DRUGS (PREVENTION OF MISUSE) ACT

CHAPTER 40:07

Act
20 of 1988
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15 of 1989
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CHAPTER 40:07

DRUGS (PREVENTION OF MISUSE) ACT

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FIRST SCHEDULE.
SECOND SCHEDULE.
THIRD SCHEDULE.
FOURTH SCHEDULE.
AN ACT to make provisions with respect to dangerous or otherwise harmful drugs and related matters, and for purposes connected therewith.

[29th December 1988]

1. This Act may be cited as the –

DRUGS (PREVENTION OF MISUSE) ACT.

2. (1) In this Act –

“Advisory Council” means the Advisory Council on the Misuse of Drugs established under this Act;

“analyst” means –

(a) the holder of any office specified in the First Schedule; and

(b) any other person appointed as such by the Minister by Notice in the Gazette;

“cannabis” (except in the expression “cannabis resin”) means any plant of the genus Cannabis or any part of any such plant, by whatever name designated;

“cannabis resin” means the separated resin, whether crude or purified, obtained from any plant or any part of a plant of the genus Cannabis;

“chemist and druggist” means a person registered as a chemist and druggist under the Medical Act;

“contravention” includes failure to comply, and “contravene” has a corresponding meaning;

“controlled drug” has the meaning assigned by section 4;

“corresponding law” has the meaning assigned by section 35;

“dentist” means a person registered as a dentist under the Medical Act;

“doctor” means a person registered as a medical practitioner under the Medical Act;
“druggist” means a person registered as a chemist and druggist under the Medical Act;

“drug trafficking” means doing or being concerned in any of the following, whether in Dominica or elsewhere:

(a) producing or supplying a controlled drug where the production or supply contravenes section 6(1) or a corresponding law;

(b) transporting or storing a controlled drug where possession of the drug contravenes section 7(1) or a corresponding law;

(c) importing or exporting a controlled drug where the importation or exportation is prohibited by section 5(1) or a corresponding law;

(d) entering into or being otherwise concerned in an arrangement whereby –

(i) the retention or control by or on behalf of another of the proceeds of drug trafficking by him is facilitated; or

(ii) the proceeds of drug trafficking by another are used to secure that funds are placed at his disposal or are used for his benefit to acquire property by way of investment;

“drug trafficking offence” means any of the following:

(a) an offence under section 5(3), 6(1) or (2) or (3) or 7(3);

(b) an offence under section 17;

(c) an offence under section 30 or section 38 of the Customs (Control and Management) Act in connection with a prohibition or restriction on importation or exportation having effect by virtue of section 5;

(d) conspiracy to commit any of the offences in paragraphs (a) to (c) above;

(e) an offence of attempting to commit any of those offences;

(f) an offence of inciting to commit any of those offences; and

(g) aiding, abetting, counselling or procuring the commission of those offences;
“Minister” means the Minister responsible for Health;

“practitioner” (except in the expression “veterinary practitioner”) means a doctor, a dentist or a veterinary practitioner;

“prepared opium” means opium prepared for smoking and includes dross and any other residues remaining after opium has been smoked;

“produce” where the reference is to producing a controlled drug, means producing it by manufacture, cultivation or any other method and “production” has a corresponding meaning;

“supplying” includes distributing;

“veterinary practitioner” means any person in Dominica holding a qualification entitling him to practise as a veterinary surgeon in any part of the Commonwealth or any person who within Dominica is engaged in the practice and profession of veterinary surgery.

(2) References in this Act to misusing a drug are references to misusing it by taking it; and the reference in the foregoing provision to the taking of a drug is a reference to the taking of it by a human being by way of any form of self administration, whether or not involving assistance by another.

(3) For the purpose of this Act the things which a person has in his possession shall be taken to include any thing subject to his control which is in the custody of another.

3. (1) There shall be constituted in accordance with the Second Schedule an Advisory Council on the Misuse of Drugs (in this Act referred to as “the Advisory Council”); and the supplementary provisions contained in that Schedule shall have effect in relation to the Advisory Council.

(2) It shall be the duty of the Advisory Council to keep under review the situation in Dominica with respect to drugs which are being or appear to them likely to be misused and of which the misuse is having or appears to them capable of having harmful effects sufficient to constitute a social problem, and to give to the Minister, where either the Advisory Council consider it expedient to do so or they are consulted by the Minister, advice on measures (whether or not involving alteration of the law) which in the opinion of the Advisory Council ought to be taken for preventing the misuse of such drugs or dealing with social problems arising out of the misuse of such drugs.
problems connected with their misuse, and in particular on measures which in the opinion of the Advisory Council, ought to be taken –

(a) for restricting the availability of such drugs or supervising the arrangements for their supply;

(b) for enabling persons affected by the misuse of such drugs to obtain proper advice, and for securing the provision of proper facilities and services for the treatment, rehabilitation and after-care of such persons;

(c) for promoting co-operation between the various professional and community services which in the opinion of the Advisory Council have a part to play in dealing with social problems connected with the misuse of such drugs;

(d) for educating the public (and in particular the young) in the dangers of misusing such drugs, and for giving publicity to these dangers; and

(e) for promoting research into, or otherwise obtaining information about, any matter which in the opinion of the Advisory Council is of relevance for the purpose of preventing the misuse of such drugs or dealing with any social problem connected with their misuse.

(3) It shall also be the duty of the Advisory Council to consider any matter relating to drug dependence or the misuse of drugs which may be referred to them by the Minister and to advise the Minister thereon.

4. (1) In this Act –

(a) the expression "controlled drugs" means any substance or product for the time being specified in Parts I, II or III of the Third Schedule;

(b) the expressions "Class A drug", "Class B drug" and "Class C drug" mean any of the substances and products for the time being specified respectively in Part I, Part II and Part III of the Third Schedule;

(2) Part IV of the Third Schedule shall have effect with respect to the meanings of expressions used in that Schedule.
(3) The Minister may after consultation with or on the recommendation of the Advisory Council by Order make such amendments to the Third Schedule as may be requisite for the purpose of adding any substance or product to, or removing any substance or product from, any of Parts I to III of that Schedule.

(4) An Order under this section may amend Part IV of the Third Schedule, and may do so whether or not it amends any other Part of that Schedule.

5. (1) Subject to subsection (2) –

(a) the importation of a controlled drug; and

(b) the exportation of a controlled drug;

are hereby prohibited.

(2) Subsection (1) does not apply –

(a) to the importation or exportation of a drug which is for the time being excepted from paragraph (a) or, as the case may be, paragraph (b) of subsection (1) by Regulations under section 9; or

(b) to the importation or exportation of a controlled drug under and in accordance with the terms of a licence issued by the Minister and in compliance with any conditions attached thereto.

(3) Any person who imports or exports controlled drugs contrary to subsection (1) is guilty of an offence and liable to the penalties laid down in section 27 and the Fourth Schedule.

6. (1) Subject to any Regulations under section 9 for the time being in force, it shall not be lawful for a person –

(a) to produce a controlled drug; or

(b) to supply or offer to supply a controlled drug to another.

(2) Subject to section 30, it is an offence for a person –

(a) to produce a controlled drug in contravention of subsection (1); or

(b) to be concerned in the production of such a drug by another in contravention of that subsection.
(3) Subject to section 30, it is an offence for a person –

(a) to supply or offer to supply a controlled drug to another in contravention of subsection (1);

(b) to be concerned in the supply of such a drug to another in contravention of subsection (1); or

(c) to be concerned in the making to another in contravention of subsection (1), of an offer to supply such a drug.

7. (1) Subject to any Regulations under section 9 for the time being in force, it shall not be lawful for a person to have a controlled drug in his possession.

(2) Subject to subsection (5) and to section 30, it is an offence for a person to have a controlled drug in his possession in contravention of subsection (1).

(3) Subject to section 30, it is an offence for a person to have a controlled drug in his possession, whether lawfully or not, with intent to supply it to another in contravention of section 6(1).

(4) Subject to subsection (1), a person found in possession of the following controlled drugs in quantities of more than –

(a) two grammes of diacetylmorphine (heroin);

(b) one grammme of cocaine;

(c) fifty-five grammes of opium;

(d) three grammes of morphine; or

(e) fifteen grammes of cannabis or cannabis resin,

shall be deemed to be in possession of such controlled drug for the purpose of supplying it to another or for drug trafficking in contravention of section 6(1) unless the contrary is proved, the burden of proof being on the accused.

(5) In any proceedings for an offence under subsection (2) or (3) in which it is proved that the accused had a controlled drug in his possession, it shall be a defence for him to prove –

(a) that, knowing or suspecting it to be a controlled drug, he took possession of it for the purpose of preventing another from committing or continuing to commit an
offence in connection with that drug and that as soon as possible after taking possession of it he took all such steps as were reasonably open to him to destroy the drug or to deliver it into the custody of a person fully entitled to take custody of it; or

(b) that, knowing or suspecting it to be a controlled drug, he took possession of it for the purpose of delivering it into the custody of a person lawfully entitled to take custody of it and that as soon as possible after taking possession of it he took all such steps as were reasonably open to him to deliver it into the custody of such a person.

(6) Subsection (5) shall apply in the case of proceedings for an offence under section 20 consisting of an attempt to commit an offence under subsection (2) or (3) as it applies in the case of proceedings for an offence under subsection (2) or (3), subject to the following modifications, that is to say –

(a) for the references to the accused having in his possession and to his taking possession of a controlled drug there shall be substituted respectively references to his attempting to get, and to his attempting to take, possession of such a drug; and

(b) in paragraphs (a) and (b) of subsection (5) the words from “and that as soon as possible” onwards shall be omitted.

(7) Nothing in subsection (5) or (6) shall prejudice any defence which is open to a person charged with an offence under this section to raise apart from those subsections.

8. (1) Subject to any Regulations under section 9 for the time being in force, it shall not be lawful for a person to cultivate any plant of the genus Cannabis.

(2) Subject to section 30, it is an offence to cultivate any such plant in contravention of subsection (1).

9. (1) The Minister may by Regulations –

(a) except from section 5(1)(a) or (b), 6(1)(a) or (b) or 7(1) such controlled drugs as may be specified in the Regulations; and
(b) make such other provision as he thinks fit for the purpose of making it lawful for persons to do things which under sections 6(1), 7(1) and 8(1), would otherwise be unlawful for them to do.

(2) Without prejudice to the generality of paragraph (b) of subsection (1), Regulations under that subsection authorising the doing of any such thing as is mentioned in that paragraph may in particular provide for the doing of that thing to be lawful—

(a) if it is done under and in accordance with the terms of a licence or other authority issued by the Minister and in compliance with any conditions attached thereto; or

(b) if it is done in compliance with such conditions as may be prescribed.

(3) Subject to subsection (4), the Minister shall so exercise his powers to make Regulations under subsection (1) as to secure—

(a) that it is not unlawful under section 6(1) for a doctor, dentist or veterinary practitioner acting in his capacity as such, to prescribe, administer, manufacture, compound or supply a controlled drug, or for a druggist or a person lawfully conducting a retail pharmacy business, acting in either case in his capacity as such, to manufacture, compound or supply a controlled drug; and

(b) that it is not unlawful under section 7(1) for a doctor, dentist, veterinary practitioner, druggist or person lawfully conducting a retail pharmacy business to have a controlled drug in his possession for the purpose of acting in his capacity as such.

(4) If in the case of a controlled drug the Minister is of the opinion that it is in the public interest—

(a) for production, supply and possession of that drug to be either wholly unlawful or unlawful except for purposes of research or other special purposes; or

(b) for it to be unlawful for practitioners, druggists and persons lawfully conducting retail pharmacy business to do in relation to that drug any of the things mentioned in subsection (3) except under a licence or other authority issued by the Minister,
He may by Order designate that drug as a drug to which this subsection applies; and while there is in force an Order under this subsection designating a controlled drug as one to which this subsection applies, subsection (3) shall not apply as regards that drug.

(5) An Order made under subsection (4) shall be subject to negative resolution of the House of Assembly.

(6) The Minister shall not make any Order under subsection (4) except after consultation with the Advisory Council.

(7) References in this section to a person "doing" things include references to his having things in his possession.

10. A person commits an offence if, being the occupier or concerned in the management of any premises, he knowingly permits or suffers any of the following activities to take place on those premises, that is to say —

(a) producing or attempting to produce a controlled drug in contravention of section 6(1);

(b) supplying or attempting to supply a controlled drug to another in contravention of section 6(1), or offering to supply a controlled drug to another in contravention of section 6(1);

(c) preparing cannabis, cannabis resin or opium for smoking;

(d) smoking cannabis, cannabis resin or prepared opium.

11. Subject to section 30, it is an offence for a person —

(a) to smoke or otherwise use prepared opium; or

(b) to frequent a place used for the purpose of opium smoking; or

(c) to have in his possession —

(i) any pipes or other utensils made or adapted for use in connection with the smoking of opium, being pipes or utensils which have been used by him or with his knowledge and permission in that connection or which he intends to use or permit others to use in that connection; or
Power to make 
regulations for 
preventing 
misuse of 
controlled drugs.

(ii) any utensils which have been used by him or with 
his knowledge and permission in connection with 
the preparation of opium for smoking.

12. (1) Subject to this Act, the Minister may by Regulations make 
such provisions as appear to him necessary or expedient for preventing 
the misuse of controlled drugs.

(2) Without prejudice to the generality of subsection (1), Regulations under this section may in particular make provisions –

(a) for requiring precautions to be taken for the safe custody 
of controlled drugs;

(b) for imposing requirements as to the documentation of 
transactions involving controlled drugs, and for requiring 
copies of documents relating to such transactions to 
be furnished to the prescribed authority;

(c) for requiring the keeping of records and the furnishing 
of information with respect to controlled drugs in such 
circumstances and in such manner as may be pre-
scribed;

(d) for the inspection of any precautions taken or records 
kept in pursuance of Regulations under this section;

(e) as to the packing and labelling of controlled drugs;

(f) for regulating the transport of controlled drugs and the 
methods used for destroying or otherwise disposing of 
such drugs when no longer required;

(g) for regulating the issue of prescriptions containing 
controlled drugs and the supply of controlled drugs on 
prescriptions, and for requiring persons issuing or dis-
spensing prescriptions containing such drugs to furnish 
to the prescribed authority such information relating to 
those prescriptions as may be prescribed;

(h) for requiring any doctor who attends to a person he 
considers, or has reasonable grounds to suspect, is 
addicted (within the meaning of the Regulations) to 
controlled drugs of any description to furnish to the 
prescribed authority such particulars with respect to 
that person as may be prescribed;
(i) for prohibiting any doctor from administering, supplying and authorising the administration and supply to persons so addicted, and from prescribing for such persons, such controlled drugs as may be prescribed, except under and in accordance with the terms of a licence issued by the Minister in pursuance of the Regulations.

13. (1) Without prejudice to any requirement imposed by Regulations made in pursuance of section 12(2)(a), the Minister may by notice in writing served on the occupier of any premises on which controlled drugs are or are proposed to be kept give directions as to the taking of precautions or further precautions for the safe custody of any controlled drugs of a description specified in the notice which are or are proposed to be kept on those premises.

(2) It is an offence to contravene any directions given under subsection (1).

14. (1) Where a person who is a practitioner or druggist has, after the coming into operation of this subsection, been convicted of an offence under this Act, the Minister may give a direction in writing under subsection (2) in respect of that person.

(2) A direction under this subsection in respect of a person shall—

(a) if that person is a practitioner, be a direction prohibiting him from having in his possession, prescribing, administering, manufacturing, compounding and supplying and from authorising the administration and supply of such controlled drugs as may be specified in the direction;

(b) if that person is a druggist, be a direction prohibiting him from having in his possession, manufacturing, compounding and supplying and from supervising and controlling the manufacture, compounding and supply of such controlled drugs as may be specified in the direction.
(3) The Minister may at any time give a direction cancelling or suspending any direction given by him under subsection (2), or cancelling any direction of his under this subsection by which a direction so given is suspended.

(4) The Minister shall cause a copy of any direction given by him under this section to be served on the person to whom it applies, and shall cause notice of any such direction to be published in the Gazette.

(5) A direction under this section shall take effect when a copy of it is served on the person to whom it applies.

(6) It is an offence to contravene a direction given under subsection (2).

15. (1) In the event of a contravention by a doctor of Regulations made in pursuance of paragraph (h) or (i) of section 12(2) or of the terms of a licence issued under Regulations made in pursuance of the said paragraph (i), the Minister may give a direction in respect of the doctor concerned prohibiting him from prescribing, administering and supplying and from authorising the administration and supply of such controlled drugs as may be specified in the direction.

(2) If the Minister is of the opinion that a practitioner is or has after the coming into operation of this subsection been prescribing, administering or supplying or authorising the administration or supply of any controlled drugs in an irresponsible manner, the Minister may, with the approval of the Advisory Council, give a direction in respect of the practitioner concerned prohibiting him from prescribing, administering and supplying such controlled drugs as may be specified in the direction.

(3) A contravention such as is mentioned in subsection (1) does not as such constitute an offence, but it is an offence to contravene a direction given under subsection (1) or (2).

16. (1) A person who commits the offence of trafficking in a substance other than a controlled drug which he represents or holds out to be a controlled drug is liable –

(a) upon summary conviction to a fine of five thousand dollars and to imprisonment for two years; or

(b) upon conviction on indictment to a fine of twenty-five thousand dollars and to imprisonment for five years.
(2) A person who commits a drug trafficking offence or the offence of being in possession of a controlled drug for the purpose of drug trafficking in any school, prison or military premises is liable –

(a) on summary conviction –

(i) to a fine of one hundred and fifty thousand dollars or where there is evidence of the street value of the controlled drug, three times the street value of the controlled drug whichever is greater; and

(ii) to imprisonment for a term which may extend to fifteen years but which shall not be less than seven years; or

(b) upon conviction on indictment to imprisonment for life.

(3) Subject to any Regulations under section 9 for the time being in force, a person found in possession of a controlled drug in any school premises is deemed to have the controlled drug for the purpose of drug trafficking unless the contrary is proved, the burden of proof being on the accused.

(4) In subsections (2) and (3) “school premises” includes buildings, playing fields or other premises established or maintained by a school for the benefit of its pupils whether or not such buildings, playing fields or other premises are within the curtilage of the school.

(5) A person who commits a drug trafficking offence or the offence of being in possession of a controlled drug for the purpose of drug trafficking and who in the course of committing the offence uses violence, corrupts or attempts to corrupt an official of the State or uses a child as carrier is liable –

(a) on summary conviction –

(i) to a fine of one hundred and fifty thousand dollars or where there is evidence of the street value of the controlled drug, three times the street value of the controlled drug whichever is greater; and

(ii) to imprisonment for a term which may extend to fifteen years but which shall not be less than seven years; and

(b) upon conviction on indictment to imprisonment for life.
17. (1) If a person enters into or is otherwise concerned in an arrangement whereby —

(a) the retention or control by or on behalf of another (call him “A”) of the proceeds of drug trafficking by A is facilitated (whether by concealment, removal from the jurisdiction, transfer to nominees or otherwise), or

(b) the proceeds of drug trafficking by A —

(i) are used to secure that funds are placed at A’s disposal; or

(ii) are used for A’s benefit to acquire property by way of investment,

knowing or suspecting or having reasonable grounds to suspect that A is a person who carries on or has carried on drug trafficking, he is guilty of an offence.

(2) In this section, references to the proceeds of drug trafficking by any person include a reference to any property which directly or indirectly represents in his hands the proceeds of drug trafficking by him.

(3) In proceedings against a person for an offence under this section, it is a defence to prove —

(a) that he did not know or suspect that the arrangement related to the proceeds of drug trafficking by A; or

(b) that he did not know or suspect that by the arrangement the retention or control by or on behalf of A of those proceeds was facilitated or, as the case may be, that by the arrangement those proceeds were used as mentioned in subsection (1).

(4) A person convicted of an offence under this section is liable—

(a) on summary conviction to a fine of five thousand dollars and to imprisonment for two years; or

(b) on conviction on indictment to a fine of fifty thousand dollars and to imprisonment for fourteen years.
18. (1) If it appears to the Minister that there exists in any area in Dominica a social problem caused by the extensive misuse of dangerous or otherwise harmful drugs in that area, he may by notice in writing served on any doctor or druggist practising in or in the vicinity of that area, or on any person lawfully conducting a retail pharmacy business at any premises situated in or in the vicinity of that area, require him to furnish to the Minister, with respect to any such drug specified in the notice and as regards any period so specified, such particulars as may be so specified relating to the quantities in which and the number and frequency of the occasions on which those drugs—

(a) in the case of a doctor, were prescribed, administered or supplied by him;

(b) in the case of a druggist, were supplied by him; or

(c) in the case of a person conducting a retail pharmacy business, were supplied in the course of that business at any premises so situated which may be specified in the notice.

(2) A notice under subsection (1) may require any such particulars to be furnished in such manner and within such time as may be specified in the notice and, if served on a druggist or person conducting a retail pharmacy business, may require him to furnish the names and addresses of doctors on whose prescriptions any dangerous or otherwise harmful drugs to which the notice relates were supplied, but shall not require any person to furnish any particulars relating to the identity of any person for or to whom any such drug has been prescribed, administered or supplied.

(3) A person commits an offence if without reasonable excuse (proof of which shall lie on him) he fails to comply with any requirements to which he is subject by virtue of subsection (1).

(4) A person commits an offence if, in purported compliance with a requirement imposed under this section, he gives any information which he knows to be false in a material particular or recklessly gives any information which is so false.

19. (1) It is an offence for a person to contravene any Regulations made under this Act other than Regulations made in pursuance of section 12(2)(h) or (i).
(2) It is an offence for a person to contravene a condition or other term of a licence issued under section 5 or of a licence or other authority issued under Regulations made under this Act, not being a licence issued under Regulations made in pursuance of section 12(2)(i).

(3) A person commits an offence if, in purported compliance with any obligation to give information to which he is subject under or by virtue of Regulations made under this Act, he gives any information which he knows to be false in a material particular or recklessly gives any information which is so false.

(4) A person commits an offence if, for the purpose of obtaining, whether for himself or another, the issue or renewal of a licence or other authority under this Act or under any Regulations made under this Act, he –

(a) makes any statement or gives any information which he knows to be false in a material particular or recklessly gives any information which is so false; or

(b) produces or otherwise makes use of any book, record or other document which to his knowledge contains any statement or information which he knows to be false in a material particular.

20. It is an offence for a person to attempt to commit an offence under this Act or to incite or attempt to incite another to commit such an offence.

21. A person commits an offence if in Dominica he assists in or induces the commission in any place outside Dominica of an offence punishable under the provisions of a corresponding law in force in that place.

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

23. The Minister may by Regulations make provision –
(a) for excluding in such cases as may be prescribed the application of any provision of this Act which create an offence;

(b) for the application of any provisions of this Act or Regulations or Orders thereunder to servants or agents of the State, subject to such exceptions, adaptations and modifications as may be prescribed.

24. (1) A member of the Police Service or other person authorised in that behalf by a general or special order of the Minister shall, for the purposes of the execution of this Act, have power to enter the premises of a person carrying on business as a producer or supplier of any controlled drug and to demand the production of, and to inspect, any books or documents relating to dealings in any such drugs and to inspect any stocks of any such drugs.

(2) If a member of the Police Service has reasonable grounds to suspect that any person is in possession of a controlled drug in contravention of this Act or of any Regulations made thereunder the member of the Police Service may, subject to subsections (3), (6) and (7) –

(a) search that person, and detain him for the purpose of searching him;

(b) search any ship, vessel, boat, aircraft, vehicle or other means of conveyance of any description in which the member of the Police Service suspects that the drug may be found, and for that purpose require the person in control of the ship, vessel, aircraft, vehicle or other means of conveyance of any description to stop it;

(c) seize and detain for the purposes of proceedings under this Act, anything found in the course of the search which appears to the member of the Police Service to be evidence of an offence under this Act.

(3) Nothing in subsection (2) shall derogate from any power of search or any power to seize or detain property which is otherwise exercisable by a member of the Police Service.

(4) If a Magistrate or a Justice of the Peace is satisfied by information on oath that there is reasonable ground for suspecting –

(a) that any controlled drugs are, in contravention of this
Act or of any Regulations made thereunder, in the possession of a person on any premises or in any place; or

(b) that a document directly or indirectly relating to, or connected with, a transaction or dealing which was, or an intended transaction or dealing which would if carried out be, an offence under this Act, or in the case of a transaction or dealing carried out or intended to be carried out in a place outside Dominica an offence against a corresponding law in force in that place, is in the possession of a person on any premises or in any place,

he may issue a warrant authorising any member of the Police Service at any time or times within one month from the date of the issue of warrant to enter, if need be by force, the premises or place named in the warrant, and to search the premises or place and any persons found therein and, if there is reasonable ground for suspecting that an offence under this Act has been committed in relation to any controlled drugs found on the premises or place or in the possession of any such persons, or that a document so found is such a document as is mentioned in paragraph (b), to seize and detain those drugs or that document, as the case may be.

(5) A person commits an offence if he –

(a) intentionally obstructs a person in the exercise of his powers under this section;

(b) being the person in control of the ship, vessel, boat, aircraft, vehicle or other means of conveyance of any description fails to stop it when required to do so by a member of the Police Service under subsection (2)(b);

(c) being a person being conveyed in a ship, vessel, boat, aircraft, vehicle or other means of conveyance of any description prevents or intimidates the person in control of or any other person operating the ship, vessel, aircraft, vehicle or other means of conveyance of any description from stopping when required to do so by a member of the Police Service under subsection (2)(b);

(d) without the permission of the member of the Police Service concerned, leaves a ship, vessel, boat, aircraft, vehicle or other conveyance of any description which
has been stopped by a member of the Police Service under subsection (2)(b);

(e) conceals from a person acting in the exercise of his powers under subsection (1) any such books, documents, stocks or drugs as are mentioned in that subsection; or

(f) without reasonable excuse (proof of which shall lie on him) fails to produce any such books or documents as are so mentioned where their production is demanded by a person in the exercise of his powers under that subsection.

(6) No person may be searched by any person of the opposite sex unless the consent of the person to be searched has first been obtained, or unless the search is made in the presence of some other person, not being a member of the Police Service, of the same sex as the person to be searched.

(7) No article of a person’s clothing may be removed from his person during a search at any place other than within a police station.

(8) Where the Commissioner of Police is satisfied by the report of an analyst that any substance seized or obtained by the police is a controlled drug, he shall destroy the substance retaining only a small sample which in any prosecution involving that substance shall be sufficient evidence of the whole.

25. (1) A member of the Police Service may arrest without warrant a person who has committed, or whom the member of the Police Service, with reasonable cause, suspects to have committed, an offence under this Act.

(2) Where any controlled drug is found on any premises searched under section 24(1), or in any ship, vessel, boat, aircraft, vehicle or other means of conveyance of any description stopped under section 24(2), the member of the Police Service who has made the search or stopped the ship, vessel, boat, aircraft, vehicle or other means of conveyance of any description, as the case may be, may arrest without warrant any person in such premises or in such ship, vessel, boat, aircraft, vehicle or other means of conveyance of any description whom he has reason to believe to be guilty of an offence under this Act.

(3) This section shall not prejudice any power of arrest conferred by law apart from this section.
26. (1) Where on the summary trial of an offence committed under this Act and triable either summarily or on indictment a person who is not less than eighteen years of age is convicted of the offence, then, if on obtaining information that his character and antecedents are such that in the opinion of the Magistrate greater punishment should be inflicted for the offence than the Magistrate has power to inflict, the Magistrate may commit that person in custody to the High Court for sentence and shall, as soon as practicable, transmit to the Registrar of the High Court the record of the proceedings.

(2) On receipt of the record, the Registrar of the High Court shall issue an order to the gaoler to bring the convicted person before a Judge of the High Court at a time to be fixed by the Judge of the High Court.

(3) The Judge of the High Court shall enquire into the circumstances of the case and shall have power to deal with the convicted person as if he had just been convicted of the offence on indictment before the High Court.

27. (1) Except where expressly otherwise provided the Fourth Schedule shall have effect, in accordance with subsection (2), with respect to the way in which a person guilty of an offence under this Act is punishable on conviction.

(2) In relation to an offence under this Act specified in the first column of the Fourth Schedule –

(a) the second column describes the general nature of the offence;

(b) the third column shows whether the offence is punishable on summary conviction or on indictment or in either way;

(c) the fourth, fifth, and sixth columns show respectively the punishments which may be imposed on a person convicted of the offence in the way specified in relation thereto in the third column (that is to say, summarily or on indictment) according to whether the controlled drug in relation to which the offence was committed was a Class A drug, a Class B drug or a Class C drug; and

(d) the seventh column shows the punishments which may be imposed on a person convicted of the offence in the way specified in relation thereto in the third column
and in the fourth, fifth, sixth and seventh columns a reference to a period gives the maximum term of imprisonment and a reference to a sum of money the maximum fine.

(3) An offence under section 20 shall be punished on summary conviction, on indictment or in either way according to whether, under the Fourth Schedule, the substantive offence is punishable on summary conviction, on indictment or in either way; and the punishments which may be imposed on a person convicted of an offence under that section are the same as those which, under that Schedule, may be imposed on a person convicted of the substantive offence.

(4) In subsection (3), "the substantive offence" means the offence under this Act to which the attempt or, as the case may be, the incitement or attempted incitement mentioned in section 20 was directed.

(5) Notwithstanding section 62 of the Magistrates Code of Procedure Act a Magistrate may try an information or complaint for an offence under this Act if the information or complaint was made at any time within twelve months from the commission of the offence.

28. (1) Where a person is convicted of an offence under this Act other than a drug trafficking offence, the Court may order forfeiture to the Government of Dominica of any opium pipe or other article or the controlled drug in respect of which the offence was committed and all receptacles of any kind whatsoever found containing the controlled drug and any ship, vessel, boat, aircraft, vehicle or other means of conveyance of any description proved to have contained the opium pipe or other article or controlled drug or anything shown to the satisfaction of the Court to relate to the offence.

(2) Without prejudice to subsection (1), where a person is convicted of a drug trafficking offence the Court shall in passing sentence order forfeiture to the Government of Dominica of –

(a) any article;

(b) any money; or
(c) any valuable consideration, relating to the offence.

(3) Forfeiture shall extend –

(a) to any property which there is reason to believe has been obtained from the proceeds of anything relating to the offence for which a person is convicted under this Act or to a conspiracy to commit any such offence; or

(b) to any thing into which any such property has been converted.

(4) Subject to subsection (5), forfeitures under this section shall be applied to treatment, rehabilitation, education and preventative measures related to drug abuse.

(5) Controlled drugs forfeited under this section shall be disposed of by a member of the Police Service or other person authorised in that behalf by a general or special order of the Minister.

29. (1) Subject to subsection (2), notwithstanding any other law, a certificate of an analyst purporting to be signed by him stating that he has analysed or examined a substance and stating the result of the analysis or examination is admissible in evidence in any prosecution under this Act and in the absence of evidence to the contrary is proof of the facts contained in the certificate without proof of the signature or the official character of the person appearing to have signed the certificate.

(2) In any prosecution under this Act either of the parties may require the attendance of an analyst to give evidence and in such case the costs of his attendance shall, unless the Judge or Magistrate orders otherwise, be payable by the party so requiring.

30. (1) This section applies to offences under section 6(2) and (3), section 7(2) and (3), and section 8(2) and section 11.

(2) Subject to subsection (3), in any proceedings for an offence to which this section applies it shall be a defence for the person charged to prove that he neither knew of nor suspected nor had reason to suspect the existence of some fact alleged by the prosecution which it is necessary for the prosecution to prove if he is to be convicted of the offence charged.

(3) Where in any proceedings for an offence to which this
section applies it is necessary if the accused is to be convicted of the offence charged for the prosecution to prove that some substance or product involved in the alleged offence was the controlled drug which the prosecution alleges it to have been, and it is proved that the substance or product in question was that controlled drug, the accused —

(a) shall not be acquitted of the offence charged by reason only of proving that he neither knew, nor suspected nor had reason to suspect that the substance or product in question was the particular controlled drug alleged; but

(b) shall be acquitted thereof —

(i) if he proves that he neither believed nor suspected nor had reason to suspect that the substance or product in question was a controlled drug; or

(ii) if he proves that he believed the substance or product in question to be a controlled drug, or a controlled drug of a description, such that if it had in fact been that controlled drug or a controlled drug of that description, he would not at the material time have been committing any offence to which this section applies.

(4) Nothing in this section shall prejudice the raising of any available defence which is open to a person charged with an offence to which this section applies.

31. (1) Any notice or other document required or authorised by this Act to be served on any person may be served on him either by delivering it to him or by leaving it at his proper address or by sending it by post.

(2) Any notice or other document so required or authorised to be served on a body corporate shall be duly served if it is served on the secretary or clerk of that body.

(3) For the purposes of this section the proper address of any person shall, in the case of the secretary or clerk of a body corporate, be that of the registered or principal office of that body, and in any other case shall be the last address of the person to be served which is known to the Minister.

32. A licence or other authority issued by the Minister for the purposes of this Act or of Regulations made under this Act, may be to any degree, general or specific, issued on such terms and subject to such
conditions (including in the case of a licence the payment of a prescribed fee) as the Minister thinks proper, and may be modified or revoked by him at any time.

33. (1) The Minister may from time to time by Regulations add to, rescind or alter the Drugs (Prevention of Misuse) Regulations and the Drugs (Notification of and Supply to Addicts) Regulations; and any Regulations made by the Minister under this Act –

(a) may make different provisions in relation to different controlled drugs, different classes of persons, different provisions of this Act or other different cases or circumstances; and

(b) may make the opinion, consent or approval of a prescribed authority or of any person authorised in a prescribed manner material for purposes of any provision of the Regulations; and

(c) may contain such supplementary, incidental and transitional provisions as appear expedient to the Minister.

(2) The Minister shall not make any Regulations under this Act except after consultation with the Advisory Council.

(3) Any Regulations made under this Act by the Minister shall be subject to negative resolution of Parliament.

34. The Minister may conduct or assist in conducting research into any matter relating to the misuse of dangerous or otherwise harmful drugs.

35. (1) In this Act the expression “corresponding law” means a law stated in a certificate purporting to be issued by or on behalf of the government of a country outside Dominica to be a law providing for the control and regulation in that country of the production, supply, use, export and import of drugs and other substances in accordance with the provisions of the Single Convention on Narcotic Drugs signed at New York on 30th March, 1961 or a law providing for the control and regulation in that country of the production, supply, use, export and import of dangerous or otherwise harmful drugs in pursuance of any treaty, convention or other agreement or arrangement to which the government of that country and the Government of Dominica are for the time being parties.

(2) A statement in any such certificate as aforesaid to the effect
that any facts constitute an offence against the law mentioned in the certificate shall be conclusive evidence of the matters stated.

FIRST SCHEDULE
COMMONWEALTH OF DOMINICA

Government analyst Commonwealth of Dominica.

SECOND SCHEDULE
CONSTITUTION, ETC., OF ADVISORY COUNCIL
ON THE MISUSE OF DRUGS

1. (1) The members of the Advisory Council, of whom there shall be not more than eleven nor less than seven, shall be appointed by the Minister after consultation with such organisations as he considers appropriate, and shall include-

(a) in relation to each of the activities specified in subparagraph (2) at least one person appearing to the Minister to have wide and recent experience of that activity; and

(b) persons appearing to the Minister to have wide and recent experience of social problems connected with the misuse of drugs.

(2) The activities referred to in subparagraph (1)(a) are –

(a) the practice of medicine (other than veterinary medicine);
(b) the practice of dentistry;
(c) the practice of veterinary medicine;
(d) the practice of pharmacy.

(3) The Minister shall appoint one of the members of the Advisory Council to be Chairman of the Advisory Council.

2. The Advisory Council may appoint committees, which may consist in part of persons who are not members of the Advisory Council to consider and report to the Advisory Council on any matter referred to them by the Advisory Council.

3. At meetings of the Advisory Council the quorum shall be four, and subject to that the Advisory Council may determine its own procedure.
THIRD SCHEDULE.
CONTROLLED DRUGS
Part I
CLASS A DRUGS

The following substances and products, namely –

Acetorphine
Allylprodine
Alphacetylmethadol
Alphameprodine
Alphamethadol
Alphadrodine
Anileridine
Benzethidine
Benzylmorphine
(3-benzylmorphine)
Betacetylmethadol
Betameprodine
Betamethadol
Bezitramide
Buffogenine
Cannabinol, except
where contained in
cannabis or cannabis
resin
Cannabinol derivatives
Clonitazene
Coca leaf
Cocaine
Desomorphine
Dextromoramide
Diamorphine
Diethylthiambutene
Diethylamiobutene
D-carboxymethylxime
Dihydromorphine
Dimenoxadol
Dimepheptanol
Dimethylthiambutene
Dioxaphetyl butyrate
Diphenoxylate
Dipipanone

Egonine, and any
derivative of egonine which is
convertible to egonine or to
cocaine
Ethylmethylthiambutene
Etonitazene
Etorphine
Extoeridine
Fentanyl
Furethidine
Hydrocodone
Hydromorphone
Hydroxpethidine
Isomethadone
Ketobemidone
Levomethorphan
Levophenacylmorphan
Levorphanol
Lysergamide
Lysergide and other
Morpheridine
Morphine methobromide,
morphine N-oxide
and other pentavalent
nitrogen morphine
derivatives
Mescaline
Metazocine
Methadone
Methyldesorphine
Methyldihydromorphine
(6methyldihyromorphine)
Metopon
Morphine
Myrophine
Noracymethadol  
Oxycodone  
Nicodomine (6-nicotinylidihydrodeine)  
Nicodimorphine (3, 6-dinicotoninylmorphine)  
Normorphine  
Oxymorphone  
Phenadoxone  
Phenazocine  
Phenoperidine  
Proheptazine  
Properidine (1-methyl 4-phenyl-1 piperidine  
4-carboxylic acid isopropyl ester)  
Pethidine  
Phenamapromide  
Pipinodine  
Pipinodine  

2. Any stereoisomeric form of a substance for the time being specified in paragraph 1 not being dextromethorphan or dextrophan.

3. Any ester of a substance for the time being specified in paragraph 1 or 2.

4. Any salt of a substance for the time being specified in any of paragraphs 1 to 3.

5. Any preparation or other product containing a substance or product for the time being specified in any of paragraphs 1 to 4.

6. Any preparation designed for administration by injection which includes a substance or product for the time being specified in any of paragraphs 1 to 3 or Part II.

1. The following substances and products, namely—

- Acetyldihydrocodeine
- Amphetamine
- Cannabis and cannabis resin
- Codeine
- Dexamphetamine
- Diphydrocodeine
- Ethylmorphine
- (3-ethylomorphine).

2. Any stereoisomeric form of a substance for the time being specified in paragraph 1.

3. Any salt of a substance for the time being specified in paragraph 1 or 2.

4. Any preparation or other product containing a substance or product for the time being specified in any of paragraphs 1 or 2, not being a preparation falling within paragraph 6 of Part I.

Part III

CLASS C DRUGS

1. The following substances, namely—

- Benzphetamine
- Chlorphentermine
- Fencamfamin
- Mephentermine
- Methaqualone
- Pemoline
- Phendimetrazine
- Phentermine
- Pipradrol
- Prolintane.

2. Any stereoisomeric form of a substance for the time being specified in paragraph 1.

3. Any salt of a substance for the time being specified in paragraph 1 or 2.

4. Any preparation or other product containing a substance for the time being specified in any of paragraphs 1 to 3.
Part IV

MEANING OF CERTAIN EXPRESSIONS
USED IN THIS SCHEDULE

For the purposes of this Schedule the following expressions (which are not among those defined in section 2 of this Act) have the meanings hereby assigned to them respectively, that is to say –

“cannabinol derivatives” means the following substances, except where contained in cannabis or cannabis resin, namely tetrahydro derivatives of cannabinol and 3-alkyl homologues of cannabinol or of its tetrahydro derivatives;

“coca leaf” means the leaf of any plant of the genus Erythroxylon from whose leaves cocaine can be extracted either directly or by chemical transformation;

“concentrate of poppy-straw” means the material produced when poppy-straw has entered into a process for the concentration of its alkaloids;

“medicinal opium” means raw opium which has undergone the process necessary to adapt it for medicinal use in accordance with the requirements of the British Pharmacopoeia, whether it is in the form of powder or is granulated or is in any other form, and whether it is or is not mixed with neutral substances;

“opium poppy” means the plant of the species Papaver somniferum L;

“poppy straw” means all parts, except the seeds, of the opium poppy after mowing;

“raw opium” includes powdered or granulated opium but does not include medicinal opium.
## FOURTH SCHEDULE

### PROSECUTION AND PUNISHMENT OF OFFENCES

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<td>3 years and $100,000</td>
<td>2 years and $75,000</td>
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<td>(b) On indictment</td>
<td>14 years and $200,000</td>
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<tr>
<td>Section 6(2)</td>
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<td>(a) Summary</td>
<td>3 years and $100,000</td>
<td>3 years and $100,000</td>
<td>2 years and $75,000</td>
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<tr>
<td>Section 6(3)</td>
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<td>(a) Summary</td>
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<td>3 years and $100,000</td>
<td>2 years and $75,000</td>
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<td>Section 7(2)</td>
<td>Having possession of a controlled drug</td>
<td>(a) Summary</td>
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<td>12 months and $10,000</td>
<td>6 months and $10,000</td>
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<td>Section 7(3)</td>
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<td>(a) Summary</td>
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<tr>
<td>Section 10</td>
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<tr>
<td>Section 13(2)</td>
<td>Contravention of directions relating to safe custody of controlled drugs</td>
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| Section 18(3)             | Failure to comply with notice requiring information relating to prescribing, supplying, etc., of drugs | (a) Summary | Class A Drug involved
|                           |                                                                                         |            | Class B Drug involved
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|                           |                                                                                         | $ 75,000   | General
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Drugs (Prevention of Misuse) Regulations

REGULATION

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FIRST SCHEDULE.
SECOND SCHEDULE.
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FOURTH SCHEDULE.
DRUGS (PREVENTION OF MISUSE) REGULATIONS

made under section 33

[29th December 1988]

Citation.

1. These Regulations may be cited as the –

   DRUGS (PREVENTION OF MISUSE) REGULATIONS.

Interpretation.

2. (1) In these Regulations –

   “authorised as a member of a group” means authorised by virtue of being a member of a class as respects which the Minister has granted an authority under and for the purposes of regulation 7(3), 8(3) or 9(3) which is in force, 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;

   “master” in relation to a ship, includes any person having or taking the charge or command of the ship;

   “officer of customs” means any person employed in the department of Customs;

   “prescription” means a prescription issued by a doctor 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 pharmacy” means a pharmacy under the management or control of a chemist or druggist registered under the Medical Act;

   “retail dealer” means a person lawfully conducting a retail pharmacy business;

   “wholesale dealer” means a person who carries on the business of selling drugs to persons who buy to sell again.
(2) In these Regulations any reference to any written law shall be construed as a reference to that written law, and as including a reference thereto as extended or applied by or under any other written law.

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

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

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

4. Where any person is authorised by a licence of the Minister 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(4) of the Act 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.

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

(a) a member of the police service 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) an officer of customs and excise 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.
6. (1) Any person may administer to another any drug specified in the First Schedule.

(2) A doctor or dentist may administer to a patient any drug specified in the First or Second Schedule.

(3) Any person other than a doctor or dentist may administer to a patient, in accordance with the directions of a doctor or dentist, any drug specified in the First or Second Schedule.

7. (1) Notwithstanding section 6(1)(a) of the Act –

(a) a practitioner or druggist, acting in his capacity as such, may manufacture or compound any drug specified in the First or Second Schedule.

(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 the First or Second Schedule.

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

(a) a practitioner;

(b) a druggist;

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

(d) the matron of a hospital or nursing home;

(e) in the case of a drug supplied to her by a person responsible for the dispensing and supply of medicines at the hospital or nursing home, the 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 university, hospital or to any other institution approved for the purpose by the Minister;
(g) an analyst, may when acting in his capacity as such, supply or offer to supply any drug specified in the First or Second Schedule to any person who may lawfully have that drug in his possession; but nothing in this subregulation authorises –

(h) the matron of a hospital or nursing home having a druggist responsible for the dispensing and supply of medicines, to supply or offer to supply any drug;

(i) a 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 doctor or druggist.

(3) Notwithstanding section 6(1)(b) of the Act, 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 the First or Second Schedule to any person who may lawfully have that drug in his possession.

(4) Notwithstanding section 6(1)(b) of the Act, a person whose name is for the time being entered in the register kept for the purposes of this subregulation by the Minister may, at the premises in respect of which his name is so entered and in compliance with any condition subject to which his name is so entered, supply or offer to supply any drug specified in the First Schedule to any person who may lawfully have that drug in his possession.

(5) Notwithstanding section 6(1)(b) of the Act the owner of a ship, or the master of a ship which does not carry a doctor on board as part of her complement, may supply or offer to supply any drug specified in the First or Second Schedule –

(a) to any member of the crew;

(b) to any person who may lawfully supply that drug; or

(c) to any member of the police service for the purpose of destruction.

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

(a) a practitioner or druggist, acting in his capacity as such, may manufacture or compound any drug specified in the Third Schedule;
(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 the Third Schedule;

(c) a person whose name is for the time being entered in the register kept for the purposes of this paragraph by the Minister may produce, at the premises in respect of which his name is so entered and in compliance with any conditions subject to which his name is so entered, any drug specified in the Third Schedule.

(2) Notwithstanding section 6(1)(b) of the Act, any of the following persons, that is to say —

(a) a practitioner;

(b) a druggist;

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

(d) the matron of a hospital or nursing home;

(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 for the time being in charge of a ward, theatre or other department in a hospital or nursing home;

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

(g) an analyst,

may, when acting in his capacity as such, supply or offer to supply any drug specified in the Third Schedule to any person who may lawfully have that drug in his possession; but nothing in this subregulation authorises —

(h) the matron of a hospital or nursing home, having a druggist responsible for the dispensing and supply of medicines, to supply or offer to supply any drug;

(i) a 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 doctor or dentist.
(3) Notwithstanding section 6(1)(b) of the Act, 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 the Third Schedule to any person who may lawfully have that drug in his possession.

(4) Notwithstanding section 6(1)(b) of the Act –

(a) a person whose name is for the time being entered in the register kept for the purposes of this paragraph by the Minister may, at the premises in respect of which his name is so entered and in compliance with any conditions subject to which his name is so entered, supply or offer to supply any drug specified in the Third Schedule to any person who may lawfully have that drug in his possession;

(b) a person whose name is for the time being entered in the register kept for the purposes of subregulation (1)(c) by the Minister may supply or offer to supply any drug which he may, by virtue of his name being so entered, lawfully produce to any person who may lawfully have that drug in his possession.

(5) Notwithstanding section 6(1)(b) of the Act, the owner of a ship, or the master of a ship which does not carry a doctor on board as part of her complement, may supply or offer to supply any drug specified in the Third Schedule –

(a) to any member of the crew; or

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

9. (1) Notwithstanding section 7(1) of the Act –

(a) a person specified in regulation 7(2) may have in his possession any drug specified in the Second Schedule.

(b) a person specified in regulation 8(2) may have in his possession any drug specified in the Third Schedule, for the purpose of acting in his capacity as such.

(2) Notwithstanding section 7(1) of the Act a person may have in his possession any drug specified in the Second or Third Schedule for administration for medical, dental or veterinary purposes in accordance with the Act.
with the directions of a practitioner; but this subregulation shall not have effect in the case of a person to whom the drug has been supplied by or on the prescription of a doctor if –

(a) that person was then being supplied with any controlled drug by or on the prescription of another doctor and failed to disclose that fact to the first-mentioned doctor 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 section 7(1) of the Act, 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 the Second or Third Schedule in his possession.

(4) Notwithstanding section 7(1) of the Act –

(a) a person whose name is for the time being entered in the register kept for the purposes of this paragraph by the Minister may, in compliance with any conditions subject to which his name is so entered, have in his possession any drug specified in the Third Schedule;

(b) a person whose name is for the time being entered in the register kept for the purposes of regulation 8(1)(c) by the Minister may have in his possession any drug which he may, by virtue of his name being so entered lawfully produce;

(c) a person whose name is for the time being entered in the register kept for the purposes of regulation 8(4)(a) by the Minister may have in his possession any drug which he may, by virtue of his name being so entered, lawfully supply or offer to supply.

(5) Notwithstanding section 7(1) of the Act –

(a) the owner of a ship, or the 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 the Second or Third Schedule so far as necessary for the equipment of the ship and compliance with the Merchant Shipping Act;
(b) the master of a foreign ship which is in a port in Dominica may have in his possession any drug specified in the Second or Third Schedule so far as necessary for the equipment of the ship.

10. (1) Notwithstanding sections 6(1)(b) and 7(1) of the Act, a registered midwife who has notified the Chief Medical Officer of her intention to practise, 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 aforesaid, administer pethidine; and

(c) surrender to the Chief Medical Officer any stocks of pethidine in her possession which are no longer required by her.

(2) Nothing in subregulation (1) authorises a midwife to have in her possession pethidine which has been obtained otherwise than on an order in writing specifying the name and occupation of the midwife obtaining the pethidine, the purpose for which it is required and the total quantity to be obtained signed by a doctor.

11. Where any person is authorised by a licence of the Minister issued under this regulation and for the time being in force to cultivate plants of the genus Cannabis, it shall not by virtue of section 8 of the Act 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.

12. Section 10 of the Act (which makes it an offence for the occupier of premises to permit certain activities there) shall not have effect in relation to the smoking of cannabis or cannabis resin for the purposes of research on any premises for the time being approved for the purposes by the Minister.

13. (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—
Drugs (Prevention of Misuse) Regulations

(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 authorised by regulations other than regulation 5(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 genuine.

(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);

(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,

but 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 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 subregulation (2) are –

(a) a practitioner;
(b) the 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 of a ship, or the master of a ship which does not carry a doctor on board as part of her complement;
(e) the master of a foreign ship in a port in Dominica.

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

(a) where furnished by the matron of a hospital or nursing home, be signed by a doctor or dentist employed or engaged in that hospital or nursing home;
(b) when furnished by the master of a foreign ship, contain a statement, signed by the proper officer of the port health authority within whose jurisdiction the ship is, that the quantity of the drug 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 for the time being in charge of any ward, theatre or other department of 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 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 drugs specified in the First Schedule or poppy-straw.

14. (1) Subject to this regulation, a person shall not issue a prescription containing a controlled drug other than a drug specified in the First Schedule.
Schedule 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 practitioner for a total quantity intended to be dispensed by instalments contain a direction specifying the amount of the instalments and 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 (whether personally or as a member of a class) for the purposes of this subregulation by the Minister.

(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.

15. (1) A person shall not supply a controlled drug other than a drug specified in the First Schedule on a prescription —

(a) unless the prescription complies with regulation 14;

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

(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 reasonable 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 the First Schedule 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.

(3) In the case of a prescription containing a controlled drug other than a drug specified in the First Schedule, 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 paragraph (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;

(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”. 

L.R.O. 1/1991
16. (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 drugs specified in the First Schedule or poppy-straw or in relation to the supply of a controlled drug by or on the prescription of a practitioner.

17. (1) Subject to subregulation (3) and regulation 19 every person authorised by or under regulation 4 or 7 to supply any drug specified in the Second or Fourth Schedule shall comply with the following requirements, that is to say –

(a) he shall in accordance with this regulation and regulation 18, keep a register and shall enter therein in chronological sequence particulars of every quantity of such a drug supplied (whether by way of administration or otherwise) by him whether to persons within or outside Dominica;

(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, 3 and 6 of the Second Schedule and paragraphs 1 and 3 of the Fourth Schedule together with it 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 drugs or strengths of drugs comprised within the class of drugs to which that register or separate part relates.

(3) Subregulations (1) and (2) shall not have effect in relation to—

(a) a person licensed under regulation 4 to supply drugs, where the licence so directs; or

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

18. Any person required to keep a register under regulation 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 regulation 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 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 Minister or by any person authorised in writing by the Minister in that behalf—

(i) furnish such particulars as may be requested in respect of the obtaining or supplying by him of any
drug specified in the Second or Fourth Schedule, or in respect of any stock of such drugs in his possession;

(ii) produce the said register and such other books or documents in his possession relating to any dealings in drugs specified in the Second or Fourth Schedule 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 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 Minister, be kept in respect of each department of the business carried on by him;

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

19. (1) Where a drug specified in the Second Schedule is supplied in accordance with regulation 7(5)(a) to a member of the crew of a ship, an entry in the official log book of the ship 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 record specified the drug supplied.

(2) A midwife authorised 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.
20. (1) A producer of any drug specified in the First Schedule 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 drug specified in the First Schedule 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; and 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 subregulation as if it were the keeping of the original document.

21. (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 drug specified in the Second or Fourth Schedule 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 subregulation by the Minister (hereafter in this regulation referred to as an “authorised person”).

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

(3) Where a drug specified in the Second or Fourth Schedule 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 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 the Second Schedule which he no longer requires, he shall not destroy the drug or cause it to be destroyed but shall dispose of it to the Minister.
Drugs (Prevention of Misuse) Regulations

CONTROLLED DRUGS EXCEPTED FROM THE PROHIBITION ON IMPORTATION, EXPORTATION AND POSSESSION

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 compound 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, micodidodine (6-nicotinoyldihydrocodeine), norcodeine, pholcodine and their respective salts.

2. 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 compound 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.

3. Any preparation of difenoxin (1-(3-cyano-3, 3-diphenylpropyl) -4-phenylpiperidine-4-carboxylic acid) containing, per dosage unit, not more than 0.5 milligrammes of difenoxin and a quantity of atropine sulphate equivalent to at least 5 per cent of the dose of difenoxin.

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

6. 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 powdered ingredient containing no controlled drug.

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

1. The following substances and products, namely:

<table>
<thead>
<tr>
<th>Acetorphine</th>
<th>Ethylmethylthiambutene</th>
</tr>
</thead>
<tbody>
<tr>
<td>Allylprodine</td>
<td>Etonitazene</td>
</tr>
<tr>
<td>Alphacetylmethadol</td>
<td>Etorphine</td>
</tr>
<tr>
<td>Alphameprodine</td>
<td>Etoxeridine</td>
</tr>
<tr>
<td>Alphamethadol</td>
<td>Fentanyl</td>
</tr>
<tr>
<td>Alphaprodine</td>
<td>Furethidine</td>
</tr>
<tr>
<td>Anileridine</td>
<td>Hydromorphan</td>
</tr>
<tr>
<td>Benzethidine</td>
<td>Hydroxyethidine</td>
</tr>
<tr>
<td>Benzylmorphine</td>
<td>Isomethadone</td>
</tr>
<tr>
<td>(3-benzylmorphine)</td>
<td>Ketobemidone</td>
</tr>
<tr>
<td>Betacetylmethadol</td>
<td>Levomethorphan</td>
</tr>
<tr>
<td>Betameprodine</td>
<td>Levomoramide</td>
</tr>
<tr>
<td>Betamethadol</td>
<td>Levophenacylmorphan</td>
</tr>
<tr>
<td>Betaprodine</td>
<td>Levarphanol</td>
</tr>
<tr>
<td>Bezitramide</td>
<td>Medicinal opium</td>
</tr>
<tr>
<td>Clonitazene</td>
<td>Metazocine</td>
</tr>
<tr>
<td>Cocaine</td>
<td>Methadone</td>
</tr>
<tr>
<td>Desomorphine</td>
<td>Methadyl acetate</td>
</tr>
<tr>
<td>Dextromoramide</td>
<td>Methyldesorphine</td>
</tr>
<tr>
<td>Diamorphine</td>
<td>Methyldihydromorphine</td>
</tr>
<tr>
<td>Diampromide</td>
<td>(6-methyldihydromorphine)</td>
</tr>
<tr>
<td>Diethylthiambutene</td>
<td>Metopon</td>
</tr>
<tr>
<td>Dihydromorphine</td>
<td>Morpheridine</td>
</tr>
<tr>
<td>Dimenoxadole</td>
<td>Morphine methobromide,</td>
</tr>
<tr>
<td>Diphephtanol</td>
<td>morphine</td>
</tr>
<tr>
<td>Dimethyllaibutene</td>
<td>Myrophine</td>
</tr>
<tr>
<td>Difefoxin (4-(3-cyano-</td>
<td>Nicomorphine</td>
</tr>
<tr>
<td>3, 3-diphenyl-propyl)</td>
<td>Noracymethadol</td>
</tr>
<tr>
<td>-4-phenylpiperidine-</td>
<td>Norlevorphanol</td>
</tr>
<tr>
<td>-4-carboxylic acid)</td>
<td>Normorphine</td>
</tr>
<tr>
<td>Dioxaphetyl butyrate</td>
<td>Norpipanone</td>
</tr>
<tr>
<td>Diphenoxy</td>
<td>N-oxide and other</td>
</tr>
<tr>
<td>Dipipanone</td>
<td>pentavalent nitrogen</td>
</tr>
<tr>
<td>Drotebanol (3, 4-dimet</td>
<td>1-Cyano-2-dimethyl-</td>
</tr>
<tr>
<td>morphinan-6B, 14-diol</td>
<td>morphine derivatives</td>
</tr>
<tr>
<td>Ecgonine, and any derivative</td>
<td>Oxycodone</td>
</tr>
<tr>
<td>of ecgonine which is</td>
<td>Oxymorphine</td>
</tr>
<tr>
<td>convertible to ecgonine</td>
<td>1-Cyano-2-dimethyl-</td>
</tr>
<tr>
<td>or to cocaine</td>
<td>amino-4, 4-diphenylbutane</td>
</tr>
</tbody>
</table>
4-Cyano-4-methyl-4-phenylpiperidine
4-Cyano-4-methyl-4-phenylpiperidine-4-carboxylic acid
2-Methyl-3-morpholino-4,1-diphenyl-propanecarboxylic acid
4-Phenylpiperidine-4-carboxylic acid ethyl ester
Pethidine
Phendoxone
Phenampromide

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 the First Schedule.

6. The following substances and products, namely:

   Acetyldihydrocodeine
   Amphetamine
   Codeine
   Dexamphetamine
   Dicyclomine
   Ethylmorphine
   (3-ethylmorphine)
   Methaqualone
   Methylandroxetine
   Methylnaltrexone
   Methylphenidate
   Methylphenidate (6-methoxy)
   Methylphenidate (6-nitro)
   Methylphenidate (6-p-tolyl)
   Nicocodine
   Nicocodine (6-
   nicotinoyl-dihydro-codeine)
   Norcodeine
   Phenmetrazine
   Pholcodine
   Propiram.


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 the First Schedule.
THIRD SCHEDULE

1. The following substances, namely —
   Benzphetamine
   Phendimetrazine
   Chlorphenamine
   Pipradrol
   Mephentermine.

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 specified in any of paragraphs 1 to 3, not being a preparation specified in the First Schedule.

FOURTH SCHEDULE

1. The following substances and products, namely —
   (a) Bufoteine
   Cannabinol
   Cannabinol derivatives
   Cannabis and cannabis resin
   Coca leaf
   Concentrate of poppy-straw
   Lysergamide
   Lysergide and other N-alkyl derivatives of lysergamide
   Mescaline
   Psilocin
   Raw opium
   4-Bromo-2,5-dimethoxy-x-methylphenethylamine
   N, N-Diethyltryptamine
   N, N-Dimethyltryptamine
   2,5-Dimethoxy-x-4-dimethylphenethylamine;
   (b) any compound (not being a compound for the time being specified in subparagraph (a) above structurally derived from tryptamine or from a ring-hydroxytryptamine by substitution 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;

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

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

3. Any ester or other 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 the First Schedule.
DRUGS (NOTIFICATION OF AND SUPPLY TO ADDICTS) REGULATIONS

made under section 33

[29th December 1988]

1. These Regulations may be cited as the –

DRUGS (NOTIFICATION OF AND SUPPLY TO ADDICTS) REGULATIONS.

2. (1) In these Regulations, “drug” means a controlled drug specified in the Schedule.

(2) For the purposes of these Regulations, a person shall be regarded as being addicted to a drug if, and only if, he has as a result of repeated administration become so dependent upon the drug that he has an overpowering desire for the administration of it to be continued.

(3) In these Regulations any reference to any written law shall be construed as a reference to that written law, and as including a reference thereto as extended or applied, by or under any other written law.

3. (1) Subject to subregulation (2), any doctor who attends a person whom he considers or has reasonable grounds to suspect is addicted to any drug shall, within seven days of the attendance, furnish in writing to the Chief Medical Officer such of the following particulars with respect to that person as are known to the doctor, that is to say, the name, address, sex and date of birth of that person, the date of the attendance and the name of the drug or drugs concerned.

(2) It shall not be necessary for a doctor who attends a person to comply with subregulation (1) in respect of that person if –

(a) the doctor is of the opinion, formed in good faith, that the continued administration of the drug or drugs concerned is required for the purpose of treating organic disease or injury; or

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the particulars which, apart from this subregulation, would have been required under these provisions to be furnished have, during the period of twelve months ending with the date of the attendance, been furnished in compliance with those provisions –

(i) by the doctor; or

(ii) if the doctor is a partner in or employed by a firm of general practitioners, by a doctor who is a partner in or employed by that firm; or

(iii) if the attendance is on behalf of another doctor, whether for payment or otherwise, by that doctor; or

(iv) if the attendance is at a hospital, by a doctor on the staff of that hospital.

4. (1) Subject to subregulation (2), a doctor shall not administer or supply to a person whom he considers, or has reasonable grounds to suspect, is addicted to any drug, or authorise the administration or supply to such a person of any substance specified in subregulation (3) or prescribe for such a person any such substance, except –

(a) for the purpose of treating organic disease or injury; or

(b) under and in accordance with the terms of a licence issued by the Minister in pursuance of these Regulations.

(2) Subregulation (1) shall not apply to the administration or supply by a doctor of a substance specified in subregulation (3) if the administration or supply is authorised by another doctor under and in accordance with the terms of a licence issued to him in pursuance of these Regulations.

(3) The substance referred to in subregulations (1) and (2) are –

(a) cocaine, its salts and any preparation or other product containing cocaine or its salts other than a preparation falling within paragraph 2 of the First Schedule to the Drugs (Prevention of Misuse) Regulations;

(b) diamorphine, its salts and any preparation or other product containing diamorphine or its salts.
CONTROLLED DRUGS TO WHICH THESE REGULATIONS APPLY

1. The following substances and products, namely —
   Cocaine       Hydromorphone       Oxycodone
   Dextromoramide  Levorphanal       Pethidine
   Diamorphine  Methadone         Phenazocine
   Dipipanone     Morphine         Piritramide
   Hydrocodone    Opium

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

3. Any ester or ether of a substance specified in paragraph 1 or 2 above not being a substance for the time being specified in Part II of the Third Schedule to the Drugs (Prevention of Misuse) Act.

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

5. Any preparation or other product containing a substance or product specified in any of paragraphs 1 to 4 above.