The Misuse of Drugs (Bailiwick of Guernsey) Ordinance,

1997

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The Misuse of Drugs (Bailiwick of Guernsey) Ordinance, 1997

THE STATES, in pursuance of their Resolution of 29th March, 1989, and in exercise of the powers conferred upon them by sections 6, 9, 21 and 30 of the Misuse of Drugs (Bailiwick of Guernsey) Law, 1974, as amended, hereby order:

Interpretation

1. In this Ordinance, unless the context otherwise requires:

"authorised analyst" means a person employed by the States of Guernsey or by the States of Alderney and approved by or on behalf of the Board as an analyst of drugs;

"authorised as a member of a group" means authorised by virtue of being a member of a class as respects which the Board has granted an authority which is for the time being in force under and for the purposes of section 8(3), 9(3) or 10(3); and "his group authority", in relation to a person who is a member of such a class, means the authority so granted to that class;

"Chief Officer of Police" means the Chief Officer of the salaried police force of the Island of Guernsey

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\(a\) Article 13 of Billet d'État No. VII of 1989.

"hospital" means an infirmary or other medical institution wholly or mainly maintained by the States of Guernsey or by the States of Alderney;

"the Law" means the Misuse of Drugs (Bailiwick of Guernsey) Law, 1974;

"master" includes every person (except a general pilot) having command or charge of a ship, and, in relation to a fishing vessel, means the skipper;

"Medical Officer of Health" means the Medical Officer of Health of the States of Guernsey and includes any Deputy Medical Officer of Health of the States of Guernsey;

"medical prescription" means a prescription issued by a medical practitioner or a dentist under Part V of the Health Service (Benefit) (Guernsey) Law, 1990;

"medicinal product" has the same meaning as in the Medicines Act 1968;

"the Merchant Shipping Laws", in relation to any ship, means the Loi relative à la Marine Marchande of 1916, and so much of the Merchant Shipping Acts 1894 to 1994 and the Merchant Shipping Act 1995 as applies to that ship;

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"midwife's supply order" means an order in writing specifying the name and occupation of the midwife obtaining a supply of a controlled drug, the purpose for which it is required and the total quantity to be supplied;

"nursing home" means a nursing home within the meaning of the Nursing Homes and Residential Homes (Guernsey) Law, 1976 or within the meaning of Part I of the Nursing and Residential Homes (Registration and Occupation) (Alderney) Law, 1987;

"officer of customs and excise" means an officer within the meaning of the Customs and Excise (General Provisions) (Bailiwick of Guernsey) Law, 1972, as amended;

"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 surgeon 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 midwife" mean a person for the time being authorised to practise as a midwife pursuant to the provisions of the Nurses, Midwives and Health Visitors Ordinance, 1987;

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h Recueil d'Ordonnances, Tome XXIV, p. 238.
"retail dealer" means a person lawfully conducting a retail pharmacy business;

"sampling officer" means a person authorised by the Board under the 
Food and Drugs (Guernsey) Law, 1970 to exercise such powers of procuring 
samples for analysis or for bacteriological or other examination as are conferred by 
section 26 of that Law;

"seamen" includes every person (except masters and general pilots) 
employed or engaged in any capacity on board any ship;

"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,

and other expressions have the same meanings as in the Law.

(2) In this Ordinance any reference to a section or Schedule is to 
the section or Schedule so numbered in this Ordinance; and any reference in a 
section or Schedule to a subsection or paragraph is to the subsection or paragraph so 
numbered in that section or Schedule.

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(3) In this Ordinance, unless the context otherwise requires, any reference to any other enactment is to that enactment as repealed and replaced, amended, extended or applied by or under any other enactment.

(4) The Interpretation (Guernsey) Law, 1948, as amended, applies to the interpretation of this Ordinance throughout the Bailiwick.

**Specification of controlled drugs for purposes of Ordinance**

2. Schedules 1 to 5 shall have effect for the purpose of specifying the controlled drugs to which certain provisions of this Ordinance apply.

**Exceptions for drugs in Schedules 4 and 5 and poppy-straw**

3. (1) Section 2(1) of the Law (which prohibits the importation and exportation of controlled drugs) shall not have effect in relation to the drugs specified in Schedules 4 and 5.

(2) Section 4(1) of the Law (which prohibits the possession of controlled drugs) shall not have effect in relation to:

   (a) any drug specified in Schedule 4 which is contained in a medicinal product;

   (b) the drugs specified in Schedule 5.

(3) Section 3(1) (which prohibits the production and supply of controlled drugs) and 4(1) of the Law shall not have effect in relation to poppy-straw.

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Licences to produce etc. controlled drugs

4. The Board may issue a licence under this section authorising a person to produce, supply, offer to supply or have in his possession any controlled drug; and where a person is for the time being so authorised it shall not by virtue of section 3(1) or 4(1) of the Law be unlawful for that person to produce, supply, offer to 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 supply and possess

5. (1) Notwithstanding the provisions of section 3(1)(b) of the Law, any person who is lawfully in possession of a controlled drug may supply that drug to the person from whom he obtained it.

(2) Notwithstanding the provisions of section 3(1)(b) of the Law, any person who has in his possession a drug specified in Schedule 2, 3, 4 or 5 which has been supplied by or on the prescription of a practitioner for the treatment of that person, or of a person whom he represents, may supply that drug to any medical practitioner, dentist or pharmacist for the purpose of its destruction.

(3) Notwithstanding the provisions of section 3(1)(b) of the Law, any person who is lawfully in possession of a drug specified in Schedule 2, 3, 4 or 5 which has been supplied by or on the prescription of a veterinary surgeon for the treatment of animals may supply that drug to any veterinary surgeon or pharmacist for the purpose of its destruction.

(4) Notwithstanding the provisions of section 3(1)(b) of the Law, any of the persons specified in paragraph (6) may supply any controlled drug to any person who may lawfully have that drug in his possession.
(5) Notwithstanding the provisions of section 4(1) of the Law, any of the persons so specified may have any controlled drug in his possession.

(6) The persons referred to in paragraphs (4) and (5) are:

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

(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 States Post Office when acting in the course of that business;

(d) an officer of customs and excise when acting in the course of his duty as such;

(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 who may lawfully have that drug in his possession.

Administration of drugs in Schedules 2, 3, 4 and 5

6. (1) Any person may administer to another any drug specified in Schedule 5.

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(2) A medical practitioner or dentist may administer to a patient any drug specified in Schedule 2, 3 or 4.

(3) Any person other than a medical practitioner or dentist may administer to a patient, in accordance with the directions of a medical practitioner or dentist, any drug specified in Schedule 2, 3 or 4.

**Production and supply of drugs in Schedules 2 and 5**

7. (1) Notwithstanding the provisions of section 3(1)(a) of the Law:

   (a) a practitioner or pharmacist, acting in his capacity as such, may manufacture or compound any drug specified in Schedule 2 or 5;

   (b) a person lawfully conducting a retail pharmacy business and acting in his capacity as such may, at the pharmacy at which he carries on that business, manufacture or compound any drug specified in Schedule 2 or 5.

(2) Notwithstanding the provisions of section 3(1)(b) of the Law, any of the following persons, that is to say:

   (a) a practitioner;

   (b) a pharmacist;

   (c) a person lawfully conducting a retail pharmacy business;
(d) the person in charge or acting person in charge of a hospital, or of a nursing home which is wholly or mainly maintained by the States of Guernsey or Alderney;

(e) in the case of such a 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 as aforesaid;

(f) an authorised analyst;

(g) a sampling officer;

may, when acting in his capacity as such, supply or offer to supply any drug specified in Schedule 2 or 5 to any person who may lawfully have that drug in his possession:

Provided that nothing in this subsection authorises:

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

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

(3) Notwithstanding the provisions of section 3(1)(b) of the Law, a person who is authorised as a member of a group may, under and in accordance with the terms of his group authority and in compliance with any conditions attached thereto, supply or offer to supply any drug specified in Schedule 2 or 5 to any person who may lawfully have that drug in his possession.

(4) Notwithstanding the provisions of section 3(1)(b) of the Law, a person who is authorised by a written authority issued by the Board under and for the purposes of this paragraph and for the time being in force may, at the premises specified in that authority and in compliance with any conditions so specified, supply or offer to supply any drug specified in Schedule 5 to any person who may lawfully have that drug in his possession.

(5) Notwithstanding the provisions of section 3(1)(b) of the Law, the owner of a ship, or the master of a ship which does not carry a medical practitioner among the seamen employed in it may supply or offer to supply any drug specified in Schedule 2 or 5:

(i) for the purpose of compliance with any of the provisions specified in paragraph (6), to any person on that ship;

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

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

(6) The provisions referred to in paragraph (5) are any provision of, or of any instrument which is in force under:

(a) the Merchant Shipping Laws;

(b) the Health and Safety at Work etc. (Guernsey) Law, 1979k.

Production and supply of drugs in Schedules 3 and 4

8. (1) Notwithstanding the provisions of section 3(1)(a) of the Law:

(a) a practitioner or pharmacist, acting in his capacity as such, may manufacture or compound any drug specified in Schedule 3 or 4;

(b) a person lawfully conducting a retail pharmacy business and acting in his capacity as such may, at the pharmacy at which he carries on that business, manufacture or compound any drug specified in Schedule 3 or 4;

(c) a person who is authorised by a written authority issued by the Board under and for the purposes of this subsection and for the time being in force may, at the
premises specified in that authority and in compliance with any conditions so specified, produce any drug specified in Schedule 3 or 4.

(2) Notwithstanding the provisions of section 3(1)(b) of the Law, any of the following persons, that is to say:

(a) a practitioner;

(b) a pharmacist;

(c) a person lawfully conducting a retail pharmacy business;

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

(e) an authorised analyst;

(f) a sampling officer;

(g) an inspector appointed for the purposes of the Poisons and Pharmacy Ordinance, 1970;  

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l Recueil d'Ordonnances, Tome XVI, p. 236; Tome XVII, p. 121; Tome XIX, pp. 73 and 195; Tome XX, p. 342; Tome XXI, p. 13; Tome XXII, p. 355; Tome XXIII, p. 427; Tome XXIV, p. 79.

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may, when acting in his capacity as such, supply or offer to supply any drug specified in Schedule 3 or 4 to any person who may lawfully have that drug in his possession.

(3) Notwithstanding the provisions of section 3(1)(b) of the Law:

(a) a person who is authorised as a member of a group, under and in accordance with the terms of his group authority and in compliance with any conditions attached thereto,

(b) the person in charge or acting person in charge of a hospital or nursing home,

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

may, when acting in his capacity as such, supply or offer to supply any drug specified in Schedule 3, or any drug specified in Schedule 4 which is contained in a medicinal product, to any person who may lawfully have that drug in his possession:

Provided that nothing in this subsection authorises:

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

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

(4) Notwithstanding the provisions of section 3(1)(b) of the Law:

(a) a person who is authorised by a written authority issued by the Board under and for the purposes of this subsection and for the time being in force may, at the premises specified in that authority and in compliance with any conditions so specified, supply or offer to supply any drug specified in Schedule 3 or 4 to any person who may lawfully have that drug in his possession;

(b) a person who is authorised under subsection (1)(c) may supply or offer to supply any drug which he may, by virtue of being so authorised, lawfully produce to any person who may lawfully have that drug in his possession.

(5) Notwithstanding the provisions of section 3(1)(b) of the Law the owner of a ship, or the master of a ship which does not carry a medical
practitioner among the seamen employed in it, may supply or offer to supply any
drug specified in Schedule 3, or any drug specified in Schedule 4 which is
contained in a medicinal product:

(i) for the purpose of compliance with any of the
provisions specified in section 7(6), to any
person on that ship; or

(ii) to any person who may lawfully supply that
drug to him.

(6) Notwithstanding the provisions of section 3(1)(b) of the Law,
a person in charge of a laboratory may, when acting in his capacity as such, supply
or offer to supply any drug specified in Schedule 3 which is required for use as a
buffering agent in chemical analysis to any person who may lawfully have that drug
in his possession.

Possession of drugs in Schedules 2, 3 and 4

9. (1) Notwithstanding the provisions of section 4(1) of the Law:

(a) a person specified in one of subsections (a) to (g) of
section 7(2) may have in his possession any drug
specified in Schedule 2;

(b) a person specified in one of subsections (a) to (e) of
section 8(2) may have in his possession any drug
specified in Schedule 3 or 4;
(c) a person specified in section 8(3)(b) or (c) or section 8(6) may have in his possession any drug specified in Schedule 3,

for the purpose of acting in his capacity as such a person:

Provided that nothing in this paragraph authorises:

(i) a person specified in subsection (e) of section 7(2);

(ii) a person specified in subsection (c) of section 8(3); or

(iii) a person specified in section 8(6),

to have in his possession any drug other than such a drug as is mentioned in the subsection in question specifying him.

(2) Notwithstanding the provisions of section 4(1) of the Law, a person may have in his possession any drug specified in Schedule 2 or 3 for administration for medical, dental or veterinary purposes in accordance with the directions of a practitioner:

Provided that this paragraph shall not have effect in the case of a person to whom the 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 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 4(1) of the Law, a person who is authorised as a member of a group may, under and in accordance with the terms of his group authority and in compliance with any conditions attached thereto, have any drug specified in Schedule 2 or 3 in his possession.

(4) Notwithstanding the provisions of section 4(1) of the Law:

(a) a person who is authorised by a written authority issued by the Board under and for the purposes of this subsection and for the time being in force may, at the premises specified in that authority and in compliance with any conditions so specified, have in his possession any drug specified in Schedule 3 or 4;

(b) a person who is authorised under section 8(1)(c) may have in his possession any drug which he may, by virtue of being so authorised, lawfully produce;

(c) a person who is authorised under section 8(4)(a) may have in his possession any drug which he may, by virtue of being so authorised, lawfully supply or offer to supply.
(5) Notwithstanding the provisions of section 4(1) of the Law:

(a) any person may have in his possession any drug specified in Schedule 2 or 3 for the purpose of compliance with any of the provisions specified in section 7(6);

(b) the master of a ship which is in a port in the Bailiwick may have in his possession any drug specified in Schedule 2 or 3 so far as necessary for the equipment of the ship.

(6) The foregoing provisions of this section are without prejudice to the provisions of section 3(2)(a).

Exemption for midwives

10. (1) Notwithstanding the provisions of sections 3(1)(b) and 4(1) of the Law, a registered midwife may, subject to the provisions of this section:

(a) so far as necessary to her professional practice, have in her possession,

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

(c) surrender to the Chief Pharmacist of the Board of Health such stocks in her possession as are no longer required by her of,
any controlled drug which she may lawfully administer under and in accordance with section 8 of the Nurses, Midwives and Health Visitors Ordinance, 1987 (g).

(2) Nothing in subsection (1) authorises a midwife to have in her possession a controlled drug which has been obtained otherwise than on a midwife's supply order signed by the designated officer within the meaning of the Nurses, Midwives and Health Visitors Ordinance, 1987.

Cultivation under licence of Cannabis plant

11. The Board may issue a licence under this section authorising a person to cultivate plants of the genus Cannabis; and where a person is for the time being so authorised it shall not by virtue of section 5 of the Law be unlawful for that person to cultivate any such plant in accordance with the terms of the licence and in compliance with any conditions attached to the licence.

Documents to be obtained by supplier of controlled drugs

12. (1) Where a person ("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 ("the recipient"); and

(b) is not authorised by any provision of this Ordinance other than the provisions of sections 5(5) and 5(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 ("the supplier") supplies a controlled drug, otherwise than on a prescription or by way of administration, to any of the persons specified in subsection (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 ("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 subsection (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 next 24 hours.

(3) A person who has given such an undertaking as aforesaid 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 subsection (2) are:

(a) a practitioner;

(b) the person in charge or acting person in charge of a hospital or nursing home;

(c) a person who is in charge of a laboratory;

(d) the owner of a ship, or the master of a ship which does not carry a medical practitioner among the seamen employed in it;

(e) the master of a ship which is in a port in the Bailiwick.

(5) A requisition furnished for the purposes of paragraph (2) shall:

(a) where furnished by the person in charge or acting person in charge 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 ship, contain a statement, signed by the Medical Officer of Health, 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 ("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 paragraph 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 section shall have effect in relation to:

(a) the drugs specified in Schedules 4 and 5 or poppy straw;
Form of prescriptions

13. (1) Subject to the provisions of this section, a person shall not issue a prescription containing a controlled drug, other than a drug specified in Schedule 4 or 5 or temazepam, unless the prescription complies with the following requirements, that is to say it shall:

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

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

(c) except in the case of a medical prescription, 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 surgeon, a declaration that the controlled drug is prescribed for an animal or herd under his care;

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

(f) specify the doses 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 supplied by instalments, contain a direction specifying the amount of the instalments of the total amount which may be supplied and the intervals to be observed when supplying.

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(2) Paragraph (1)(b) shall not have effect in relation to:

(a) a prescription issued by a person approved (whether personally or as a member of a class) for the purposes of this paragraph by the Board; or

(b) a prescription containing no controlled drug other than:

(i) phenobarbitone;

(ii) phenobarbitone sodium; or

(iii) a preparation containing a drug specified in paragraph (i) or (ii) above.

(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 subsection (1)(e) if the prescription is written on the patient's bed card or case sheet.

**Provisions as to supply on prescription**

14. (1) A person shall not supply a controlled drug other than a drug specified in Schedule 4 or 5 on a prescription:

(a) unless the prescription complies with the provisions of section 13;
(b) unless the address specified in the prescription as the address of the person issuing it is an address within the Bailiwick;

(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 paragraph (3), later than thirteen weeks after the date specified in the prescription.

(2) Subject to paragraph (3), a person supplying on prescription a controlled drug other than a drug specified in Schedule 4 or 5 shall, at the time of the supply, mark on the prescription the date on which the drug is supplied and, unless it is a medical prescription, shall retain the prescription on the premises from which the drug was supplied.

(3) In the case of a prescription containing a controlled drug other than a drug specified in Schedule 4 or 5, which contains a direction that specified instalments of the total amount may be supplied at stated intervals, the person supplying the drug shall not do so otherwise than in accordance with that direction and:

(a) subsection (1) shall have effect as if for the requirement contained in paragraph (e) thereof there were substituted a requirement that the occasion on

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which the first instalment is supplied shall not be later than thirteen weeks after the date specified in the prescription;

(b) subsection (2) shall have effect as if for the words "at the time of the supply" there were substituted the words "on each occasion on which an instalment is supplied".

Exemption for certain prescriptions

15. Nothing in sections 13 and 14 shall have effect in relation to a prescription issued for the purposes of the Food and Drugs (Guernsey) Law, 1970.

Marking of bottles and other containers

16. (1) Subject to paragraph (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 section shall have effect in relation to:

(a) the drugs specified in Schedules 4 and 5 or poppy-straw;

(b) any drug specified in Schedule 3 contained in or comprising a preparation which:

(i) is required for use as a buffering agent in chemical analysis,

(ii) has present in it both a substance specified in paragraph 1 or 2 of that Schedule and a salt of that substance, and

(iii) is premixed in a kit;

(c) the supply of a controlled drug by or on the prescription of a practitioner.

Record-keeping requirements in respect of drugs in Schedules 1 and 2

17. (1) Subject to subsection (3) and section 19, every person authorised by or under section 4 or 7 to supply any drug specified in Schedule 1 or 2 shall comply with the following requirements, that is to say:
(a) he shall, in accordance with the provisions of this section and of section 18, keep a register and shall enter therein in chronological sequence in the form specified in Part I or Part II of Schedule 6, as the case may require, particulars of every quantity of a drug specified in Schedule 1 or 2 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 the Bailiwick;

(b) he shall use a separate register or separate part of the register for entries made in respect of each class of drugs, and each of the drugs specified in paragraphs 1 and 3 of Schedule 1 and 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 paragraph (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 drugs or strengths of drugs comprised within the class of drugs to which that register or separate part relates.

(3) The foregoing provisions of this section shall not have effect in relation to:

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(a) in the case of a drug supplied to him for the purpose of destruction in pursuance of section 5(2) or (3), a practitioner or pharmacist;

(b) a person licensed under section 4 to supply any drug, where the licence so directs; or

(c) 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

18. Any person required to keep a register under section 17 shall comply with the following requirements, that is to say:

(a) the class of 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 section 17 in such a register shall be made on the day on which the 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 next 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 this Ordinance;

(f) 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 drugs in respect of which he is required to keep a separate register, so, however, that a separate register may, with the approval of the Board, be kept in respect of each department of the business carried on by him;

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

**Record-keeping requirements in respect of drugs in Schedule 2 in particular cases**

19. (1) Where a drug specified in Schedule 2 is supplied in accordance with section 7(5)(i) to any person on a ship, an entry in the official log book required to be kept under the Merchant Shipping Laws or, in the case of a ship which is not required to carry such an official logbook, a report signed by the master of the ship, shall, notwithstanding anything in this Ordinance, 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 Medical Officer of Health.

(2) A midwife authorised by section 10(1) to have in her possession any drug specified in Schedule 2 shall:

(a) on each occasion on which she obtains a supply of such a drug, enter in a book kept by her and used solely for the purposes of this subsection 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 such a drug to a patient, enter in that book as soon as practicable the name and address of the patient, the amount administered and the form in which it was administered.

Record-keeping requirements in respect of drugs in Schedules 3 and 4

20. (1) Every person who is authorised under section 4 or 8(1)(c) to produce any drug specified in Schedule 3 or 4 shall make a record of each quantity of such a drug produced by him.

(2) Every person who is authorised by or under any provision to the Law to import or export any drug specified in Schedule 3 shall make a record of each quantity of such a drug imported or exported by him.

(3) Every person who is authorised under section 8(4) to supply any drug specified in Schedule 4 shall make a record of each quantity of such a drug imported or exported by him.
Paragraph (2) shall not have effect in relation to a person licensed under the Law to import or export any drug where the licence so directs.

Preservation of registers, books and other documents

21. (1) All registers and books kept in pursuance of section 17 or 19(2) shall be preserved for a period of two years from the date on which the last entry therein is made.

(2) Every record made in pursuance of section 20 shall be preserved for a period of two years from the date on which the record was made.

(3) Every requisition, order or prescription (other than a medical prescription) on which a controlled drug is supplied in pursuance of this Ordinance 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 Schedules 3 and 5

22. (1) A producer of any drug specified in Schedule 3 or 5 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 person who is authorised under section 8(4)(a) to supply any drug specified in Schedule 3 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.

(3) A retail dealer in any drug specified in Schedule 3, a person in charge or acting person in charge of a hospital or nursing home and a person in charge or acting person in charge of any dispensary or pharmacy in charge of a hospital or nursing home 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.
charge of a laboratory 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.

(4) A retail dealer in any drug specified in Schedule 5 shall keep every invoice or other like record issued in respect of each quantity of such a drug obtained by him.

(5) Every invoice or other record which is required by this section to be kept in respect of a drug specified in Schedule 3 shall contain information sufficient to identify the date of the transaction and the person by whom or to whom the drug was supplied.

(6) Every document kept in pursuance of this section (other than a medical prescription) 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 the said period of two years shall be treated for the purposes of this paragraph as if it were the keeping of the original document.

**Furnishing of information with respect to controlled drugs**

23. (1) The persons specified in paragraph (2) shall on demand made by the Board or by any person authorised in writing by the Board in that behalf:

(a) furnish such particulars as may be requested in respect of the producing, obtaining or supplying by him of any controlled drug or in respect of any stock of such drugs in his possession;
(b) for the purpose of confirming any such particulars, produce any stock of such drugs in his possession;

(c) produce any register, book or document required to be kept under this Ordinance relating to any dealings in controlled drugs which is in his possession.

(2) The persons referred to in paragraph (1) are:

(a) any person authorised by or under this Ordinance to produce any controlled drug;

(b) any person authorised by or under any provision of the Law to import or export any controlled drug;

(c) a wholesale dealer;

(d) a retail dealer;

(e) a practitioner;

(f) the person in charge or acting person in charge of a hospital or nursing home;

(g) a person who is in charge of a laboratory;

(h) a person who is authorised under section 8(4)(a) to supply any controlled drug.
(3) Nothing in this section shall require the furnishing of personal records which a person has acquired or created in the course of his profession or occupation and which he holds in confidence; and in this paragraph "personal records" means documentary and other records concerning an individual (whether living or dead) who can be identified from them and relating to his physical or mental health.

**Destruction of controlled drugs**

**24.** (1) No person who is required by any provision of, or by any term or condition of a licence having effect under, this Ordinance to keep records with respect to a drug specified in Schedule 1, 2, 3 or 4 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 authorised (whether personally or as a member of a class) for the purposes of this paragraph by the Board (an "authorised person").

(2) An authorised person may, for the purposes of analysis, take a sample of a drug specified in Schedule 1, 2, 3 or 4 which is to be destroyed.

(3) Where a drug specified in Schedule 1, 2, 3 or 4 is destroyed in pursuance of paragraph (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, this Ordinance to keep a record in respect of the obtaining or supply of that drug, that record shall include particulars of the date of its destruction and the quantity destroyed and shall be signed by the authorised person in whose presence the drug is destroyed.

(4) Where the master or owner of a ship has in his possession a 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 an officer of police or to a person who may lawfully supply that drug to him.
(5) Nothing in paragraph (1) or (3) shall apply to any person who is required to keep records only by virtue of section 20(2) or (3) or 22(3).

(6) Nothing in paragraph (1) or (3) shall apply to the destruction of a drug which has been supplied to a practitioner or pharmacist for that purpose in pursuance of section 5(2) or (3).

Repeals, amendment, and transitional provisions

25. (1) The Ordinances listed in Schedule 7 are repealed.

(2) In section 4 of the Health Service (Benefit) Ordinance, 1990\(^m\) for "section 12 of the Misuse of Drugs (Bailiwick of Guernsey) Ordinance 1976" there is substituted "section 14 of the Misuse of Drugs (Bailiwick of Guernsey) Ordinance 1997".

(3) Notwithstanding subsection (1), any register, record, book, prescription or other document required to be preserved under section 18 or 19 of the Misuse of Drugs (Bailiwick of Guernsey) Ordinance 1976\(^n\) shall be preserved for the same period of time as if this Ordinance had not been made.

(4) In the case of a prescription issued before this Ordinance comes into force:

(a) paragraphs (a) and (b) of section 14(1) does not apply;

but

\(^{m}\) Recueil d'Ordonnances Tome XXV, p. 191.

\(^{n}\) Recueil d'Ordonnances, Tome XX, p.272.
(b) the prescription must comply with the requirements of the Misuse of Drugs (Bailiwick of Guernsey) Ordinance, 1976.

**Extent**

26. This Ordinance shall apply throughout the Bailiwick.

**Citation**

27. This Ordinance may be cited as the Misuse of Drugs (Bailiwick of Guernsey) Ordinance 1997.

**Commencement**

28. This Ordinance shall come into force on 1st June 1997.
CONTROLLED DRUGS SUBJECT TO THE REQUIREMENTS OF
SECTIONS 12,13,14,16,17,18,21,23 and 24.

1. The following substances and products, namely:

(a) Bufotenine
Cannabinol
Cannabinol derivatives, not being dronabinol or its stereoisomers
Cannabis and cannabis resin
Cathinone
Coca leaf
Concentrate of poppy-straw
Eticyclidine
Lysergamide
Lysergide and other N-alkyl derivatives of lysergamide
Mescaline
Psilocin
Raw opium
Rolicyclidine
Tenocyclidine
4-Bromo-2,5-dimethoxy-a-methylphenethylamine
N,N-Diethyltryptamine
N,N-Dimethyltryptamine
2,5-Dimethoxy-a,4-dimethylphenethylamine
N-Hydroxy-tenamphetamine
4-Methyl-aminorex;
(b) any compound (not being a compound for the time being specified in subparagraph (a) above) structurally derived from tryptamine or from a ring-hydroxy tryptamine by substitution at the nitrogen atom of the sidechain with one or more alkyl substituents but no other substituent;

(c) any compound (not being methoxyphenamine or a compound for the time being specified in subparagraph (a) above) structurally derived from phenethylamine, an N-alkylphenethylamine, a-
methylphenethylamine, an N-alkyl-a-methylphenethylamine, a-
ethylphenethylamine, or an N-alkyl-a-ethylphenethylamine by substitution in the ring to any extent with alkyl, alkoxy, alkenylenedioxy or halide substituents, whether or not further substituted in the ring by one or more other univalent substituents;

(d) any compound (not being a compound for the time being specified in Schedule 2) structurally derived from fentanyl by modification in any of the following ways, that is to say:

(i) by replacement of the phenyl portion of the phenethyl group by any heteromonocycle whether or not further substituted in the heterocycle;

(ii) by substitution in the phenethyl group with alkyl, alkenyl, alkoxy, hydroxy, halogeno, haloalkyl, amino or nitro groups;

(iii) by substitution in the piperidine ring with alkyl or alkenyl groups;
(iv) by substitution in the aniline ring with alkyl, alkoxy, alkenyinedioxy, halogeno or haloalkyl groups;

(v) by substitution at the 4-position of the piperidine ring with any alkoxy carbonyl or alkoxyalkyl or acyloxy group;

(vi) by replacement of the N-propionyl group by another acyl group;

(e) any compound (not being a compound for the time being specified in Schedule 2) structurally derived from pethidine by modification in any of the following ways, that is to say:

(i) by replacement of the 1-methyl group by an acyl, alkyl whether or not unsaturated, benzyl or phenethyl group, whether or not further substituted;

(ii) by substitution in the piperidine ring with alkyl or alkenyl groups or with a propano bridge, whether or not further substituted;

(iii) by substitution in the 4-phenyl ring with alkyl, alkoxy, aryloxy, halogeno or haloalkyl groups;

(iv) by replacement of the 4-ethoxycarbonyl by any other alkoxy carbonyl or alkoxylalkyl or acyloxy group;

(v) by formation of an N-oxide or of a quaternary base.

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 5.
SCHEDULE 2

Sections 5, 6, 7, 9, 17(1)(b), 19, 24(4)

CONTROLLED DRUGS SUBJECT TO THE REQUIREMENTS OF
SECTIONS 12, 13, 14, 16, 17, 18, 19, 21, 23 and 24.

1. The following substances and products, namely:

   Acetorphine
   Ecgonine, and any derivative
   of ecgonine which is
   or to cocaine
   Allyprodine
   convertible to ecgonine
   Alphacetylmethadol
   Ethylmethylthiambutene
   Alphameprodine
   Etonitazene
   Alphaprodine
   Etorphine
   Anilertidine
   Etoxeridine
   Benzethidine
   Fentanyl
   Benzylmorphine
   Furethidine
   (3-benzylmorphine)
   Hydrocodone
   Betacetylmethadol
   Hydromorphinol
   Betameprodine
   Hydromorphone
   Betamethadol
   Hydroxypethidine
   Betaprodine
   Isomethadone
   Bezitramide
   Ketobemidone
   Carfentanil
   Levomethorphan
   Clonitazene
   Levomoramide
   Cocaine
   Levophenacylmorphan
   Desomorphone
   Levorphanol
   Dextromoramide
   Lofentanil
Diamorphine  
Diampropamide  
Diethylthiambutene  
Difenoxin  
Dihydrocodeinone  
O-carboxymethyloxime  
Dihydromorphine  
(6-methylidihydromorphine)  
Dimenoxadole  
Dimepexanol  
Dimethylthiambutene  
Dioxaphetyl butyrate  
Diphenoxylate  
Dipipanone  
Dronabinol  
Drotebanol  
Nicomorphine  
Noracymethadon  
Normethadon  
Normorphine  
Norpipanone  
Oxycodone  
Oxymorphone  
Pethidine  
Phenadoxone  
Phenampronide  
Phencyclidine  
Phenomorphin  
Phenoperidine  
Medicinal opium  
Metazocine  
Methadone  
Methadyl acetate  
Methyldesoxphine  
Methyldihydromorphine  
Metopon  
Morpheridine  
Morphine  
Morphine methobromide,  
morphine N-oxide and other  
pentavalent nitrogen  
morphine derivatives  
Myrophine  
Properidine  
Racemethorphan  
Racemoramide  
Racemorphan  
Sufentanil  
Thebacon  
Thebaine  
Tilidate  
Trimeperidine  
4-Cyano-2-dimethylamino-4,  
4-diphenylbutane  
4-Cyano-1-methyl-4-  
phenylpiperidine  
1-Methyl-4-phenylpiperidine  
-4-carboxylic acid  

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Piminodine 2-Methyl-3-morpholino-1,
Piritramide 1-diphenylpropanecarboxylic
Proheptazine 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 5.

6. The following substances and products, namely:

   Acetyldihydrocodeine  Methaqualone
   Amphetamine  Methylamphetamine
   Codeine  Methylphenidate
   Dextropropoxyphene  Nicocodine
   Dihydrocodeine  Nicodicodine (6-
   Ethylmorphine nicotinoyldihydrocodeine)
   (3-ethylmorphine)
   Fenethylline  Norcodeine
   Glutethimide  Phenmetrazine
   Lefetamine  Pholcodine
   Propiram

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Mecloqualone                        Quinalbarbitone


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

9. Any preparation or other product containing a substance or product specified in any of paragraphs 6 to 8, not being a preparation specified in Schedule 5.
CONTROLLED DRUGS SUBJECT TO THE REQUIREMENTS OF
SECTIONS 12,13 (EXCEPT TEMAZEPAM),14,16,20,21,22,23 and 24.

1. The following substances, namely:

(a) Benzphetamine  Mephentermine
    Buprenorphine  Meprobamate
    Cathine  Methylphenobarbitone
    Chlorphentermine  Methyprylone
    Diethylpropion  Pentazocine
    Ethchlorvynol  Phendimetrazine
    Ethinamate  Phentermine
    Mazindol  Pipradrol
    Temazepam

(b) any 5,5 disubstituted barbituric acid not being quinalbarbitone.

2. Any stereoisomeric form of a substance specified in paragraph 1 not
   being phenylpropanolamine.

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

4. Any preparation or other product containing a substance specified in
   any of paragraphs 1 to 3, not being a preparation specified in Schedule 5.
CONTROLLED DRUGS EXCEPTED FROM THE PROHIBITION ON IMPORTATION, EXPORTATION AND, WHEN IN THE FORM OF A MEDICINAL PRODUCT, POSSESSION, AND SUBJECT TO THE REQUIREMENTS OF SECTIONS 20, 21, 23 AND 24.

1. The following substances and products, namely:

- Alprazolam
- Loprazolam
- Bromazepam
- Lorazepam
- Camazepam
- Lormetazepam
- Chlordiazepoxide
- Medazepam
- Clobazam
- Mefenorex
- Clonazepam
- Midazolam
- Clorazepic acid
- Nimetazepam
- Clotiazepam
- Nitrazepam
- Cloxazolam
- Nordazepam
- Delorazepam
- Oxazepam
- Diazepam
- Oxazolam
- Estazolam
- Pemoline
- Ethyl loflazepate
- Pinazepam
- Fencamfamin
- Prazepam
- Fenproporex
- Fludiazepam
- Pyrovalerone
- Flunitrazepam
- Flurazepam
- Tetrazepam
- Halazepam
- Triazolam
- Haloxazolam
- N-Ethylamphetamine

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Ketazolam

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

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

4. Any preparation or other product containing a substance of product specified in any of paragraphs 1 to 3, not being a preparation specified in Schedule 5.
CONTROLLED DRUGS EXCEPTED FROM PROHIBITION ON IMPORTATION, EXPORTATION AND POSSESSION AND SUBJECT TO THE REQUIREMENTS OF SECTIONS 22 and 23.

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 or 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 that 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 would constitute a risk to health.

3. Any preparation of medicinal 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 dextropropoxyphene, being a preparation designed for oral administration, containing not more than 135 milligrams of dextropropoxyphene (calculated as base) per dosage unit or with a total concentration of not more than 2.5 per cent (calculated as base) in undivided preparations.

5. Any preparation of difenoxin containing, per dosage unit, not more than 0.5 milligrams of difenoxin and a quantity of atropine sulphate equivalent to at least 5 per cent of the dose of difenoxin.

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

7. Any preparation of propiram containing, per dosage unit, not more than 100 milligrams of propiram calculated as base and compounded with at least the same amount (by weight) of methylcellulose.

8. Any powder of ipecacuanha and opium comprising-
   10 per cent 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.
9. Any mixture containing one or more of the preparations specified in paragraphs 1 to 8, being a mixture of which none of the other ingredients is a controlled drug.
**Section 17**

**SCHEDULE 6**

**FORM OF REGISTER**

**PART I**

*Entries to be made in case of obtaining*

<table>
<thead>
<tr>
<th>Date on which supply received</th>
<th>NAME</th>
<th>ADDRESS</th>
<th>Amount obtained</th>
<th>Form in which obtained</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Of person or firm from whom obtained</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**PART II**

*Entries to be made in case of supply*

<table>
<thead>
<tr>
<th>Date on which the transaction was effected</th>
<th>NAME</th>
<th>ADDRESS</th>
<th>Particulars as to licence or authority of person or firm supplied to be in possession</th>
<th>Amount supplied</th>
<th>Form in which supplied</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
SCHEDULE 7

Section 25(1)

Repeals

The Misuse of Drugs (Bailiwick of Guernsey) Ordinance, 1976

The Misuse of Drugs (Bailiwick of Guernsey) Law, 1974 (Modification) Ordinance, 1976⁰

The Misuse of Drugs (Amendment) (Bailiwick of Guernsey) Ordinance, 1983ᵖ

The Misuse of Drugs (Bailiwick of Guernsey) Law, 1974 (Modification) Ordinance, 1983⁹

The Misuse of Drugs (Bailiwick of Guernsey) Law, 1974 (Modification) Ordinance, 1988ʳ

The Misuse of Drugs (Bailiwick of Guernsey) Law, 1974 (Modification) Ordinance, 1989ˢ

The Misuse of Drugs (Amendment) (Bailiwick of Guernsey) Ordinance, 1989ᶜ

The Misuse of Drugs (Bailiwick of Guernsey) Law, 1974 (Modification) Ordinance, 1991ᵘ

The Misuse of Drugs (Amendment) (Bailiwick of Guernsey) Ordinance, 1991ᵛ

⁰ Recueil d'Ordonnances Tome XX, p.271.
ᵖ Recueil d'Ordonnances Tome XXII, p.480
⁹ Recueil d'Ordonnances Tome XXII, p.483
ʳ Recueil d'Ordonnances Tome XXIV, p.477
ˢ Recueil d'Ordonnances Tome XXV, p.38
ᶜ Recueil d'Ordonnances Tome XXV, p.43
ᵘ Recueil d'Ordonnances Tome XXV, p.325
ᵛ Recueil d'Ordonnances Tome XXV, p.326

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