public Health Specialist in Community Medicine**,** or a medical practitioner authorized by him in writing to issue such orders, specifying the name of the midwife obtaining the pethidine, the purpose for which it is required and the quantity to be obtained.

Documents to be obtained by supplier of controlled drugs.

11.(1) Where a person (hereafter in this subregulation referred to as "the supplier"), not being a practitioner, supplies a controlled drug otherwise than on a prescription, the supplier shall not deliver the drug to a person who–

(a) purports to be sent by or on behalf of the person to whom it is supplied (hereafter in this subregulation referred to as "the recipient"); and

(b) is not authorized by any provision of these Regulations other than the provisions of regulation 6(f) to have that drug in his possession,

unless that person produces to the supplier a statement in writing signed by the recipient to the effect that he is empowered by the recipient to receive that drug on behalf of the recipient, and the supplier is reasonably satisfied that the document is a genuine document.

(2) Where a person (hereafter in this subregulation referred to as "the supplier") supplies a controlled drug, otherwise than on a prescription or by

way of administration, to any of the persons specified in subregulation (4), the supplier shall not deliver the drug–

(a) until he has obtained a requisition in writing which–

(i) is signed by the person to whom the drug is supplied (hereafter in this subregulation referred to as "the recipient");

(ii) states the name, address and profession or occupation of the recipient;

(iii) specifies the purpose for which the drug supplied is required and the total quantity to be supplied; and

(iv) where appropriate, satisfies the requirements of subregulation (5); and

(b) unless he is reasonably satisfied that the signature is that of the person purporting to have signed the requisition and that that person is engaged in the profession or occupation specified in the requisition:

Provided that where the recipient is a practitioner and he represents that he urgently requires a controlled drug for the purpose of his profession, the supplier may, if he is reasonably satisfied that the recipient so requires the drug and is, by reason of some emergency, unable before delivery to furnish to the supplier a requisition in writing duly signed, deliver the drug to the recipient on an undertaking by the recipient to furnish such a requisition within the twenty-four hours next following.

(3) A person who has given such an undertaking shall deliver to the person by whom the controlled drug was supplied a signed requisition in accordance with the undertaking.

(4) The persons referred to in subregulation (2) are–

(a) a practitioner;

(b) the matron or acting matron of a hospital or nursing home;

(c) a person who is in charge of a laboratory the recognised activities of which consist in, or include, the conduct of scientific education or research;

(d) the owner or master of a ship which does not carry a doctor on board as part of her complement; and

(e) the master of a foreign ship in port in Gibraltar.

(5) A requisition furnished for the purposes of subregulation (2) shall–

(a) where furnished by the matron or acting matron of a hospital or nursing home, be signed by a medical practitioner or dentist employed or engaged in that hospital or nursing home;

(b) where furnished by the master of a foreign ship, contain a statement, signed by a medical practitioner that the quantity of the drug to be supplied is the quantity necessary for the equipment of the ship.

(6) Where the person responsible for the dispensing and supply of medicines at any hospital or nursing home supplies a controlled drug to the sister or acting sister for the time being in charge of any ward, theatre or other department in that hospital or nursing home (hereafter in this subregulation referred to as "the recipient") he shall–

(a) obtain a requisition in writing, signed by the recipient, which specifies the total quantity of the drug to be supplied; and

(b) mark the requisition in such manner as to show that it has been complied with,

and any requisition obtained for the purposes of this subregulation shall be retained in the dispensary at which the drug was supplied and a copy of the requisition or a note of it shall be retained or kept by the recipient.

(7) Nothing in this regulation shall have effect in relation to the controlled drugs specified in Schedule 1 or poppy-straw.

Form of prescription.

12. (1) Subject to the provisions of this regulation, a person shall not issue a prescription containing a controlled drug other than a drug specified in Schedule 1 unless the prescription complies with the following requirements, that is to say, it shall–

(a) be in ink or otherwise so as to be indelible and be signed by the person issuing it with his usual signature and dated by him;

(b) in so far as it specifies the information required by paragraphs (e) and (f) below to be specified, be written by the person issuing it in his own handwriting;

(c) specify the address of the person issuing it;

(d) have written thereon, if issued by a dentist, the words "for dental treatment only" and, if issued by a veterinary practitioner, the words "for animal treatment only";

(e) specify the name and address of the person for whose treatment it is issued or, if it is issued by a veterinary practitioner, of the person to whom the controlled drug prescribed is to be delivered;

(f) specify the dose to be taken and–

(i) in the case of a prescription containing a controlled drug which is a preparation, the form and, where appropriate, the strength of the preparation, and either the total quantity (in both words and figures) of the preparation or the number (in both words and figures) of dosage units, as appropriate, to be supplied;

(ii) in any other case, the total quantity (in both words and figures) of the controlled drug to be supplied;

(g) in the case of a prescription for a total quantity intended to be dispensed by instalments, contain a direction specifying the amount of the instalments of the total amount which may be dispensed and the intervals to be observed when dispensing.

(2) Subregulation (1)(b) shall not have effect in relation to a prescription issued by a person approved for the purposes of this subregulation by the Public Health Specialist in Community Medicine .

(3) In the case of a prescription issued for the treatment of a patient in a hospital or nursing home, it shall be a sufficient compliance with subregulation (1)(e) if the prescription is written on the patient's bed card or case sheet.

Provisions as to supply on prescription.

13.(1) A person shall not supply a controlled drug, other than a controlled drug specified in Schedule 1, on a prescription–

(a) unless the prescription complies with the provisions of regulation 12;

(b) unless the address specified in the prescription as the address of the person issuing it is an address within Gibraltar;

(c) unless he either is acquainted with the signature of the person by whom it purports to be issued and has no reason to suppose

that it is not genuine, or has taken reasonably sufficient steps to satisfy himself that it is genuine;

(d) before the date specified in the prescription;

(e) subject to subregulation (3), later than thirteen weeks after the date specified in the prescription.

(2) Subject to subregulation (3), a person dispensing a prescription containing a controlled drug, other than a drug specified in Schedule 1, shall, at the time of dispensing it, mark thereon the date on which it is dispensed and shall retain it on the premises on which it was dispensed:

Provided that in the case of a health prescription the prescription shall be sent to the Public Health Specialist in Community Medicine .

(3) In the case of a prescription containing a controlled drug, other than a drug specified in Schedule 1, which contains a direction that specified instalments of the total amount may be dispensed at stated intervals, the person dispensing it shall not supply the drug otherwise than in accordance with that direction and–

(a) subregulation (1) shall have effect as if for the requirement contained in subparagraph (e) thereof there were substituted a requirement that the occasion on which the first instalment is dispensed shall not be later than thirteen weeks after the date specified in the prescription; and

(b) subregulation (2) shall have effect as if for the words “at the time of dispensing it” there were substituted the words “on each occasion on which an instalment is dispensed”.

Marking of bottles and other containers.

14.(1) Subject to subregulation (2), no person shall supply a controlled drug otherwise than in a bottle, package or other container which is plainly marked–

(a) in the case of a controlled drug other than a preparation, with the amount of the drug contained therein;

(b) in the case of a controlled drug which is a preparation–

(i) made up into tablets, capsules or other dosage units, with the amount of each component (being a controlled drug) of the preparation in each dosage unit and the number of dosage units in the bottle, package or other container;

(ii) not made up as aforesaid, with the total amount of the preparation in the bottle, package or other container and the percentage of each of its components which is a controlled drug.

(2) Nothing in this regulation shall have effect in relation to the controlled drugs specified in Schedule 1 or poppy-straw or in relation to the supply of a controlled drug by or on the prescription of a practitioner.

Keeping of registers.

15. (1) Subject to subregulation (3) and regulation 17, every person authorized by or under regulation 5 or 8 to supply any controlled drug specified in Schedule 2 shall comply with the following requirements, that is to say–

(a) he shall, in accordance with the provisions of this regulation and of regulation 16, keep a register and shall enter therein in chronological sequence in the form specified in Part I or Part II of Schedule 4 as the case may require, particulars of every quantity of a controlled drug specified in Schedule 2 or 3 obtained by him and of every quantity of such a drug supplied (whether by way of administration or otherwise) by him whether to persons within or outside Gibraltar; and

(b) he shall use a separate register or separate part of the register for entries made in respect of each class of controlled drugs, and each of the controlled drugs specified in paragraphs 1, 3 and 6 of Schedule 2 together with its salts and any preparation or other product containing it or any of its salts shall be treated as a separate class, so however that any stereoisomeric form of a drug or its salts shall be classed with that drug.

(2) Nothing in subregulation (1) shall be taken as preventing the use of a separate section within a register or separate part of a register in respect of different controlled drugs or strengths of such drugs comprised within the class of controlled drugs to which that register or separate part relates.

(3) The foregoing provisions of this regulation shall not have effect in relation to–

(a) a person licensed under regulations 5 to supply any controlled drug, where the licence so directs; or

(b) the sister or acting sister for the time being in charge of a ward, theatre or other department in a hospital or nursing home.

Requirements as to registers.

16. Any person required to keep a register under regulation 15 shall comply with the following requirements, that is to say–

(a) the class of controlled drugs to which the entries on any page of any such register relate shall be specified at the head of that page;

(b) every entry required to be made under regulation 15 in such a register shall be made on the day on which the controlled drug is obtained or, as the case may be, on which the transaction in respect of the supply of the drug by the person required to make the entry takes place or, if that is not reasonably practicable, on the day next following that day;

(c) no cancellation, obliteration or alteration of any such entry shall be made, and a correction of such an entry shall be made only by way of marginal note or footnote which shall specify the date on which the correction is made;

(d) every such entry and every correction of such an entry shall be made in ink or otherwise so as to be indelible;

(e) such a register shall not be used for any purpose other than the purposes of these Regulations;

(f) the person so required to keep such a register shall on demand made by the Public Health Specialist in Community Medicine or by any person authorized in writing by the Public Health Specialist in Community Medicine in that behalf–

(i) furnish such particulars as may be required in respect of the obtaining or supplying by him of any controlled drug specified in Schedule 2 or 3 or in respect of any stock of such drugs in his possession;

(ii) for the purpose of confirming any such particulars, produce any stock of such controlled drugs in his possession;

(iii) produce the register and such other books or documents in his possession relating to any dealings in controlled drugs specified in Schedule 2 or 3 as may be requested;

(g) a separate register shall be kept in respect of each premises at which the person required to keep the register carries on his business or occupation, but subject to that not more than one register shall be kept at one time in respect of each class of

drug in respect of which he is required to keep a separate register, so, however, that a separate register may, with the approval of the Public Health Specialist in Community Medicine, be kept in respect of each department of the business carried on by him; and

(h) every such register in which entries are currently being made shall be kept at the premises to which it relates.

Record-keeping requirements in particular cases.

17. (1) Where a controlled drug specified in Schedule 2 is supplied in accordance with regulation 8(3) to a member of the crew of a ship, an entry in the official log book required to be kept under the Merchant Shipping Ordinance or, in the case of a ship which is not required to carry such an official log book, a report signed by the master of the ship, shall, notwithstanding anything in these Regulations, be a sufficient record of the supply if the entry or report specifies the drug supplied and, in the case of a report, it is delivered as soon as may be to the Captain of the Port.

(2) A midwife authorized by regulation 10(1) to have pethidine in her possession shall–

(a) on each occasion on which she obtains a supply of pethidine, enter in a book kept by her and used solely for the purposes of this subregulation the date, the name and address of the person from whom the drug was obtained, the amount obtained and the form in which it was obtained; and

(b) on administering pethidine to a patient, enter in the said book as soon as practicable the name and address of the patient, the amount administered and the form in which it was administered.

Preservation of registers, books and other documents.

18. (1) All registers and books kept in pursuance of regulations 15 or 17 (2) shall be preserved for a period of two years from the date on which the last entry therein is made.

(2) Every requisition, order or prescription on which a controlled drug is supplied in pursuance of these Regulations shall be preserved for a period of two years from the date on which the last delivery under it was made.

Preservation of records relating to drugs in Schedule 1.

19. (1) A producer of any controlled drug specified in Schedule 1 and a wholesale dealer in any such drug shall keep every invoice or other like

record issued in respect of each quantity of such a drug obtained by him and in respect of each quantity of such a drug supplied by him.

(2) A retail dealer in any controlled drug specified in Schedule I shall keep every invoice or other like record issued in respect of each quantity of such a drug obtained by him.

(3) Every document kept in pursuance of this regulation shall be preserved for a period of two years from the date on which it is issued:

Provided that the keeping of a copy of the document made at any time during those two years shall be treated for the purposes of this subregulation as if it were the keeping of the original document.

Destruction of controlled drugs.

20. (1) No person who is required by any provision of, or by any term or condition of a licence having effect under, these Regulations to keep records with respect to a controlled drug specified in Schedule 2 or 3 shall destroy such a drug or cause such a drug to be destroyed except in the presence of and in accordance with any directions given by a person authorized for the purposes of this subregulation by the Director (hereafter in this regulation referred to as an "authorized person").

(2) An authorized person may, for the purpose of analysis, take a sample of a controlled drug specified in Schedule 2 or 3 which is to be destroyed.

(3) Where a controlled drug specified in Schedule 2 or 3 is destroyed in pursuance of subregulation (1) by or at the instance of a person who is required by any provision of, or by any term or condition of a licence having effect under, these Regulations to keep a record in respect of the obtaining or supply of that drug, that record shall include particulars of the date of destruction and the quantity destroyed and shall be signed by the authorized person in whose presence the drug is destroyed.

(4) Where the master or owner of a ship has in his possession a controlled drug specified in Schedule 2 which he no longer requires, he shall not destroy the drug or cause it to be destroyed but shall dispose of it to a person who may lawfully supply it.

Forms.

21. The Forms contained in Schedule 5 shall be used for the purposes to which they relate.

Transitional provisions.

22. *Omitted.*

Repeal.

23. *Omitted.*

DRUGS (MISUSE) REGULATIONS
SCHEDULE 1

Regulations 4, 7 ,8, 11, 12, 13, 14, and 19

CONTROLLED DRUGS EXCEPTED FROM THE PROHIBITION ON IMPORTATION, EXPORTATION AND POSSESSION AND SUBJECT TO THE REQUIREMENTS OF REGULATION 19.

1. (1) Any preparation of one or more of the substances to which this paragraph applies, not being a preparation designed for administration by injection, when compounded with one or more other active or inert ingredients and containing a total of not more than 100 milligrammes of the substance or substances (calculated as base) per dosage unit and with a total concentration of not more than 2.5 per cent (calculated as base) in undivided preparations.

(2) The substances to which this paragraph applies are acetyldihydrocodeine, codeine, dihydrocodeine, ethylmorphine, nicocodine, nicodicodine (6 nicotinoyldihydrocodeine), norcodeine pholcodine and their respective salts.

2. Any preparation of cocaine containing not more than 0.1 per cent of cocaine calculated as cocaine base, being a preparation compounded with one or more other active or inert ingredients in such a way that the cocaine cannot be recovered by readily applicable means or in a yield which constitute a risk to health.

3. Any preparation of medicial opium or of morphine containing (in either case) not more than 0.2 per cent, of morphine calculated as anhydrous morphine base, being a preparation compounded with one or more other active or inert ingredients in such a way that the opium or, as the case may be, the morphine, cannot be recovered by readily applicable means or in a yield which would constitute a risk to health.

4. Any preparation of diphenoxylate containing, per dosage unit, not more than 2.5 milligrammes of diphenoxylate calculated as base, and a quantity of atropine sulphate equivalent to at least 1 per cent of the dose of diphenoxylate.

5. Any powder of ipecacuanha and opium comprising 10 percent, opium, in powder, 10 per cent, ipecacuanha root, in powder, well mixed with 80 per cent of any other powdered ingredient containing no controlled drug.

6. Any mixture containing one or more of the preparations specified in paragraphs 1 to 5, being a mixture of which none of the other ingredients is a controlled drug.

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DRUGS (MISUSE) REGULATIONS
SCHEDULE 2

Regulations 7, 8, 9, 15, 16 and 20

CONTROLLED DRUGS SUBJECT TO THE REQUIREMENTS OF REGULATIONS 11, 12, 13, 14, 15, 16, 17 AND 20.

I. The following substances and products, namely–

Acetorphine
Allylprodine
Alphacetylmethadol
Alphameprodine
Alphamethadol
Alphaprodine
Anileridine
Benzethidine
Benzylmorphine
(3-benzylmorphine)
Betacetylmethadol
Betameprodine
Betamethadol
Betaprodine
Bezitramide
Clonitazene
Cocaine
Desomorphine
Dextromoramide
Diamorphine
Diampromide
Diethylthiambutene
Hydrocodone
Hydromorphinol
Hydromorphone
Dihydrocodeinon
O-carboxymethyloxime
Dihydromorphine
Dimenoxadole
Dimepheptanol
Dimethylthiambutene
Dioxaphetyl butyrate
Diphenoxylate
Dipipanone
Drotebanol (3, 4-dimethoxy-
17-methylmorphinan-6B, 14-diol)
Ecgonine, and any derivative
of ecgonine which is
convertible to ecgonine or to cocaine

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Ethylmethylthiambutene
Etonitazene
Etorphine
Etoxeridine
Fentanyl
Furethidine
Norpipanone
Oxycodone
Oxymorphone
Hydroxypethidine
Isomethadone
Ketobemidoine
Levomethorphan
Levomoramide
Levophenacylmorphan
Levorphanol
Medicinal opium
Metazocine
Methadone
Methadyl acetate
Methyldesorphine
Methyldihydromorphine,
(6-methyldihydromor-phine)
Metopon
Morpheridine
Morphine
Morphine methobromide,
morphine N-oxide and
other pentavalent nitrogen
morphine derivatives
Myrophine
Nicomorphine
Noracymethadol
Norlevorphanol
Normethadone
Normorphine
Pethidine
Phenadoxone
Phenampromide
Phenazocine
Phenomorphan
Phenoperidine
Piminodine
Piritramide
Proheptazine
Properidine
Racemethorphan
Racemoramide

Racemorphan
Thebacon
Thebaine
Trimeperidine
4-Cyano-2-dimethylamino-4,
4-diphenylbutane
4-Cyano-1-methyl-
4-phenylpiperidine
1-Methyl-4-phenylpiperdine-
4-carboxylic acid
2-Methyl-3-morpholino-1,
1 -diphenylpropanecar-
boxylic acid
4-Phenylpiperidine
4-carboxylic acid
ethyl ester

2. Any stereoisomeric form of a substance specified in paragraph 1 not being dextromethorphan or dextrorphan.

3. Any ester or ether of a substance specified in paragraph 1 or 2, not being a substance specified in paragraph 6.

4. Any salt of a substance specified in any of paragraphs 1 to 3.

5. Any preparation or other product containing a substance or product specified in any of paragraphs 1 to 4, not being a preparation specified in Schedule 1.

6. The following substances and products, namely–

Acetyldihydrocodeine
Amphetamine
Benzphetamine
Chiorphentermine
Codeine
Dexamphetamine
Diethylpropion
Dihydro codeine
Ethylmorphine (3-ethyl- morphine)
Mephentermine
Methaqualone
Methylamphetamine
Methyiphenidate
Nicocodine
Nicodicodine (6-nicotinoyldi-hydrocodeine)
Norcodeine

Phendimetrazine
Phenmetrazine
Pholcodine
Pipradrol
Propiram.

7. Any stereoisomeric form of substance specified in paragraph 6.

8. Any salt of a substance specified in paragraphs 6 or 7.

9. Any preparation or other product contatining a substance or product specified in any of paragraphs 6 to 8 not being a preparation specified in Schedule.

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DRUGS (MISUSE) REGULATIONS

SCHEDULE 3

Regulations 15 and 20

CONTROLLED DRUGS SUBJECT TO THE REQUIREMENTS OF REGULATIONS 11, 12, 13, 14, 15, l6 AND 20.

1. The following substances and products, namely–

Bufotenine
Cannabinol
Cannabinol derivatives
Cannabis and cannabis resin
Coca leaf
Concentrate of poppy-straw
Lysergamide
Lysergide and other N-alkyl derivatives of lysergamide
Mescaline
Raw opium
Psilocin
N,N-Diethyltryptamine
N ,N-Dimethyltryptamine, and
2,5-Dimethoxy-a, 4-dimethylphenethylamine.

2. Any stereoisomeric form of a substance specified in paragraph 1.

3. Any ester or ether of a substance specified in paragraph 1 or 2.

4. Any salt of a substance specified in any of paragraphs 1 to 3.

5. Any preparation or other product containing a substance or product specified in any of paragraphs 1 to 4, not being a preparation specified in Schedule 1.

DRUGS (MISUSE) REGULATIONS
SCHEDULE 4

Regulation 15

FORM OF REGISTER

PART I

ENTRIES TO BE MADE IN CASE OF OBTAINING

Date on which supply received	Name	Address	Amount	Form in which obained
	Of person or firm from whom obtained			

PART II
ENTRIES TO BE MADE IN CASE OF SUPPLY.

Date on which the transaction was effected	Name	Address	Particulars as to licence or authority of person or firm supplied to be in possession	Amount supplied	Form in which supplied
	Of person or firm supplied				

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DRUGS (MISUSE) REGULATIONS
SCHEDULE 5.

Regulation 21.

FORM A.

DRUGS (MISUSE) ORDINANCE.

Licence No .....…….....
File No.....…….....…...
Applicant's .....…….....
Refce No.....…… ...….....

LICENCE TO IMPORT CONTROLLED DRUGS.

In accordance with the provisions of the Drugs (Misuse) Ordinance and of the powers delegated to him by the Governor, the Specialist in Community Medicine of the Gibraltar Health Authority (hereinafter called `the Specialist') hereby grants to
(hereinafter called `the licensee') a licence to import from the following controlled drugs, namely:
subject to the following conditions:

l. The licence does not authorise the licensee, to obtain, be in possession of or deal with, the controlled drugs named herein other than in accordance with the provisions of the Ordinance.

2. The licence is available only for the controlled drugs of the exact quantity, kind and form specified above.

3. The licence does not relieve the importer from compliance with any regulations in force for the time being relating to the importation of goods into, or transhipment of goods in, Gibraltar or with any Post Office regulations for the time being in force in Gibraltar nor from any rules or regulations respecting the transmission of articles by post which may for the time being be in force, whether within Gibraltar or elsewhere.

4. The licence is valid only for the licensee and may be revoked at any time by the Specialist, to whom it shall in that event be immediately surrended. It shall be produced for inspection when required by any duly authorised officer.

5. The licence, unless sooner revoked, shall continue in force for six calendar months from the date hereof. It shall be produced to the Specialist at the time of importation and when the last consignment of drugs is imported. If not used it shall be surrended to the Specialist within seven days of its expiry.

6. The copy of the export licence, if any, which accompanies the drugs shall be forwarded to the Specialist immediately the importation of the drugs has been effected.

Date ....…….....

Specialist in Community Medicine
Gibraltar Health Authority
St. Bernard's Hospital
Gibraltar

Note: If any alteration is desired in this licence it must be returned with request for amendment and a statement of the reason therefor. No unauthorised alteration is permissible.

FORM B.

DRUGS (MISUSE) ORDINANCE.

Licence No .....…….....
File No.....…….....
Applicant's .....…….....
Refce No ...…….....

LICENCE TO EXPORT CONTROLLED DRUGS.

In accordance with the provisions of the Drugs (Misuse) Ordinance and the powers delegated to him by the Governor, the Specialist in Community Medicine of the Gibraltar Health Authority (hereinafter called `the Specialist') hereby authorises

(hereinafter called `the licensee') a licence to export from: 'the port/airport of Gibraltar by
Gibraltar by parcel post in
parcels from the General Post Office at Gibraltar
to
by virtue of Import Certificate No.: dated issued by
the following controlled drugs, namely:

subject to the following conditions-

1. The licence does not authorise the licensee to obtain, be in possession of or deal with the controlled drugs named herein other than in accordance with the provisions of the Ordinance.

2. The licence is available only for controlled drugs of the exact quantity, kind and form specified above.

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DRUGS (MISUSE) REGULATIONS

3. The licence does not relieve the licensee from compliance with any regulations in force for the time being relating to the exportation of goods from Gibraltar nor from any provision of the Post Office Ordinance, or of any Post Office regulations for the time being in force, nor from any rules or regulations respecting the transmission of articles by post which may for the time being be in force, whether within Gibraltar or elsewhere.

4. If the drugs are authorised to be exported by ship the attached duplicate copy shall accompany the consignment to the place of destination and for this purpose the licensee shall cause it to be delivered to the master of the vessel by which the consignment is despatched. (See Note 2).

5. If the drugs are authorised to be exported by post the attached duplicate copy shall be placed inside the outer wrapper of the parcel containing the drugs. If the drugs are contained in more than one parcel, the duplicate copy shall be placed inside the outer wrapper of one of them; the parcels shall be consecutively numbered on the outer wrapper, and on each parcel there shall be legibly stated the number of the parcel in which the duplicate copy is to be found. (See Note 3).

6. The licensee, if so required by the Specialist, shall produce to him within such time as he may allow, proof to his satisfaction that the drugs were at the destination named in this licence, and in the event of non-compliance with this condition the authorisation shall be deemed void and of no effect.

7. The licensee shall furnish to the Specialist such returns of the goods exported by him in pursuance of this licence as may from time to time be required.

8. The licence is valid only for the licensee and may be revoked at any time by the Specialist. It shall he produced for inspection when required by any duly authorised officer.

9. The licence, unless sooner revoked, shall continue in force for. three calendar months from the date hereof. It must be produced at the time of export to

*the customs officer at
*the Director of Postal Services,

who will retain it. If not used it shall be surrendered to the Specialist within seven days of the date of its expiry.

Date ..…….....

Specialist in Community Medicine
Gibraltar Health Authority
St. Bernard's Hospital

DRUGS (MISUSE) REGULATIONS

Gibraltar

**Strike out the words not applicable.*

Note: 1, If any alteration is desired in this licence it must be returned with a request for amendment and a statement of the reasons therefor. No unauthorised alteration is permissible.

2. In the case of drugs exported by ship, this document is required in pursuance of Article IS of the International Opium Convention 1925 and Article 31 of the Single Convention on Narcotic drugs 1961 to be produced to the competent authorities of any country through which the consignment passes, whether it is transhipped or not. Failure to comply with condition 4 of the licence may lead to delay or confiscation of the consignment.

3. In the case of drugs exported by post, failure to comply with condition 5 of the licence may lead to delay or confiscation of the parcels in the country of destination.

ENDORSEMENT.
(by Government of Imparting Country).

Please return to:

THE SPECIALIST IN COMMUNITY MEDICINE
GIBRALTAR HEALTH AUTHORITY
ST, BERNARD'S HOSPITAL
GIBRALTAR.

Certified that the controlled drugs detailed in this licence have been duly imported.

Signed .....…….....