Number 12 of 1977

MISUSE OF DRUGS ACT 1977

REVISED

Updated to 25 May 2018

This Revised Act is an administrative consolidation of the Misuse of Drugs Act 1977. It is prepared by the Law Reform Commission in accordance with its function under the Law Reform Commission Act 1975 (3/1975) to keep the law under review and to undertake revision and consolidation of statute law.

All Acts up to and including Data Protection Act 2018 (7/2018), enacted 24 May 2018, and all statutory instruments up to and including Data Protection Act 2018 (Establishment Day) Order 2018 (S.I. No. 175 of 2018), made 24 May 2018, were considered in the preparation of this Revised Act.

Disclaimer: While every care has been taken in the preparation of this Revised Act, the Law Reform Commission can assume no responsibility for and give no guarantees, undertakings or warranties concerning the accuracy, completeness or up to date nature of the information provided and does not accept any liability whatsoever arising from any errors or omissions. Please notify any errors, omissions and comments by email to revisedacts@lawreform.ie.
Introduction

This Revised Act presents the text of the Act as it has been amended since enactment, and preserves the format in which it was passed.

Related legislation

**Health Acts 1947 to 2015**: this Act is one of a group of Acts included in this collective citation, to be construed together as one (Health (General Practitioner Service) Act 2015 (19/2015), s. 4(2)). The Acts in this group are:

- Health Act 1947 (28/1947)
- Health Act 1953 (26/1953) (citation only)
- Health (Fluoridation of Water Supplies) Act 1960 (46/1960) (citation only)
- Health Act 1970 (1/1970)
- Misuse of Drugs Act 1977 (12/1977), ss. 36 and 42 (in so far as it amends the Health Acts 1947 to 1970) (citation only)
- Health (Family Planning) Act 1979 (20/1979)
- Health (Nursing Homes) Act 1990 (23/1990)
- Health (Amendment) Act 1991 (15/1991), other than s. 8
- Health (Amendment) Act 1996 (15/1996)
- Health (Amendment) (No. 2) Act 1996 (23/1996)
- Health (Amendment) (No. 3) Act 1996 (32/1996), other than ss. 21 and 22
- Health (Eastern Regional Health Authority) Act 1999 (13/1999)
- Health (Miscellaneous Provisions) Act 2001 (14/2001), except in so far as it relates to the Tobacco (Health Promotion and Protection) Act 1988 (citation only)
- Health Act 2004 (42/2004)
- Health (Amendment) Act 2005 (3/2005), in so far as it amends the Health Acts 1947 to 2004
- Health (Repayment Scheme) Act 2006 (17/2006)
- Hepatitis C Compensation Tribunal (Amendment) Act 2006 (22/2006), s. 6
- Health (Nursing Homes) (Amendment) Act 2007 (1/2007)
- Health Act 2007 (23/2007)
- Medical Practitioners Act 2007 (25/2007), s. 57(9) (citation only)
- Health Act 2008 (21/2008)
- Health (Miscellaneous Provisions) Act 2009 (25/2009), s. 64
• Health (Amendment) Act 2010 (15/2010) (citation only)
• Health (Amendment) (No. 2) Act 2010 (20/2010)
• Child Care (Amendment) Act 2011 (19/2011), ss. 35 and 36 (citation only)
• Health (Alteration of Criteria for Eligibility) Act 2013 (10/2013)
• Health (Pricing and Supply of Medical Goods) Act 2013 (14/2013), s. 30 (citation only)
• Health Service Executive (Governance) Act 2013 (23/2013)
• Health (Alteration of Criteria for Eligibility) (No. 2) Act 2013 (42/2013) (citation only)
• Local Government Reform Act (1/2014), ss. 1(14) and 5(6) (in so far as it amends the Health (Fluoridation of Water Supplies) Act 1960) and sch. 2 part 6
• Health Service Executive (Financial Matters) Act 2014 (17/2014)
• Health (General Practitioner Service) Act 2014 (28/2014)
• Health (General Practitioner Service) Act 2015 (19/2015)

Acts previously included in the group but now repealed are:

• Health Act 1954 (23/1954)
• Health and Mental Treatment Act 1957 (16/1957), s. 1
• Health and Mental Treatment (Amendment) Act 1958 (37/1958), s.1
• Health (Homes For Incapacitated Persons) Act 1964 (8/1964)
• Health and Mental Treatment (Amendment) Act 1966 (2/1966), s. 1
• Health (Mental Services) Act 1981 (17/1981)
• Health (Family Planning) (Amendment) Act 1985 (4/1985)
• Health (Amendment) Act 2004 (19/2004)

Misuse of Drugs Acts 1977 to 2017: this Act is one of a group of Acts included in this collective citation, to be construed together as one (Misuse of Drugs (Supervised Injecting Facilities) Act 2017 (7/2017), s. 13(2)). The Acts in this group are:

• Misuse of Drugs Act 1977 (12/1977)
• Misuse of Drugs Act 1984 (18/1984)
• Criminal Justice Act 1999 (10/1999), Part II
• Criminal Justice Act 2006 (26/2006), ss. 80-85
• Criminal Justice Act 2007 (29/2007), s. 33
• Misuse of Drugs (Amendment) Act 2015 (6/2015)
• Misuse of Drugs (Amendment) Act 2016 (9/2016)
• Misuse of Drugs (Supervised Injecting Facilities) Act 2017 (7/2017)

Pharmacopoeia Acts 1931 and 1977: this Act is one of a group of Acts included in this collective citation, to be construed together as one (Misuse of Drugs Act 1977 (12/1977), s. 43(4)). The Acts in this group are:

• Pharmacopoeia Act 1931 (22/1931)
• Misuse of Drugs Act 1977 (12/1977), ss. 35 and 42 (in so far as s. 42 amends the Pharmacopoeia Act 1931)

Poisons Acts 1961 and 1977: this Act is one of a group of Acts included in this collective citation, to be construed together as one (Misuse of Drugs Act 1977 (12/1977), s. 43(6)). The Acts in this group are:

• Poisons Act 1961 (12/1961)
• Misuse of Drugs Act 1977 (12/1977), s. 33

Pharmacy Acts 1875 to 1977: this Act is one of a group of Acts previously included in this collective citation, to be construed together as one (Misuse of Drugs Act 1977 (12/1977), s. 43(3)). The substantive provision of this Act (s. 32) as well as legislation amended by provisions of this Act included in the collective citation (s. 34) were repealed (29.11.2008) by Pharmacy Act 2007 (20/2007), s. 4 and sch. 4, S.I. No. 487 of 2008.

Annotations

This Revised Act is not annotated and only shows textual amendments. An annotated version of this revision is also available which shows textual and non-textual
amendments and their sources. It also shows editorial notes including statutory
instruments made pursuant to the Act and previous affecting provisions.

Material not updated in this revision

Where other legislation is amended by this Act, those amendments may have been
superseded by other amendments in other legislation, or the amended legislation
may have been repealed or revoked. This information is not represented in this
revision but will be reflected in a revision of the amended legislation if one is
available. A list of legislative changes to any Act, and to statutory instruments from
1986, may be found linked from the page of the Act or statutory instrument at
www.irishstatutebook.ie.
ARRANGEMENT OF SECTIONS

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SCHEDULE

ACTS REFERRED TO

<p>| Dentists Act, 1928 | 1928, No. 25 |
| Pharmacy Act, 1951 | 1951, No. 30 |
| Amendment Act, 1890 | 1890, c. 48 |
| Medical Practitioners Act, 1927 | 1927, No. 25 |
| Pharmacy Act (Ireland), 1875 | 1875, c. 57 |
| Veterinary Surgeons Act, 1931 | 1931, No. 36 |
| Pharmacy Act, 1962 | 1962, No. 14 |
| Pharmacopoeia Act, 1931 | 1931, No. 22 |
| Health Act, 1947 | 1947, No. 28 |
| Health Act, 1953 | 1953, No. 26 |</p>
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<td>Companies Act, 1963</td>
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<td>Dangerous Drugs Act, 1934</td>
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BE IT ENACTED BY THE OIREACHTAS ASfollows:

Interpretation.

1.—(1) In this Act—

['business’ includes a profession;]

['cannabis’ (except in ‘cannabis resin’) means any plant of the genus Cannabis or any part of any such plant (by whatever name designated) but includes neither cannabis resin nor any of the following products after separation from the rest of any such plant, namely—

(a) mature stalk of any such plant,

(b) fibre produced from such mature stalk, or

(c) seed of any such plant;]

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

“the Dental Board” means the Dental Board established under the Dentists Act, 1928;

“duly issued prescription” has the meaning assigned to it by section 18 of this Act;

“forged prescription” has the meaning assigned to it by section 18 of this Act;

['Irish Medicines Board’ means the Irish Medicines Board established under section 3 of the Irish Medicines Board Act 1995;]

“land” includes land covered wholly or partly with water;

“the Medical Registration Council” means the Medical Registration Council established under the Medical Practitioners Acts, 1927 to 1961;
“the Minister” means the Minister for Health;

[‘opium poppy’ means a plant of the species Papaver somniferum L or Papaver bracteatum Lindl;]

“pharmacist” means a registered pharmaceutical chemist, a registered dispensing chemist and druggist and a registered druggist;

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

[“practitioner” means a registered medical practitioner, a registered dentist, a registered veterinary surgeon, a registered nurse and a registered midwife;]

“prescribed” means prescribed by regulations made by the Minister under this Act;

“registered dentist” means a person registered in the register established under the Dentists Act, 1928;

“registered dispensing chemist and druggist” means a person registered in the register of dispensing chemists and druggists established under the Pharmacy Act, 1951;

“registered druggist” means a person registered in the register of registered druggists in Ireland established under the Pharmacy Act (Ireland), 1875, Amendment Act, 1890;

“registered medical practitioner” means a person registered in the register established under the Medical Practitioners Act, 1927;

[“registered midwife” means a midwife whose name is entered in the midwives division of the register of nurses and midwives under the Nurses and Midwives Act 2011;]

[“registered nurse” means a nurse whose name is entered in the nurses division of the register of nurses and midwives under the Nurses and Midwives Act 2011;]

“registered pharmaceutical chemist” means a person registered in the register of pharmaceutical chemists for Ireland established under the Pharmacy Act (Ireland), 1875;

“registered veterinary surgeon” means a person registered in the register established under the Veterinary Surgeons Act, 1931;

“registration authority” means such one of the following as the context requires namely, the Dental Board, the Medical Registration Council and the Veterinary Council;

“the respondent” in relation to a reference under section 8 or section 9 of this Act means the practitioner in respect of whom the reference is made;

“special direction” has the meaning assigned to it by section 7 (2) of this Act;

“supply” includes giving without payment;

[‘temporary direction’ means a direction under section 9 of this Act;]

[‘vessel’ includes a hovercraft;]

“the Veterinary Council” means the Veterinary Council established under the Veterinary Surgeons Act, 1931.

(2) For the purposes of this Act any controlled drug, pipe, utensil or document of which a person has control and which is in the custody of another who is either under the person’s control or, though not under the person’s control, acts on his behalf, whether as an agent or otherwise, shall be regarded as being in the possession of the person, and the provisions of section 16 and section 18 together with the provisions
of this Act relating to the possession of controlled drugs shall be construed and have
effect in accordance with the foregoing.

Controlled drugs. 2.—(1) In this Act “controlled drug” means any substance, product or preparation
(other than a substance, product or preparation specified in an order under subsection
(3) of this section which is for the time being in force) which is either specified in the
Schedule to this Act or is for the time being declared pursuant to subsection (2) of
this section to be a controlled drug for the purposes of this Act.

(2) The Government may by order declare any substance, product or preparation
(not being a substance, product or preparation specified in the Schedule to this Act)
to be a controlled drug for the purposes of this Act and so long as an order under this
subsection is in force, this Act shall have effect as regards any substance, product or preparation specified in the order as if the substance, product or preparation were
specified in the said Schedule.

(3) The Government may by order declare that the provisions of this Act shall not
apply in relation to a substance, product or preparation specified both in the order
and in the Schedule to this Act, and so long as an order under this subsection is in
force, this Act shall not apply in relation to a substance, product or preparation
specified in the order.

(4) The Government may by order amend or revoke an order under this section
(including an order made under this subsection).

Restriction on
possession of
controlled drugs. 3.—(1) Subject to subsection (3) of this section and section 4 (3) of this Act, a person
shall not have a controlled drug in his possession.

(2) A person who has a controlled drug in his possession in contravention of
subsection (1) of this section shall be guilty of an offence.

(3) The Minister may by order declare that subsection (1) of this section shall not
apply to a controlled drug specified in the order, and for so long as an order under
this subsection is in force the prohibition contained in the said subsection (1) shall
not apply to a drug which is a controlled drug specified in the order.

(4) The Minister may by order amend or revoke an order under this section
(including an order made under this subsection).

Regulations
permitting
possession of
controlled drugs. 4.—(1) The Minister may make regulations enabling any person, or persons of a
prescribed class or description, in prescribed circumstances or for prescribed
purposes, to possess a controlled drug subject to such conditions (if any), or subject
to and in accordance with such licence, as may be prescribed.

(2) Subject to section 13 of this Act, the Minister shall exercise his power to make
regulations under this section so as to secure that it is not unlawful under this Act
for a practitioner or pharmacist to have a controlled drug in his possession for the
purpose of his profession or business.

(3) It shall be lawful for any person, or a person of a class or description specified
in regulations under this section, to have in his possession in prescribed circumstances
or for prescribed purposes, as may be appropriate, a controlled drug specified
therein, provided that any conditions specified in the regulations or attached to a
licence granted under this Act and applicable in the particular case are complied with
by him.

Regulations to
prevent misuse of
controlled drugs. 5.—(1) For the purpose of preventing the misuse of controlled drugs, the Minister
may make regulations—
(a) prohibiting absolutely, or permitting subject to such conditions or exceptions as may be specified in the regulations, or subject to any licence, permit or other form of authority as may be so specified—

(i) the manufacture, production or preparation of controlled drugs,

(ii) the importation or exportation of controlled drugs,

(iii) the supply, the offering to supply or the distribution of controlled drugs,

(iv) the transportation of controlled drugs,

(b) requiring prescribed documents to be used in a prescribed manner in relation to prescribed transactions concerning controlled drugs and requiring copies of such documents to be furnished to prescribed persons, or to persons of a prescribed class or description,

(c) requiring prescribed precautions to be taken for the purpose of ensuring the safe custody of controlled drugs,

(d) requiring prescribed records to be kept in relation to controlled drugs and regulations under this section may specify the manner in which the records are to be kept and maintained and such regulations may also provide for the furnishing of information relating to such records in such circumstances and in such manner as may be prescribed,

(e) providing for the inspection by prescribed persons of precautions taken or records kept in pursuance of regulations under this section,

[(f) subject to subsection (1A), regulating the issue of prescriptions for controlled drugs and the supply of controlled drugs on prescription by—

(i) registered medical practitioners, registered dentists or registered veterinary surgeons,

(ii) relevant nurses, or

(iii) relevant midwives.]

(g) requiring persons dispensing prescriptions for controlled drugs to furnish to the Minister such information relating to those prescriptions as may be prescribed,

(h) regulating or controlling the packaging and labelling of controlled drugs and such regulations may in particular require prescribed particulars relating to controlled drugs or a prescribed statement (including a warning or caution) relating to such drugs to be printed either on the outside of any packet or container used in the sale, supply or distribution of controlled drugs or on a label attached to such packet or container,

(i) requiring that any controlled drugs which, because of their condition or for any other reason, are not intended to be used shall be destroyed or disposed of in a prescribed manner,

(j) requiring any manufacturer, manufacturer's agent or wholesaler who wishes to withdraw a controlled drug from public sale to give six months notice of such proposed withdrawal unless the Minister is satisfied that it is in the public interest that such controlled drug should be withdrawn at such shorter notice as the Minister may determine.

[(1A) The Minister shall not make regulations under subsection (1)(f)(ii) or (iii) unless the Minister, having had regard to the nature and purpose of the controlled drug (including any deleterious effects which may arise from the misuse thereof), is satisfied that it is reasonably safe to permit the issue of prescriptions for that controlled drug by relevant nurses or relevant midwives.]
(2) Subject to section 13 of this Act, the Minister shall exercise his power to make regulations under this section so as to secure that it is not unlawful under this Act for—

(a) a practitioner [other than a relevant nurse or a relevant midwife], for the purpose of his profession, to prescribe, administer, manufacture, compound or supply a controlled drug,

(b) a pharmacist, for the purpose of his profession or business, to manufacture, compound or supply a controlled drug,

provided that nothing in this subsection shall be construed as enabling the Minister to make regulations under this Act authorising a registered druggist to keep open shop for the compounding or dispensing of medical prescriptions.

[(3) Subject to section 13, the Minister may make regulations under this section so as to secure that it is not unlawful under this Act for a practitioner who is a relevant nurse or a relevant midwife, for the purpose of the practitioner’s profession as a relevant nurse or a relevant midwife, to prescribe, administer or supply a controlled drug if the Minister, after having had regard to the nature and purpose of the controlled drug (including any deleterious effects which may arise from the misuse thereof), is satisfied that it is reasonably safe to permit the practitioner, for the purpose of the practitioner’s profession as a relevant nurse or a relevant midwife, to prescribe, administer or supply that controlled drug.]

[(4) In this section—

‘relevant midwife’ means a registered midwife or a class of registered midwives;

‘relevant nurse’ means a registered nurse or a class of registered nurses.]]

6.—(1) Where a practitioner or pharmacist has after the commencement of this subsection been convicted of—

(a) an offence under this Act, or

(b) an offence against the Customs Acts in relation to the importation or exportation of a controlled drug,

the Minister may give a direction under subsection (2) of this section in respect of that person.

[(1A) Where a relevant person has after the commencement of this subsection been convicted of—

(a) an offence under this Act, or

(b) an offence against the Customs Acts in relation to the importation or exportation of a controlled drug,

the Minister may give a direction under subsection (2) of this section in respect of that person.]

(2) A direction under this subsection shall—

(a) in case the direction relates to 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) in case the direction relates to a pharmacist [or relevant person, be a direction prohibiting the pharmacist or relevant person, as the case may be, from having in the pharmacist’s or relevant person’s, as the case may be,] 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) of this section, 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 Iris Oifigiúil and in such other manner (if any) as the
Minister may consider appropriate.

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

(6) Any person who contravenes a direction given under this section shall be guilty
of an offence.

[(7) In this section, ‘relevant person’ means—

(a) a person, not being a pharmacist, keeping open shop for the dispensing or
compounding of medical prescriptions in accordance with the provisions of
the Pharmacy Acts 1875 to 1977, or

(b) any director, manager, secretary or other official of a person referred to in
paragraph (a) of this definition which is a body corporate.]

Special directions
prohibiting
prescribing etc.
of controlled
drug in certain
cases.

7.—(1) If the Minister believes that a practitioner is or has been, after the
commencement of this section, prescribing, administering or supplying, or authorising
the administration or supply of any controlled drug in an irresponsible manner, subject
to the provisions of this Act, he may give a direction in respect of the practitioner
prohibiting him prescribing, administering or supplying or authorising the administra-
or supply of such controlled drugs as may be specified in the direction.

(2) A direction given pursuant to this section (in this Act subsequently referred to
as a special direction) shall come into force when a copy of it is given to the practi-
tioner to whom it relates and, subject to subsection (3) of this section and section 11
(1) of this Act, the special direction shall remain in operation until it is cancelled.

(3) The Minister may [...] suspend the operation of a special direction.

(4) The Minister may [...] cancel a special direction.

(5) Where the Minister suspends the operation of or cancels a special direction, he
shall cause notice to that effect to be given to the practitioner to whom the special
direction applies and, as soon as may be, cause notice of the suspension or cancellation
to be published in the Iris Oifigiúil.

(6) A person who contravenes a special direction shall be guilty of an offence.

Investigation of
cases where
Minister consid-
ers there are
grounds for
special direction.

8.—(1) If the Minister considers that there may be grounds for giving a special
direction, he shall forthwith establish a committee of inquiry, constituted in accordance
with any regulations under section 12 of this Act which apply to it, and as soon as
may be after such committee is established he shall refer the matter in question to
the committee for investigation and when making the reference send to the
committee a statement of such grounds, and it shall be the duty of the committee in
accordance with this section to investigate the matter referred to it and to report on
it to the Minister.

(2) Where the Minister sends a statement of grounds to a committee of inquiry
established pursuant to this section, he shall at the same time send to the respondent
a copy of the statement and invite him to submit to the committee in writing, within
the period of twenty-one days commencing on the date on which the statement is
sent to the committee, any representations relating to the matter to be investigated
which he may then wish to make.

(3) (a) Where a committee of inquiry is established under this section, a meeting
of the committee of inquiry shall be convened by the Minister who shall at
the same time fix a day for the meeting, being a day which is neither earlier
than the seventh day after the expiration of the period referred to in
subsection (2) of this section nor later than the twenty-first day after such
expiration.

(b) Where the Minister convenes a meeting under this subsection, he shall at the
same time send to the respondent not less than seven days’ notice in writing
of the date, place and time fixed by the Minister for the meeting and the
notice shall also notify the respondent that he may make representations
to, and if he so wishes appear in person before, the committee of inquiry
concerned, be assisted by another person (whether so appearing or not) in
making such representations or have such representations made by another
person (whether so appearing or not) acting on his behalf.

(4) A committee of inquiry established under this section shall report to the Minister
on its investigation as soon as may be and shall state in the report whether or not
they recommend the giving of a special direction as regards the matter being investi-
gated, and in case the committee recommends the giving of such a direction they
shall indicate in their report either the controlled drugs which the committee
considers should be specified in the relevant special direction or that the committee
considers that such direction should apply to all controlled drugs.

(5) Having considered the report of the committee of inquiry established under this
section, the Minister may—

(a) decide to give in respect of the respondent a special direction specifying all
or any of the controlled drugs indicated in a recommendation of the
committee, or

(b) decide not to give a special direction,

and in case the Minister pursuant to this section decides not to give a special
direction, he shall notify the respondent accordingly.

(6) Where the Minister gives a special direction, he shall, as soon as may be, cause
a copy of the special direction to be served on the respondent and shall cause a copy
of the direction to be published in the Iris Oifigiúil and in such other manner (if any)
as the Minister may consider appropriate.

(7) Where the Minister gives a special direction, he shall send a copy of the report
received by him from the relevant committee of inquiry and the special direction to
the respondent and also to—

(a) in case the respondent is a registered dentist, the Dental Board,

(b) in case the respondent is a registered medical practitioner, the Medical
Council,

(c) in case the respondent is a registered veterinary surgeon, the Veterinary
Council,

[(d) in case the practitioner concerned is a registered nurse, to An Bord Altranais.]

[9.—(1) Where the Minister refers a matter for investigation to a committee of
inquiry established under section 8 of this Act, he may give a direction under this
section in respect of the respondent prohibiting his prescribing, administering or
supplying or authorising the administration or supply of such controlled drugs as may

Prohibition of
prescribing etc. in
cases of urgency.

[No. 12.]

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[1977.]
be specified in the direction, and such direction shall come into force on the expiration of
the period of seven days beginning on the day on which a copy of the direction is
sent by the Minister to the respondent unless, not later than the seventh day
following the day on which such copy is sent, the respondent satisfies the Minister
that the direction should not come into force.

(2) In case a copy of a temporary direction is sent by the Minister, the Minister shall
at the same time send to the respondent a notice in writing stating that the respondent
may, within the time limit specified in subsection (1) of this section, make representa-
tions to the Minister stating why the temporary direction should not come into
force.

(3) A temporary direction shall remain in force until the expiration of the period of
twenty-eight days beginning on the day on which it is given or until the Minister makes
a decision under section 8 (5) of this Act as regards the relevant case, whichever first
occurs.

(4) The Minister may extend or further extend, in either case for a period not
exceeding twenty-eight days, the period during which a particular temporary direction
is to remain in force.

(5) Where a temporary direction is given, extended or further extended, the
Minister shall, as soon as may be, cause a notice of the temporary direction, its
extension or further extension, as may be appropriate, to be published in the Iris
Oifigiúil and in such other manner (if any) as the Minister may consider appropriate.

Investigation on
initiative of
Dental Board,
Medical Regis-
tration Council or
Veterinary Coun-
cil.

10.—[...]

Appeals.

11.—(1) Any practitioner or pharmacist who is aggrieved by a direction under section
6 (2) of this Act, a special direction or a temporary direction may, not later than three
weeks after the day on which the direction under the said section 6 (2), special
direction or temporary direction, as the case may be, comes into force, appeal to the
High Court, and that Court may—

(a) by interim order suspend the operation of the direction under the said section
6 (2), special direction or temporary direction, either generally or in a
particular respect, until the final determination of the proceedings,

(b) confirm the direction under the said section 6 (2), special direction or temporary
direction with or without modification or cancel it.

(2) Where a direction under section 6 (2) of this Act, special direction or temporary
direction is suspended or cancelled or confirmed with modifications by the High Court,
the order of the Court shall not prejudice the validity of anything done on foot of the
direction prior to the making of the order.

(3) Where a direction under section 6 (2) of this Act, special direction or temporary
direction is suspended or cancelled by the High Court, the Minister shall as soon as
may be cause notice thereof to be published in the Iris Oifigiúil.

Regulations
(committees and
panels).

12.—(1) The Minister may, after consultation with any registration authority
concerned, make regulations in relation to the constitution and procedure of
committees of inquiry [...] established pursuant to section 8 [...] of this Act.
Subject to the provisions of this Act and to any regulations made by the Minister under this section and which apply to it, a committee [... ] referred to in subsection (1) of this section may regulate its procedure and business.

13.—(1) If in the case of any controlled drug the Minister is of the opinion that it is in the public interest—

(a) for the manufacture, production, preparation, sale, supply, distribution and possession of that drug to be either wholly unlawful or unlawful except for purposes of research or for other special purposes specified in an order under this section, or

(b) for it to be unlawful for any person who is either a practitioner or a pharmacist to have in his possession or to do in relation to that drug any of the things mentioned in section 5 (2) of this Act 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 section designating a controlled drug as one to which this subsection applies, section 4 (2) of this Act and the said section 5 (2) shall not apply as regards that drug.

(2) The Minister may by order revoke or amend any order under this section (including an order under this subsection).

14.—(1) The Minister may grant licences or issue permits or authorisations for any of the purposes of this Act, attach conditions to any such licence, permit or authorisation, vary such conditions and revoke any such licence, permit or authorisation.

(2) The Minister may make regulations requiring the payment of prescribed fees in respect of the grant or issue under this section of a licence, permit or authorisation.

15.—(1) Any person who has in his possession, whether lawfully or not, a controlled drug for the purpose of selling or otherwise supplying it to another in contravention of regulations under section 5 of this Act, shall be guilty of an offence.

(2) Subject to section 29 (3) of this Act, in any proceedings for an offence under subsection (1) of this section, where it is proved that a person was in possession of a controlled drug and the court, having regard to the quantity of the controlled drug which the person possessed or to such other matter as the court considers relevant, is satisfied that it is reasonable to assume that the controlled drug was not intended for the immediate personal use of the person, he shall be presumed, until the court is satisfied to the contrary, to have been in possession of the controlled drug for the purpose of selling or otherwise supplying it to another in contravention of regulations under section 5 of this Act.

15A.—(1) A person shall be guilty of an offence under this section where—

(a) the person has in his possession, whether lawfully or not, one or more controlled drugs for the purpose of selling or otherwise supplying the drug or drugs to another in contravention of regulations under section 5 of this Act, and

(b) at any time while the drug or drugs are in the person’s possession the market value of the controlled drug or the aggregate of the market values of the controlled drugs, as the case may be, amounts to [€13,000] or more.

(2) Subject to section 29(3) of this Act (as amended by section 6 of the Criminal Justice Act, 1999), in any proceedings for an offence under this section, where—
(a) it is proved that a person was in possession of a controlled drug, and

(b) the court, having regard to the quantity of the controlled drug which the person possessed or to such other matters that the court considers relevant, is satisfied that it is reasonable to assume that the controlled drug was not intended for his immediate personal use,

he shall be presumed, until the court is satisfied to the contrary, to have been in possession of the controlled drug for the purpose of selling or otherwise supplying it to another in contravention of regulations under section 5 of this Act.

(3) If the court is satisfied that a member of the Garda Síochána or an officer of customs and excise has knowledge of the unlawful sale or supply of controlled drugs, that member or officer, as the case may be, shall be entitled in any proceedings for an offence under this section to be heard and to give evidence as to—

(a) the market value of the controlled drug concerned, or

(b) the aggregate of the market values of the controlled drugs concerned.

[(3A) In any proceedings for an offence under this section, it shall not be necessary for the prosecutor to prove that a person knew that at any time while the controlled drug or drugs concerned were in the person’s possession that the market value of that drug or the aggregate of the market values of those drugs, as the case may be, amounted to €13,000 or more or that he or she was reckless in that regard.]

(4) No proceedings may be instituted under this section except by or with the consent of the Director of Public Prosecutions.

(5) In this section—

‘market value’, in relation to a controlled drug, means the price that drug could be expected to fetch on the market for the unlawful sale or supply of controlled drugs;

‘an officer of customs and excise’ has the same meaning as in section 6 of the Criminal Justice (Drug Trafficking) Act, 1996.

15B.— (1) A person shall be guilty of an offence where—

(a) the person imports one or more controlled drugs in contravention of regulations under section 5 of this Act, and

(b) at or about the time the drug or drugs are imported the market value of the controlled drug or the aggregate of the market values of the controlled drugs, as the case may be, amounts to €13,000 or more.

(2) If the court is satisfied that a member of the Garda Síochána or an officer of customs and excise has knowledge of the unlawful sale or supply of controlled drugs, that member or officer, as the case may be, shall be entitled in any proceedings for an offence under this section to be heard and to give evidence as to—

(a) the market value of the controlled drug concerned, or

(b) the aggregate of the market values of the controlled drugs concerned.

(3) In any proceedings for an offence under this section, it shall not be necessary for the prosecutor to prove that a person knew that at the time the person imported the controlled drug or drugs concerned that the market value of that drug or the aggregate of the market values of those drugs, as the case may be, amounted to €13,000 or more or that he or she was reckless in that regard.

(4) No proceedings may be instituted under this section except by or with the consent of the Director of Public Prosecutions.
In this section ‘market value’ and ‘an officer of customs and excise’ have the meanings they have in section 15A of this Act.

15C.— (1) A person shall be guilty of an offence where—

(a) the person, other than in accordance with regulations made under section 4 of this Act, conveys a controlled drug into a prison, children detention school or remand centre or to a person in the prison, school or centre,

(b) the person, other than in accordance with regulations made under section 4 of this Act, places a controlled drug in any place inside or outside a prison, children detention school or remand centre with intent that it shall come into the possession of a person in the prison, school or centre,

(c) the person throws or projects a controlled drug into a prison, children detention school or remand centre, or

(d) the person, while in the vicinity of a prison, children detention school or remand centre, has in his or her possession a controlled drug with intent to commit an act referred to in paragraph (a), (b) or (c) of this subsection.

(2) A person may be guilty of an offence under subsection (1) of this section irrespective of the quantity of the controlled drug concerned.

(3) Subject to section 29(3) of this Act, in any proceedings for an offence under subsection (1)(d) of this section, where—

(a) it is proved that a person was in possession of a controlled drug in the vicinity of a prison, children detention school or remand centre, as the case may be, and

(b) the court (or the jury, as the case may be), having regard to all the circumstances including the person’s proximity to the prison, school or centre, as the case may be, the packaging (if any) of the controlled drug and the time of the day or night concerned, is satisfied that it is reasonable to assume that the controlled drug was not intended for his or her immediate personal use,

he or she shall be presumed, until the court (or the jury, as the case may be) is satisfied to the contrary, to have been in possession of the controlled drug with intent to commit an act referred to in paragraph (a) or (b) or, as the case may be, (c) of subsection (1) of this section.

(4) In any proceedings for an offence under subsection (1) of this section, it shall not be necessary for the prosecutor to prove that the controlled drug concerned was intended to come into the possession of any particular person in the prison, children detention school or remand centre, as the case may be.

(5) If a prison officer or an authorised member of the staff of a children detention school or remand centre reasonably suspects that a person has committed or is committing an offence under this section, he or she may, for the purpose of detecting the commission of such an offence, search the person at any time while he or she is in the prison, school or centre, as the case may be.

(6) A prison officer or an authorised member of the staff of a children detention school or remand centre may, for the purpose of performing his or her functions under subsection (5) of this section, have a controlled drug in his or her possession.

(7) A person guilty of an offence under this section shall be liable—

(a) on summary conviction, to a fine not exceeding €3,000 or imprisonment for a term not exceeding 12 months or both, or
(b) on conviction on indictment, to a fine or imprisonment for a term not exceeding 10 years or both.

(8) In this section—

‘an authorised member of the staff’—

(a) in relation to a children detention school, means a member of the staff of the school who is authorised in writing for the purposes of this section by the Director (within the meaning of section 157 of the Children Act 2001) of the school, and

(b) in relation to a remand centre, means a member of the staff of the centre who is authorised in writing for the purposes of this section by the owners or, as the case may be, the managers of the centre;

‘children detention school’ and ‘remand centre’ have the meanings they have in section 3(1) of the Children Act 2001;

‘prison’ means a place of custody administered by the Minister for Justice, Equality and Law Reform and includes Saint Patrick’s Institution and a place of detention provided under section 2 of the Prisons Act 1970, and ‘prison officer’, in relation to a prison, shall be construed accordingly.

16.—(1) A person shall not—

(a) smoke or otherwise use prepared opium,

(b) frequent a place used for the purpose of smoking or otherwise using prepared opium, or

(c) 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

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

(2) A person who contravenes a provision of subsection (1) of this section shall be guilty of an offence.

17.—(1) A person shall not cultivate opium poppy [for the production of opium], any plant of the genus Cannabis or any plant of the genus Erythroxylon except under and in accordance with a licence issued in that behalf [under section 14(1)].

[(2) Every person who cultivates opium poppy [for the production of opium], a plant of the genus Cannabis or a plant of the genus Erythroxylon in contravention of subsection (1) of this section shall be guilty of an offence.]

18.—(1) A person shall not forge a document purporting to be a prescription issued by a practitioner (which document is in this Act referred to as a forged prescription).

(2) A person shall not with intent to deceive either alter or use a prescription which has been duly issued by a practitioner (which document is in this Act referred to as a duly issued prescription).

(3) A person shall not have in his possession either a forged prescription or a duly issued prescription which has been altered with intent to deceive.
(4) The Minister may by regulations declare that in circumstances specified in the regulations subsection (3) of this section shall not apply in relation to persons who are of a prescribed class or description, and for so long as regulations under this subsection are in force the said subsection (3) shall be construed in accordance with and have effect subject to the regulations.

(5) A person who contravenes a provision of this section shall be guilty of an offence.

19.—(1) A person who is the occupier or is in control or is concerned in the management of any land, vehicle or vessel and who knowingly permits or suffers any of the following to take place on the land, vehicle or vessel, namely—

(a) the cultivation contrary to section 17 of this Act of opium poppy or any plant of the genus Cannabis,

(b) the preparation of opium for smoking,

(c) the preparation of cannabis for smoking,

(d) the smoking of cannabis, cannabis resin or prepared opium,

(e) the manufacture, production or preparation of a controlled drug in contravention of regulations made under section 5 of this Act,

(f) the importation or exportation of a controlled drug in contravention of such regulations,

(g) the sale, supply or distribution of a controlled drug in contravention of such regulations,

(h) any attempt so to contravene such regulations, or

(i) the possession of a controlled drug in contravention of section 3 of this Act,

shall be guilty of an offence.

(2) In any proceedings for an offence under subsection (1) of this section, where it is proved that an activity or contravention mentioned in the said subsection (1) took place on particular land or on a particular vehicle or vessel and that the defendant was, at the time of the alleged offence, the occupier of, or in control or concerned in the management of the land, vehicle or vessel, as the case may be, it shall be presumed until the court is satisfied to the contrary that the activity or contravention took place with the knowledge of the defendant.

19. — Occupiers etc. permitting certain activities to take place on land, vehicle or vessel to be guilty of an offence.

20.—(1) Any person who aids, abets, counsels or induces the commission in a place outside the State of an offence punishable under a corresponding law in force in that place shall be guilty of an offence.

(2) In this section “a corresponding law” means a law stated in a certificate purporting to be issued by or on behalf of the government of a country outside the State to be a law providing for the control or regulation in that country of the manufacture, production, supply, use, exportation or importation of dangerous or otherwise harmful drugs in pursuance of any treaty, convention, protocol or other agreement between states and prepared or implemented by, or under the auspices of, the League of Nations or the United Nations Organisation and which for the time being is in force.

(3) Any statement in a certificate mentioned in subsection (2) of this section as to the effect of the law mentioned in the certificate or any such statement that any facts constitute an offence against the law so mentioned shall, for the purposes of any proceedings under this Act, be evidence of the matters stated.

Offences relating to acts outside the State.
21.—(1) A person who attempts to commit an offence under this Act, or who aids, abets, counsels or procures the commission of an offence under this Act, or who solicits or incites any other person to commit an offence under this Act shall be guilty of an offence.

(2) A person who, whether by act or omission, contravenes or fails to comply with regulations under this Act shall be guilty of an offence.

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

(4) Any person who by act or omission impedes or obstructs a member of the Garda Síochána or a person duly authorised under this Act in the lawful exercise of a power conferred by this Act shall be guilty of an offence and if, in the case of a continuing offence, the impediment or obstruction is continued after conviction, he shall be guilty of a further offence.

(5) Any person who conceals from a person lawfully exercising a power under section 24 of this Act any controlled drug, or who without reasonable excuse fails to produce any book, record or other document which he has been duly required to produce under that section, shall be guilty of an offence.

(6) Any person who contravenes a condition attached to a licence, permit or authorisation granted or issued by the Minister under this Act (other than section 24) or under regulations made under this Act shall be guilty of an offence.

(7) Any person who, for the purpose of obtaining, whether for himself or another, the grant, issue or renewal of a licence, permit or authorisation under this Act or under regulations made under this Act—

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

shall be guilty of an offence.

22.—(1) In any proceedings for an offence under this Act, it shall not be necessary to negative by evidence the existence of any—

(a) order made under section 2 [or 3] of this Act,

(b) licence, permit or authorisation under this Act,

and accordingly the onus of proving the existence of any such licence, permit or authorisation shall be on the person seeking to avail himself thereof.

(2) In any proceedings for an offence under this Act it shall not be necessary for the prosecutor to prove that at the time of the offence—

(a) a defendant was not a person to whom regulations made under section 4 of this Act applied,

(b) a defendant was a person to whom an exception under regulations made under section 5 of this Act applied, and

in case a defendant claims that—

(i) by virtue of the said section 4 he had lawfully in his possession a controlled drug,
(ii) he is a person to whom such an exception applied,

the onus of proving such lawful possession, or that he is such a person, as may be appropriate, shall be on the defendant.

23.—(1) A member of the Garda Síochána who with reasonable cause suspects that a person is in possession in contravention of this Act of a controlled drug, may without warrant—

(a) search the person and, if he considers it necessary for that purpose, detain the person for such time as is reasonably necessary for making the search,

(b) search any vehicle, vessel or aircraft in which he suspects that such drug may be found [(and any substance, article or other thing on or in the vehicle, vessel or aircraft)] and for the purpose of carrying out the search may, if he thinks fit, require the person who for the time being is in control of such vehicle, vessel or aircraft to bring it to a stop and when stopped to refrain from moving it, or in case such vehicle, vessel or aircraft is already stationary, to refrain from moving it, or

(c) [examine (by opening or otherwise) and] seize and detain anything found in the course of a search under this section which with such cause appears to him to be something which might be required as evidence in proceedings for an offence under this Act.

[(1A) Where a member of the Garda Síochána decides to search a person under this section, he may require the person to accompany him to a Garda Station for the purpose of being so searched at that station.

(1B) Where a member of the Garda Síochána decides to search a vehicle, vessel or aircraft under this section he may as regards the person who appears to him to be the owner or in control or charge for the time being of the vehicle, vessel or aircraft make any one or more or all of the following requirements:

(a) require such person, pending the commencement of the search, not to remove from the vehicle, vessel or aircraft, as may be appropriate, any substance, article or other thing,

(b) in case the decision relates to a vehicle and the place at which he finds the vehicle is in his reasonable opinion unsuitable for such search, require such person forthwith to take the vehicle or cause it to be taken to a place which he considers suitable for such search and which is specified by him,

(c) require the person to be in or on or to accompany the vehicle, vessel or aircraft, as may be appropriate, for so long as the requirement under this paragraph remains in force.

(1C) Where there is a failure to comply with a requirement made under this section the following provisions shall apply—

(a) in case the requirement was made under subsection (1A) of this section, the member of the Garda Síochána concerned may arrest without warrant the person of whom the requirement was made, and

(b) in case the requirement is a requirement mentioned in paragraph (b) of subsection (1B) of this section, such member may take the vehicle concerned, or cause it to be taken, to a place which he considers suitable for a search under this section.

(1D) Where a requirement is made of a person under this section—

(a) in case the requirement is a requirement mentioned in paragraph (c) of subsection (1B) of this section, if at any time while the requirement is in
force the person of whom it was made is neither in nor on nor accompanying the vehicle, vessel or aircraft, as may be appropriate, in relation to which the requirement was made, he shall be guilty of an offence,

(b) in case of any other requirement under this section the person who fails to comply with the requirement shall be guilty of an offence.

(1E) A requirement mentioned in paragraph (c) of subsection (1B) of this section shall remain in force until the search in relation to which it is made is completed.

(1F) Where a requirement described in paragraph (a) of subsection (1B) of this section is made of a person, the search in relation to which the requirement is made shall be carried out as soon as is practicable.

(2) Nothing in this section shall operate to prejudice any power to search, or to seize or detain property which may be exercised by a member of the Garda Síochána apart from this section.

Powers to inspect and demand production of drugs, books or documents.

24.—(1) For the purpose of enforcing this Act and regulations made thereunder, a member of the Garda Síochána or a person authorised in writing in that behalf by the Minister or the Irish Medicines Board may at all reasonable times—

(a) enter any building or other premises in which a person carries on business as a producer, manufacturer, seller or distributor of controlled drugs [or as a practitioner],

(b) require any such person, or any person employed in connection with such a business, to produce any controlled drugs which are in his possession or under his control,

(c) require any such person, or any person so employed, to produce any books, records or other documents [(including those containing any data that constitutes personal data)] which relate to transactions concerning controlled drugs and which are in his possession or under his control, and

(d) inspect any controlled drug, book, record or other document produced in pursuance of a requirement under this section.

[(2) For the purposes of enforcing this Act and any statutory instruments made thereunder, and without prejudice to the generality of subsection (1) of this section, a person authorised in writing in that behalf by the Council of the Pharmaceutical Society of Ireland may at all reasonable times—

(a) enter any building or premises in which a person keeps open shop for the dispensing or compounding of medical prescriptions,

(b) require any such person, or any person employed in connection with keeping such open shop for the dispensing or compounding of medical prescriptions, to produce any controlled drugs which are in his possession or under his control,

(c) require any such person, or any person so employed, to produce any books, records or other documents [(including those containing any data that constitutes personal data)] which relate to transactions concerning controlled drugs and which are in his possession or under his control, and

(d) inspect any controlled drug, book, record or other document produced in pursuance of a requirement under this section.

(3) Where the Minister or the Irish Medicines Board authorises a person under subsection (1) of this section, then the Minister or the Irish Medicines Board, as the case may be, shall furnish the person with a warrant of his authorisation.
(4) Where the Pharmaceutical Society of Ireland authorises a person under subsection (2) of this section, then it shall furnish the person with a warrant of his authorisation.

(5) Where—

(a) a person has been authorised by the Minister under subsection (1) of this section at any time before the commencement of this subsection,

(b) the authorisation is still in force immediately before that commencement, and

(c) either—

(i) the person has, before that commencement, been issued with a certificate of his authorisation, or

(ii) the person has not, before that commencement, been issued with a certificate of his authorisation,

then the Minister shall—

(d) in a case falling within paragraph (c)(i) of this subsection, furnish the person with a warrant of his authorisation upon the surrender of his certificate of authorisation,

(e) in a case falling within paragraph (c)(ii) of this subsection, as soon as reasonably practicable after that commencement, furnish the person with a warrant of his authorisation.

(6) Where a person authorised under subsection (1) or (2) of this section—

(a) claims to exercise a power by virtue of that authorisation, and

(b) is required by a person in relation to whom the power is proposed to be exercised, to produce evidence of that authorisation,

then the person so authorised shall not exercise that power until he has produced the warrant of authorisation furnished under this section to the person in relation to whom the power is proposed to be exercised.

(7) A certificate of authorisation referred to in subsection (5)(c)(i) of this section which has not been surrendered as referred to in subsection (5)(d) of this section shall be deemed to be a warrant of authorisation furnished under this section to the person to whom the certificate of authorisation was furnished, and subsection (6) of this section shall be construed accordingly.

(8) In this section—

‘Data Protection Regulation’ means Regulation (EU) 2016/679 of the European Parliament and of the Council of 27 April 201613 on the protection of natural persons with regard to the processing of personal data and on the free movement of such data, and repealing Directive 95/46/EC (General Data Protection Regulation);

‘personal data’ means personal data within the meaning of—

(a) the Data Protection Regulation, or

(b) Part 5 of the Data Protection Act 2018.

Power of arrest. 25.—(1) Where with reasonable cause a member of the Garda Síochána suspects that an offence under section 15 of this Act has been committed and so suspects a person of having committed the offence, he may arrest the person without warrant.

(2) Where with reasonable cause a member of the Garda Síochána,
(a) suspects that an offence under this Act, other than an offence under section 15, has been committed or attempted, and

(b) suspects a person of having committed the offence or having made the attempt, then if the member,

(c) with reasonable cause suspects that the person unless he is arrested either will abscond for the purposes of evading justice or will obstruct the course of justice, or

(d) having enquired of the person, has reasonable doubts as to the person’s identity or place of abode, or

(e) having enquired of the person, knows that the person does not ordinarily reside in the State, or has reasonable doubts as to whether the person so resides,

he may arrest the person without warrant.

Search warrants.  26.—(1) If a Justice of the District Court or a Peace Commissioner is satisfied by information on oath of a member of the Garda Síochána [or if, [subject to the provisions of subsections (2) and (2A) of section 8 of the Criminal Justice (Drug Trafficking) Act 1996], a member of the Garda Síochána not below the rank of superintendent is satisfied] that there is reasonable ground for suspecting that—

(a) a person is in possession in contravention of this Act on any premises of a controlled drug, a forged prescription or a duly issued prescription which has been wrongfully altered and that such drug or prescription is on a particular premises [or other land], or

[(aa) opium poppy, a plant of the genus Cannabis or a plant of the genus Erythroxylon is being cultivated contrary to section 17 of this Act on or in any premises or other land, or]

(b) 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 the State, an offence against a provision of a corresponding law within the meaning of section 20 of this Act and in force in that place, is in the possession of a person on any premises,

[such Justice, Commissioner or, as the case may be, member] may issue a search warrant mentioned in subsection (2) of this section.

[(2) A search warrant issued under this section shall be expressed and operate to authorise a named member of the Garda Síochána, accompanied by such other members of the Garda Síochána and such other persons as may be necessary, at any time or times within one month of the date of issue of the warrant, to enter (if need be by force) the premises or other land named in the warrant, to search such premises or other land and any persons found therein, to examine any substance, article or other thing found thereon or therein, to inspect any book, record or other document found thereon and, if there is reasonable ground for suspecting that an offence is being or has been committed under this Act in relation to a substance, article or other thing found on such premises or other land or that a document so found is a document mentioned in subsection (1) (b) of this section or is a record or other document which the member has reasonable cause to believe to be a document which may be required as evidence in proceedings for an offence under this Act, to seize and detain the substance, article, document or other thing, as the case may be.
(3) Where any premises or other land is entered pursuant to a warrant issued under this section, the member of the Garda Síochána named in the warrant may do either or both of the following:

(a) arrest without warrant any person or persons found on such premises or other land for the purpose of searching him or them,

(b) so arrest any such person or persons and keep him or them, as may be appropriate, under arrest until such time as such of the powers of search or examination as he wishes to exercise pursuant to the warrant have been exercised by him.

(4) In this section—

‘land’ includes any structure on land;

‘structure’ means building, structure or any other thing constructed, erected, placed or made on, in or under any land.

Penalties.

27. — [(1) Subject to section 28 of this Act, every person guilty of an offence under section 3 of this Act shall be liable—

(a) where the relevant controlled drug is cannabis or cannabis resin and the court is satisfied that the person was in possession of such drug for his personal use:

(i) in the case of a first offence,

(I) on summary conviction, to a fine not exceeding £300, or

(II) on conviction on indictment, to a fine not exceeding £500,

(ii) in the case of a second offence,

(I) on summary conviction, to a fine not exceeding £400, or

(II) on conviction on indictment, to a fine not exceeding £1,000,

(iii) in the case of a third or subsequent offence,

(I) on summary conviction, to a fine not exceeding £1,000 or, at the discretion of the court, to imprisonment for a term not exceeding twelve months, or to both the fine and the imprisonment, or

(II) on conviction on indictment, to a fine of such amount as the court considers appropriate or, at the discretion of the court, to imprisonment for a term not exceeding three years, or to both the fine and the imprisonment;

(b) in any other case—

(i) on summary conviction, to a fine not exceeding £1,000 or, at the discretion of the court, to imprisonment for a term not exceeding twelve months, or to both the fine and the imprisonment, or

(ii) on conviction on indictment, to a fine of such amount as the court considers appropriate or, at the discretion of the court, to imprisonment for a term not exceeding seven years, or to both the fine and the imprisonment.

(2) Subject to section 28 of this Act, every person guilty of an offence under section 6, 7, 16, 17, 19 or 20 of this Act shall be liable—
(a) on summary conviction, to a fine not exceeding £1,000 or, at the discretion of the court, to imprisonment for a term not exceeding twelve months, or to both the fine and the imprisonment, or

(b) on conviction on indictment, to a fine of such amount as the court considers appropriate or, at the discretion of the court, to imprisonment for a term not exceeding fourteen years, or to both the fine and the imprisonment.

(3) Subject to section 28 of this Act, every person guilty of an offence under section 15 of this Act shall be liable—

(a) on summary conviction, to a fine not exceeding £1,000 or, at the discretion of the court, to imprisonment for a term not exceeding twelve months, or to both the fine and the imprisonment, or

(b) on conviction on indictment, to a fine of such amount as the court considers appropriate or, at the discretion of the court, to imprisonment for life or such lesser period as the court shall determine, or, at such discretion, to both such fine and such lesser period of imprisonment.

((3A) Every person guilty of an offence under section 15A or 15B of this Act shall be liable, on conviction on indictment—

(a) to imprisonment for life or such shorter term as the court may determine, subject to subsections (3C) and (3D) of this section or, where subsection (3F) of this section applies, to that subsection, and

(b) at the court’s discretion, to a fine of such amount as the court considers appropriate.

(3B) The court, in imposing sentence on a person for an offence under section 15A or 15B of this Act, may, in particular, have regard to whether the person has a previous conviction for a drug trafficking offence.

(3C) Where a person (other than a person under the age of 18 years) is convicted of an offence under section 15A or 15B of this Act, the court shall, in imposing sentence, specify a term of not less than 10 years as the minimum term of imprisonment to be served by the person.

(3D) (a) The purpose of this subsection is to provide that in view of the harm caused to society by drug trafficking, a court, in imposing sentence on a person (other than a person under the age of 18 years) for an offence under section 15A or 15B of this Act, shall specify a term of not less than 10 years as the minimum term of imprisonment to be served by the person, unless the court determines that by reason of exceptional and specific circumstances relating to the offence, or the person convicted of the offence, it would be unjust in all the circumstances to do so.

(b) Subsection (3C) of this section shall not apply where the court is satisfied that there are exceptional and specific circumstances relating to the offence, or the person convicted of the offence, which would make a sentence of not less than 10 years imprisonment unjust in all the circumstances and for that purpose the court may, subject to this subsection, have regard to any matters it considers appropriate, including—

(i) whether that person pleaded guilty to the offence and, if so—

(I) the stage at which he or she indicated the intention to plead guilty, and

(II) the circumstances in which the indication was given, and
whether that person materially assisted in the investigation of the offence.

(c) The court, in considering for the purposes of paragraph (b) of this subsection whether a sentence of not less than 10 years imprisonment is unjust in all the circumstances, may have regard, in particular, to—

(i) whether the person convicted of the offence concerned was previously convicted of a drug trafficking offence, and

(ii) whether the public interest in preventing drug trafficking would be served by the imposition of a lesser sentence.

(3E) Subsections (3C) and (3D) of this section apply and have effect only in relation to a person convicted of a first offence under section 15A or 15B of this Act (other than a person who falls under paragraph (b) of subsection (3F) of this section), and accordingly references in those first-mentioned subsections to an offence under section 15A or 15B of this Act are to be construed as references to a first such offence.

(3F) Where a person (other than a person under the age of 18 years)—

(a) is convicted of a second or subsequent offence under section 15A or 15B of this Act, or

(b) is convicted of a first offence under one of those sections and has been convicted under the other of those sections,

the court shall, in imposing sentence, specify a term of not less than 10 years as the minimum term of imprisonment to be served by the person.

(3G) The power conferred by section 23 of the Criminal Justice Act 1951 to commute or remit a punishment shall not, in the case of a person serving a sentence imposed under subsection (3A) of this section, be exercised before the expiry of the minimum term specified by the court under subsection (3C) or (3F), as may be appropriate, of this section less any reduction of that term under subsection (3H) of this section.

(3H) The rules or practice whereby prisoners generally may earn remission of sentence by industry and good conduct shall apply in the case of a person serving a sentence imposed under subsection (3A) of this section and the minimum term specified by the court under subsection (3C) of this section shall be reduced by the amount of any remission so earned by the person.

(3I) Any powers conferred by rules made under section 2 of the Criminal Justice Act 1960 to release temporarily a person serving a sentence of imprisonment shall not, in the case of a person serving a sentence imposed under subsection (3A) of this section, be exercised during the term for which the commutation or remission of his or her punishment is prohibited by subsection (3G) of this section unless for a grave reason of a humanitarian nature, and any release so granted shall be only of such limited duration as is justified by such reason.

(3J) In imposing a sentence on a person convicted of an offence under section 15A or 15B of this Act, a court—

(a) may inquire whether at the time of the commission of the offence the person was addicted to one or more controlled drugs, and

(b) if satisfied that the person was so addicted at that time and that the addiction was a substantial factor leading to the commission of the offence, may list the sentence for review after the expiry of not less than one-half of the term specified by the court under subsection (3C) or (3F), as may be appropriate, of this section.

(3K) On reviewing a sentence listed under subsection (3J)(b) of this section, the court—
(a) may suspend the remainder of the sentence on any conditions it considers fit, and
(b) in deciding whether to exercise its powers under this subsection, may have regard to any matters it considers appropriate.

(3L) Paragraph (a) of section 13(2) of the Criminal Procedure Act 1967 shall not apply in relation to an offence under section 15A or 15B of this Act, but each of those offences shall be deemed for the purposes of paragraph (b) of section 13(2) of that Act to be an offence to which section 13 of that Act applies.

(3M) The reference in subsection (3I) of this section to section 2 of the Criminal Justice Act 1960 shall be construed to include that section as applied by section 4 of the Prisons Act 1970.

(3N) In subsections (3B) and (3D) of this section ‘drug trafficking offence’ has the meaning it has in section 3(1) of the Criminal Justice Act 1994 and in subsection (3D) of this section ‘drug trafficking’ has the meaning it has in the said section 3(1).

(4) Subject to section 28 of this Act, every person guilty of an offence under section 18 of this Act shall be liable—

(a) on summary conviction, to a fine not exceeding £400 or, at the discretion of the court, to imprisonment for a term not exceeding six months, or to both the fine and the imprisonment, or

(b) on conviction on indictment, to a fine of such amount as the court considers appropriate or, at the discretion of the court, to imprisonment for a term not exceeding three years, or to both the fine and the imprisonment.

(5) Every person guilty of an offence under section 21 (1) of this Act shall be liable to be punished on summary conviction as if he were guilty of the substantive offence and in case a penalty on conviction on indictment is provided by this Act in relation to the substantive offence, he shall be liable to be proceeded against on indictment and, if convicted, punished as if he were convicted on indictment of the substantive offence.

(6) Every person guilty of an offence under section 21 (2) of this Act shall be liable—

(a) in case the regulation in relation to which the offence was committed is a regulation made pursuant to section 5 (1) (a) of this Act, other than a regulation regulating the transportation of controlled drugs,

(i) on summary conviction, to a fine not exceeding £1,000 or, at the discretion of the court, to imprisonment for a term not exceeding twelve months, or to both the fine and the imprisonment, or

(ii) on conviction on indictment, to a fine of such amount as the court considers appropriate or, at the discretion of the court, to imprisonment for a term not exceeding fourteen years, or to both the fine and the imprisonment, and

(b) in case the regulation in relation to which the offence was committed is a regulation made otherwise than under the said section 5 (1) (a) or is a regulation regulating the transportation of controlled drugs—

(i) on summary conviction, to a fine not exceeding £500 or, at the discretion of the court, to imprisonment for a term not exceeding six months, or to both the fine and the imprisonment, or

(ii) on conviction on indictment, to a fine of such amount as the court considers appropriate, or at the discretion of the court, to imprisonment for a term not exceeding two years, or to both the fine and the imprisonment.
(7) Every person guilty of an offence under section 21 of this Act, other than an offence mentioned in subsection (1) or subsection (2) of that section, shall be liable on summary conviction to a fine not exceeding £400 or, at the discretion of the court, to imprisonment for a term not exceeding six months, or to both the fine and the imprisonment.

(8) Every person guilty of an offence under paragraph (a) or (b) of subsection (1D) of section 23 of this Act, as amended by section 12 of the Misuse of Drugs Act, 1984, shall be liable on summary conviction to a fine not exceeding £200.

(9) Every person guilty of an offence under section 5 of the Misuse of Drugs Act, 1984, shall on summary conviction be liable—

(a) in case the offence is an offence under subsection (2) of that section, to a fine not exceeding £1,000,

(b) in any other case, to a fine not exceeding £500.

(10) […]

(11) […]

(12) In this section—

“relevant controlled drug” means the controlled drug in relation to which the offence was committed;

“substantive offence” means the offence under this Act to which the attempt or, as the case may be, the aiding, abetting, counselling, procuring, soliciting or incitement was directed.

Power of court to remand persons convicted under section 3, 15, 16, 17 or 18 and to obtain a report and in certain cases to arrange for the medical treatment or for the care of such persons.

28.—(1) (a) Where a person is convicted of an offence under section 3 of this Act, other than a first or second offence in relation to which a penalty may be imposed under section 27 (1) (a) of this Act, or an offence under section 15 or 16 of this Act, or of attempting to commit any such offence, [if, having regard to the circumstances of the case, the court considers it appropriate so to do, the court may] remand the person for such period as it considers necessary for the purposes of this section (being a period not exceeding eight days in the case of a remand in custody), and request [the Health Service Executive], [probation and welfare] officer or other body or person, considered by the court to be appropriate, to—

(i) cause to be furnished to the court a medical report in writing on the convicted person together with such recommendations (if any) as to medical treatment which the person making the report considers appropriate to the needs [arising because of his being dependent on drugs.] of the convicted person, and

(ii) furnish to the court a report in writing as to the vocational and educational circumstances and social background of the convicted person together with such recommendations (if any) as to care which the body or person making the report considers appropriate to the said needs.

(b) Where a person is convicted of a first or second offence under section 3 of this Act in relation to which a penalty may be imposed under the said section 27 (1) (a) or an offence under section 17 or 18 of this Act, or of attempting to commit any such offence, and the court, having regard to the circumstances of the case, considers it appropriate so to do, the court may remand the person [on bail or, unless a penalty falls to be imposed on the person under paragraph (a) of section 27 (1) of this Act, in custody] for such period as it considers necessary for the purposes of this section, and request [the Health Service Executive], [probation and welfare] officer or other body or person, considered by the court to be appropriate, to—
(i) cause to be furnished to the court a medical report in writing on the convicted person together with such recommendations (if any) as to medical treatment which the person making the report considers appropriate to the needs [arising because of his being dependent on drugs] of the convicted person, and

(ii) furnish to the court a report in writing as to the vocational and educational circumstances and social background of the convicted person together with such recommendations (if any) as to care which the body or person making the report considers appropriate to the said needs.

(2) Having considered the reports furnished pursuant to subsection (1) of this section, the court shall, if in its opinion the welfare of the convicted person warrants its so doing, instead of imposing a penalty under section 27 of this Act, but subject to subsection (8) of this section either—

(a) permit the person concerned to enter into a recognisance containing such of the following conditions as the court considers appropriate having regard to the circumstances of the case and the welfare of the person, namely—

(i) a condition that the person concerned be placed under the supervision of such body (including [the Health Service Executive]) or person as may be named in the order and during a period specified in the order,

(ii) in case the person concerned is placed under such supervision, a condition requiring such person, at the place at which he normally resides or at such other place as may be specified in the order and during such period and at such intervals as shall be so specified, to receive visits from and permit visits by—

(I) in case such person is placed under the supervision of a body, an officer of that body,

(II) in case such person is placed under the supervision of a person, that person,

(iii) a condition requiring such person to undergo medical [or other] treatment recommended in the report,

(iv) a condition requiring such person for such treatment to attend or remain in a hospital, clinic or other place specified in the order for a period so specified,

(b) order that the person be detained in custody in a designated custodial treatment centre for a period not exceeding the maximum period of imprisonment which the court may impose in respect of the offence to which the conviction relates, or one year, whichever is the shorter.

(3) A court may, if it thinks fit, consider otherwise than in public—

(a) a report under subsection (1) of this section,

(b) whether or not it will permit a person to enter into a recognisance mentioned in subsection (2) of this section, or

(c) whether or not it will make an order referred to in paragraph (b) of subsection (2) of this section.

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(4) In any proceedings in which a report furnished under subsection (1) of this section is considered, the court may, if it believes that it is in the interests of the person concerned not to know the contents of the report, withhold from him the report, but the foregoing shall not be construed as preventing any barrister or solicitor who appears on such person's behalf in the proceedings seeing the report or, if thought fit, questioning or commenting on any of its contents in the proceedings.

(5) Where it is alleged to the court that a person has been in breach of a recognisance entered into by him under subsection (2) of this section, the court, notwithstanding the decision by it under the said subsection (2), may direct that the person be brought before the court, and, if satisfied that the person has been in breach of the recognisance, may estreat the recognisance and, subject to subsection (8) of this section, either make in respect of the person an order referred to in paragraph (b) of subsection (2) of this section or proceed to deal with the case in accordance with the provisions of section 27 of this Act as if the decision had not been made.

(6) If at any time during a period of detention in a designated custodial treatment centre it appears to the court, on an application made by or on behalf either of the prosecutor or the person who is being detained, or on receipt of a message, in a form approved of by the Minister, from an authorised medical practitioner [or the person who is for the time being in charge of such centre], that the person being detained under this Act is not then, or may not then be, in further need of the treatment or care of which the court formerly considered him to be in need, or that his continued detention in custody in the designated custodial treatment centre is not then, or may not then be, in his best interests or in the best interests of other persons in that centre, the court, notwithstanding its decision under subsection (2) of this section, may order the person to be brought before the court.

(7) Where a person is brought before the court pursuant to an order under subsection (6) of this section, the court may inquire into the case and hear such evidence as it considers relevant, and if, having considered the circumstances of the case, the court is satisfied that the person is not then in further need of the treatment or care referred to in subsection (6) of this section, or that his continued detention in custody in the designated custodial treatment centre would not be in his best interests or in the best interests of other persons in that centre, the court, notwithstanding its decision under subsection (2) of this section, may revoke the relevant order made by it under the said subsection (2) and, subject to subsection (8) of this section,

(a) permit the person to enter into a recognisance described in the said subsection (2) if the court is of the opinion that the welfare of the person warrants its so doing, or

(b) order the person to be detained for a period not exceeding the unexpired portion of the period specified in the revoked order in a custodial treatment centre other than that so specified, or

(c) decide not to impose any penalty under section 27 of this Act, or

(d) where it considers it appropriate so to do and subject to subsection (9) of this section, proceed to deal with the offence in accordance with section 27 of this Act.

(8) The court shall not under this section either,

(a) permit a person to enter into a recognisance containing a condition requiring him for [medical or other treatment] to remain in a specified hospital, clinic or other place, or

(b) order a person to be detained in a custodial treatment centre,

unless, after consultation with, or consideration of a report of, either the [the medical practitioner or other person] in charge of the hospital, clinic, custodial treatment centre or other place concerned or a medical practitioner nominated by [the medical practitioner or other person] so in charge, the court is satisfied that the giving or
making of the permission or order would be an appropriate course having regard to
the needs of the person and would not prejudicially affect the ability of such hospital,
clinic, custodial treatment centre or other place to provide for the treatment or care
of persons.

(9) In case a court decides, pursuant to subsection (7) of this section, to impose a
sentence of imprisonment under section 27 of this Act, the period of imprisonment
which may be so imposed shall not exceed the period by which the maximum term
of imprisonment which that court could otherwise have imposed under the said section
27 for the offence of which the person was convicted exceeds the period already
spent by him in custody on foot of the order revoked by the court.

(10) The Minister may by order designate an institution which in his opinion is
suitable for the medical treatment or the care of persons in respect of whom an order
may be made under this section, or a specified part of such an institution, as a
designated custodial treatment centre for the purpose of this section.

(11) In this section—
“authorised medical practitioner” means a registered medical practitioner authorised
for the purposes of this section by the Minister in writing or a registered medical
practitioner of a class specified by the Minister as being authorised for the said
purposes;

[‘probation and welfare officer’ means an officer employed in the probation and
welfare service of the Department of Justice.]

[...]
and that as soon as practicable he took all such steps as were reasonably open to him to destroy the drug or document or to deliver it into the custody of such a person.

(3) In any proceedings for an offence under section 15 or 15A, or subsection (1)(d) of section 15C, of this Act, a defendant may rebut the presumption raised by subsection (2) of the said section 15 or 15A or subsection (3) of the said section 15C, as the case may be, by showing that at the time of the alleged offence, he or she was by virtue of regulations made under section 4 of this Act lawfully in possession of the controlled drug or drugs to which the proceedings relate.

(4) In any proceedings for an offence under section 19 of this Act it shall be a defence to show that the defendant took steps to prevent the occurrence or continuance of the activity or contravention to which the alleged offence relates and that, in the particular circumstances, the steps were taken as soon as practicable and were reasonable.

(5) In any proceedings for an offence under section 16, 17 or 21 (2) of this Act, it shall be a defence for the defendant to prove that he neither knew of nor suspected nor had reason to suspect the existence of some fact alleged by the prosecutor which it is necessary for the prosecutor to prove if he is to be convicted of the offence charged.

(6) In any proceedings for an attempt to commit an offence under this Act the defences mentioned in subsection (2) or (5) of this section shall, with the necessary modifications, be open to the defendant.

(7) Subject to subsection (1) of this section nothing in this section shall prevent a person raising a defence which, apart from this section, would be open to him to raise in proceedings for an offence under this Act.

Forfeiture.

30.—[(1) Subject to subsection (2) of this section, a court by which a person is convicted of an offence under this Act or a drug trafficking offence (within the meaning of the Criminal Justice Act, 1994), may order anything shown to the satisfaction of the court to relate to the offence to be forfeited and either destroyed or dealt with in such other manner as the court thinks fit.]

(2) A court shall not order anything to be forfeited under this section if a person claiming to be the owner of or otherwise interested in it applies to be heard by the court, unless an opportunity has been given to him to show cause why the order should not be made.

Offences in relation to bodies corporate.

31.—Where an offence under this Act is committed by a body corporate or by a person purporting to act on behalf of a body corporate and is proved to have been so committed with the consent, connivance or approval of, or to have been facilitated by any neglect on the part of, any director, manager, secretary or other official of such body, such person shall also be guilty of the offence.

Poisons for purposes of Pharmacy Acts, 1875 to 1962.

32.—[…]

Amendment of Poisons Act, 1961.

33.—(1) The Poisons Act, 1961, shall be amended as follows:

(a) section 4 (1) is hereby amended by—

(i) the addition of “and one of whom is a person with knowledge and experience of the manufacture of preparations containing poisons” to paragraph (b), and
(ii) the insertion of the following paragraph after paragraph (c),

“(cc) one person who is a fellow, ordinary member or licentiate of the Institute of Chemistry of Ireland,”;

(b) section 14 (3) is hereby amended by the substitution of the following paragraphs for paragraphs (j) to (l):

“(j) provide for the enforcement and execution of the provisions of the regulations—

(i) by officers of the Minister,

(ii) with the consent of the Minister for Agriculture, by officers of that Minister,

(iii) by the Pharmaceutical Society of Ireland and its officers, and

(iv) by health boards and their officers,

(k) enable any such officer (with, in the case of an officer of the Minister or the Minister for Agriculture, a written authorisation of whichever of those Ministers is appropriate, in the case of an officer of the Pharmaceutical Society of Ireland, a written authorisation of that Society, and in the case of an officer of a health board, a written authorisation of the board), at all reasonable times, for the purpose of ascertaining whether or not there is or has been a contravention of the regulations, to enter premises of a class or description specified in the regulations and to inspect any substance or article which is so specified and require the production of and inspect, and if he thinks fit take copies of any entry in, any book, record or other document which is of a class or description so specified,

(l) provide for the taking (without payment) by such officers, with such authorisation, of samples of poisons or such substances for test, examination or analysis,

(m) prescribe the certificate or other evidence to be given of the result of any such test, examination or analysis and the classes of person by whom such certificate or evidence is to be given,

(n) provide that any certificate or other evidence specified under paragraph (m) of this subsection and given in respect of the test, examination or analysis of a sample shall with regard to that sample be evidence for all purposes of the result,

(o) provide for the prosecution of offences under section 17 of this Act in relation to the regulations by the Minister, the Pharmaceutical Society of Ireland or health boards, and

(p) provide for matters ancillary to the foregoing matters.”;

(c) the following new section is hereby inserted after section 15:

“Evidence of result of certain tests, examinations or analyses.

15A.—Whenever regulations made under this Act provide that a certificate or other evidence is to be evidence for all purposes of a result of a test, examination or analysis of a sample, such certificate or other evidence shall until the contrary is shown, in relation to that sample, be accepted by a court as sufficient evidence of the result of the test, examination or analysis.”.

(2) Regulations under section 14 or section 15 of the Poisons Act, 1961, may provide that a substance which is declared to be a poison for the purposes of those regulations or a provision thereof, shall, for the purposes of whichever of the said sections is
appropriate and the regulations or provision to which the declaration relates, be regarded in circumstances specified in the regulations as not being the subject of the declaration.

Amendment of section 2 of Pharmacy Act, 1962.

34.—Section 2 of the Pharmacy Act, 1962, is hereby amended by the substitution of the following subsection for subsection (4):

“(4) It shall not be a contravention of subsection (2) of this section for a person to keep open shop for the sale of a substance which is declared to be a poison for the purposes of regulations made under section 14 of the Poisons Act, 1961, if the person is a person, or a member of a class of persons, by whom pursuant to such regulations the substance may be sold or offered or kept for sale.”.

Amendment of Pharmacopoeia Act, 1931.

35.—(1) The Pharmacopoeia Act, 1931, is hereby amended as follows:

(a) section 2 (1) shall be construed and have effect as if the reference therein to the Council were a reference to the Minister for Health;

(b) the following subsection shall be substituted for subsection (2) of section 2:

“(2) The Minister for Health may by regulations make such modifications (by way of deletion, addition or amendment) in the British Pharmacopoeia for the time being in force in Great Britain as he thinks fit.”;

(c) the following subsection shall be substituted for subsection (2) of section 3:

“(2) A certificate endorsed on a book purporting to be a copy of the British Pharmacopoeia and purporting to be signed by an officer of the Minister for Health that such book is a copy of the British Pharmacopoeia in force in Great Britain on a specified day or days, or during a specified period, shall in any legal proceedings until the contrary is shown be admitted as evidence of the facts so certified, and in such proceedings it shall not be necessary to prove the signature of the person purporting to sign the certificate or that the person was an officer of the said Minister.”; and

(d) the following new section shall be inserted after section 4:

“Regulations.

4A.—Every regulation made under this Act shall be laid before each House of the Oireachtas as soon as may be after it is made and, if a resolution annulling the regulation is passed by either such House within the next twenty-one days on which that House has sat after the regulation is laid before it, the regulation shall be annulled accordingly, but without prejudice to anything previously done thereunder.”.

(2) As on and from the specified day section 4 of the Pharmacopoeia Act, 1931, shall be construed and have effect as if—

(a) “or the European Pharmacopoeia” were inserted after “Saorstát Éireann Pharmacopoeia” in subsection (1), and

(b) the following subsection were added to the section:

“(3) In this section ‘the European Pharmacopoeia’ means the Pharmacopoeia elaborated under the auspices of the Council of Europe in pursuance of the Convention in that behalf done at Strasbourg on the 22nd day of July, 1964.”.

(3) In this section “the specified day” means the day specified for the purposes of this section in a notice published by the Minister in the Iris Oifigiúil.
36.—Section 65 of the Health Act, 1947, as amended by section 39 of the Health Act, 1953, and section 6 of the Health Act, 1970, is hereby amended by—

(a) the substitution of the following paragraphs for paragraph (j) of subsection (3):

"(j) the enforcement and execution of the regulations—

(i) by officers of the Minister,

(ii) by health boards and their officers,

(iii) with the consent of the Minister for Finance, by officers of Customs and Excise,

(iv) with the consent of the Minister for Industry and Commerce, by officers of that Minister,

(v) by the Pharmaceutical Society of Ireland and its officers;

(k) the enabling for the purpose of ascertaining whether or not there is or has been a contravention of the regulations, of any such officer (with, in the case of an officer of the Minister or the Minister for Industry and Commerce or an officer of Customs and Excise, a written authorisation of whichever of those Ministers or the Minister for Finance is appropriate, in the case of an officer of a health board, a written authorisation of the board and in the case of an officer of the Pharmaceutical Society of Ireland, a written authorisation of that Society), at all reasonable times to enter any premises which are of a class or description specified in the regulations and to inspect or examine any substance or article which is of a class or description so specified and require the production of and inspect, and if he thinks fit, to take copies of any entry in, any book, record or other document which is of a prescribed class or description;

(l) the taking (without payment) by such officers, with such authorisation, of samples of such substances or articles for test, examination or analysis;

(m) the prescribing of the certificate or other evidence to be given of the result of any such test, examination or analysis and the classes of person by whom such certificate or evidence is to be given; and

(n) providing that any certificate or other evidence specified under paragraph (m) of this subsection and given in respect of the test, examination or analysis of a sample shall with regard to that sample be evidence for all purposes of the result."

(b) the substitution of the following subsection for subsection (5):

"(5) An offence under this section may be prosecuted by the Minister, the Pharmaceutical Society of Ireland or by the health board in whose functional area the offence is committed."

37.—(1) Where a notice or other document is required or authorised by this Act or by regulations under this Act to be served on or given or sent to a person, it may be served on or given or sent to him—

(a) by delivering it to him,

(b) in the case of a person other than a body corporate, by sending it by post in an envelope addressed to him at the address at which he ordinarily resides or carries on business, or
(c) in the case of a body corporate, by sending it by post in an envelope addressed to the secretary or principal officer of the body at the address at which the body carries on business.

(2) For the purposes of subsection (1) of this section, a company registered under the Companies Act, 1963, shall be deemed to carry on business at its registered offices and every other body corporate and every unincorporated body of persons shall be deemed to carry on business at its principal office or place of business.

38.—(1) The Minister may make regulations for prescribing any matter referred to in this Act as prescribed, provided that in so far as any such regulations provide for the charging of fees they shall only be made with the consent of the Minister for Finance.

(2) Regulations under this Act may apply to controlled drugs generally, to controlled drugs of a prescribed class or description, or to one or more prescribed controlled drugs.

(3) Every regulation and every order made under this Act (other than an order under section 8 (8) or an order referred to in section 11 or section 28) shall be laid before each House of the Oireachtas as soon as may be after it is made and, if a resolution annulling the regulation or order is passed by either such House within the next twenty-one days on which that House has sat after the regulation or order is laid before it, the regulation or order, as the case may be, shall be annulled accordingly but without prejudice to the validity of anything previously done thereunder.

39.—The expenses incurred by the Minister in the administration of this Act shall, to such extent as may be sanctioned by the Minister for Finance, be paid out of moneys provided by the Oireachtas.

40.—(1) All moneys payable under regulations under this Act shall be collected and taken in such manner as the Minister for Finance may from time to time direct and shall be paid into or disposed of for the benefit of the Exchequer in accordance with the directions of the Minister for Finance.

(2) The Public Offices Fees Act, 1879, shall not apply in respect of moneys mentioned in subsection (1) of this section and payable to the Minister.

41.—(1) The Dangerous Drugs Act, 1934, is hereby repealed.

(2) In case a provision of this Act other than subsection (1) of this section, comes into force on a day which is earlier than the day on which the said subsection (1) comes into force, the following provisions shall have effect, namely, as regards the period beginning on the day on which the first-mentioned provision comes into force and ending on the day on which the said subsection (1) comes into force, an act or omission which is an offence under this Act shall not be an offence under the Dangerous Drugs Act, 1934.

42.—(1) The following are hereby repealed:

(a) (i) “by the General Council of Medical Education and Registration of the United Kingdom” in section 1 of the Pharmacopoeia Act, 1931, and

(ii) section 2 (3) of that Act; and

(b) section 78 of the Health Act, 1970.

(2) Notwithstanding subsection (1) of this section, the Minister may by regulations provide—
(a) that a register specified in the regulations and kept by him under regulations made under section 78 of the Health Act, 1970, shall be included in and shall be deemed to be part of a register to be kept by him for the purposes of regulations under this Act, or

(b) that any person, being a person whose name was, immediately before the commencement of this section, on a register specified in the regulations and kept by the Minister under the said section 78, shall be deemed to have been granted, issued or given, as may be appropriate, such licence, permit or other form of authority under this Act as may be specified in the regulations.

(3) Notwithstanding section 41 (1) of this Act, the Minister may by regulations provide that any person who was, immediately before the commencement of this section, the holder of a licence, permit or other authority granted, issued or given under the Dangerous Drugs Act, 1934, shall be deemed to have been granted, issued or given, as may be appropriate, such licence, permit or other form of authority under this Act as may be specified in the regulations.

(4) Regulations made under this section which include provisions mentioned in paragraph (b) of subsection (2) or in subsection (3) of this section may also include—

(a) provisions deeming any such licence, permit or other form of authority to have been granted, issued or given subject to conditions specified in the regulations, and

(b) provisions enabling the Minister in specified circumstances to direct that provisions of regulations under this section shall cease to apply in relation to a particular person.

(5) Where the Minister duly gives a direction referred to in subsection (4) (b) of this section, the provisions specified in the direction shall in accordance with the direction cease to apply in relation to the person to whom the direction relates.

43.—(1) This Act may be cited as the Misuse of Drugs Act, 1977.

(2) Subsection (1) of this section and section 41 (2) of this Act shall come into operation on the passing hereof and the other purposes and provisions of this Act shall come into operation on such day or days as may be fixed therefor by any order or orders of the Minister, either generally or with reference to any particular such purpose or provision and different days may be so fixed for different such purposes and different such provisions of this Act.

(3) The Pharmacy Acts, 1875 to 1962, and sections 32 and 34 of this Act may be cited together as the Pharmacy Acts, 1875 to 1977.

(4) The Pharmacopoeia Act, 1931, section 35 of this Act, and section 42 of this Act in so far as it amends that Act, may be cited together as the Pharmacopoeia Acts, 1931 and 1977.

(5) The Health Acts, 1947 to 1970, section 36 of this Act, and section 42 of this Act in so far as it amends those Acts, may be cited together as the Health Acts, 1947 to 1977.

SCHEDULE

Controlled Drugs

1 Acetorphine.
Acetyldihydrocodeine.
Acetylmethadol.
Allylprodine.
Alphacetylmethadol.
Alphameprodine.
Alphamethadol.
Alphaprodine.
Amphetamine.
Amylobarbitone.
Anileridine.
Benzethidine.
Benzphetamine.
Benzylmorphine (3-benzylmorphine).
Betacetylmethadol.
Betameprodine.
Betamethadol.
Betaprodine.
Bezitramide.
Bufotenine.
Cannabinol, except where contained in cannabis or cannabis resin.
Cannabinol derivatives.
Cannabis and cannabis resin.
Chlorphentermine.
Clonitazene.
Coca leaf.
Cocaine.
Codeine.
Codoxime.
Desomorphine.
Dexamphetamine.
Dextromoramide.
Diamorphine.
Diamppromide.
Diethylthiambutene.
Difenoxin.
Dihydrocodeine.
Dihydromorphine.
Dimenoxadole.
Dimepheptanol.
Dimethylthiambutene.
Dioxaphetyl butyrate.
Diphenoxylate.
Dipipanone.
Drotebanol.
Ecgonine, and any derivative of ecgonine which is convertible to ecgonine or to cocaine.
Ethylmethyliambutene.
Ethylmorphine (3-ethylmorphine).
Etonitazene.
Etorphine.
Etoxeridine.
Fentanyl.
Furethidine.
Hydrocodone.
Hydromorphinol.
Hydromorphone.
Hydroxypethidine.
Isomethadone.
Ketobemidone.
Levomethorphan.
Levomoramide.
Levophenacylmorphan.
Levorphanol.
Lysergamide.
Lysergide and other N-alkyl derivatives of lysergamide.
Mephentermine.
Mescaline.
Metazocine.
Methadone.
Methaqualone.
Methylamphetamine.
Methyldesorphine.
Methyldihydromorphine (6-methyldihydromorphine).
Methylphenidate.
Metopon.
Morpheridine.
Morphine.
Morphine methobromide, morphine N-oxide and other pentavalent nitrogen morphine derivatives.
Myrophine.
Nicocodine.
Nicodicodine (6-nicotinoyldihydrocodeine).
Nicomorphine.
Noracymethadol.
Norcodeine.
Norlevorphanol.
Normethadone.
Normorphine.
Norpipanone.
Opium, whether raw, prepared or medicinal.
Oxycodone.
Oxymorphone.
Pentobarbitone.
Pethidine.
Phenadoxone.
Phenampramide.
Phenazocine.
Phendimetrazine.
Phenmetrazine.
Phenobarbitone.
Phenomorphan.
Phenoperidine.
Pholcodine.
Piminodine.
Pipradrol.
Piritramide.
Poppy straw and concentrate of poppy straw.
Proheptazine.
Properidine.
Propiram.
Psilocin.
Quinalbarbitone.
Racemethorphan.
Racemoramide.
Racemorphan.
Thebacon.
Thebaine.
Trimeperidine.
4-Cyano-2-dimethylamino-4, 4-diphenylbutane.
4-Cyano-1-methyl-4-phenylpiperidine.
N, N-Diethyltryptamine.
N, N-Dimethyltryptamine.
2, 5-Dimethoxy-α, 4-dimethyl-phenethylamine.
1-Methyl-4-phenylpiperidine-4-carboxylic acid.
2-Methyl-3-morpholino-1, 1-diphenylpropanecarboxylic acid.
4-Phenylpiperidine-4-carboxylic acid ethyl ester.

[[1A. (a) N-(Adamantan-1-yl)-1-(5-fluoropentyl)-1H-indazole-3-carboxamide
Alfentanil
(3-Amino-2,2-dimethylpropyl)-4-aminobenzoate
5-(2-Aminopropyl) indole
1-(1,3-Benzodioxol-5-yl)-2-(1-pyrrolidinyl)-1-pentanone
N-(1-Benzyl-4-piperidyl) propionanilide
2-(4-Bromo-2,5-dimethoxyphenyl)-N-[(2-methoxyphenyl)methyl]ethanamine
1-(4-Bromofuro[2,3-f][1]benzofuran-8-yl)propan-2-amine
Carfentanil
Cathinone
2-(4-Chloro-2,5-dimethoxyphenyl)-N-[(2-methoxyphenyl)methyl]ethanamine
1-Cyclohexyl-4-(1,2-diphenylethyl) piperazine
3,4-Dichloro-N-[(1-dimethylamino) cyclohexyl]methyl]benzamide
Dihydroetorphine
[2,3-Dihydro-5-methyl-3-(4-morpholinylmethyl) pyrrolo[1,2,3-de]-1,4-benzoazin-6-yl]-1-naphthalenylmethanone
Dimethocaine
3-Dimethylheptyl-11-hydroxyhexahydrocannabinol
Eticyclidine
Etryptamine
1-(2-Fluorophenyl)-2-methylaminopropan-1-one
1-(3-Fluorophenyl)-2-methylaminopropan-1-one
1-(4-Fluorophenyl)-2-methylaminopropan-1-one
9-(Hydroxymethyl)-6,6-dimethyl-3-(2-methyloctan-2-yl)-6a,7,10,10a-tetrahydrobenzo[c]chromen-1-ol
[9-Hydroxy-6-methyl-3-[5-phenylpentan-2-yl]oxy-5,6,6a,7,8,9,10,10a-octahydrophenanthridin-1-yl] acetate
N-Hydroxy-tenamphetamine
2-(4-Iodo-2,5-dimethoxyphenyl)-N-[(2-methoxyphenyl) methyl]ethanamine
Khat (being the leaves of Catha edulis (Celastraceae))
Lisdexamfetamine
Lofentanil
Methcathinone
2-(3-Methoxyphenyl)-2-(ethylamino) cyclohexanone
1-(4-Methoxyphenyl)-2-(methylamino) propan-1-one
Methyl(2S,4aR,6aR,7R,10aS,10bR)-9-acetyloxy-2-(furan-3-yl)-6a,10b-dimethyl-4,10-dioxo-2,4a,5,6,7,8,9,10a-octahydro-1H-benzo[f]isochromene-7-carboxylate and any product, whether natural or otherwise, including any plant or plant material of any kind or description, which contains any proportion of the said substance
2-Methylamino-1-(3,4-methylenedioxyphenyl) butan-1-one
2-Methylamino-1-(3,4-methylenedioxyphenyl) propan-1-one
4-Methyl-aminorex
(8-Methyl-8-azabicyclo[3.2.1]octan-3-yl)-4-fluorobenzoate
Methyl 2-[[1-(cyclohexymethyl) indole-3-carbonyl]amino]-3,3-dimethylbutanoate
Methyl(E)-2-[(2S,3S,7aS,12bS)-3-ethyl-7a-hydroxy-8-methoxy-2,3,4,6,7,12b-hexahydro-1H-indolo[2,3-a]quinolizin-2-yl]-3-methoxyprop-2-enoate and any product, whether natural or otherwise, including any plant or plant material of any kind or description, which contains any proportion of the said substance
Methyl(E)-2-[(2S,3S,12bS)-3-ethyl-8-methoxy-1,2,3,4,6,7,12,12b-octahydroindolo[2,3-a]quinolizin-2-yl]-3-methoxyprop-2-enoate and any product, whether natural or otherwise, including any plant or plant material of any kind or description, which contains any proportion of the said substance
4-methyl-5-(4-methylphenyl)-4,5-dihydro-1,3-oxazol-2-amine
α-Methyl-4-(methylthio) phenethylamine
1-(4-Methylphenyl)-2-methylaminopropan-1-one
Nabilone
Oripavine
Phencyclidine
1-Phenylcyclohexylamine
4-(1-Phenylcyclohexyl) morpholine
1-Piperidinocyclohexanecarbonitrile
Quinolin-8-yl 1-(5-fluoropentyl)-1H-indole-3-carboxylate
Remifentanil
Rolicyclidine
Sufentanil
Tapentadol
Tenocyclidine
N-[1-(2-Thenyl)-4-piperidyl]propionanilide
4-[1-(2-Thienyl) cyclohexyl]morpholine
1-[1-(2-Thienyl) cyclohexyl]pyrrolidine
Tilidine.

(b) Any substance (not being bupropion) structurally derived from 2-amino-1-phenyl-1-propanone by modification in any of the following ways:

(i) by substitution in the phenyl ring to any extent with alkyl, alkenyl, alkynyl, alkoxy, alkylthio, alkylenedioxy, haloalkyl or halo substituents, whether or not further substituted in the phenyl ring by one or more other univalent substituents;

(ii) by substitution at the 2- or 3-position of the propanone side-chain with an alkyl substituent;

(iii) by substitution at the nitrogen atom with one or more alkyl or dialkyl groups, or by inclusion of the nitrogen atom in a cyclic structure.

(c) Any substance structurally derived from 2-amino-1-propanone by substitution at the 1-position with any monocyclic, or fused-polycyclic ring system (not being a phenyl ring or alkylenedioxyphenyl ring system), whether or not the substance is further modified in any of the following ways:

(i) by substitution in the ring system to any extent with alkyl, alkenyl, alkynyl, alkoxy, alkylthio, haloalkyl or halo substituents, whether or not further substituted in the ring system by one or more other univalent substituents;

(ii) by substitution at the 3-position with an alkyl substituent;

(iii) by substitution at the 2-amino nitrogen atom with one or more alkyl or dialkyl groups, or by inclusion of the 2-amino nitrogen atom in a cyclic structure.

(d) Any substance structurally derived from 3-(1-benzylo)indole or 3-(1-naphthoyl)indole by modification in any of the following ways:

(i) by substitution at the nitrogen atom of the indole ring by alkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl or 2-(4-morpholinyl)ethyl;

(ii) by replacement of one or more hydrogen atoms of any of the substituents referred to in clause (i), with a halo substituent;

whether or not further substituted in the indole ring to any extent and whether or not substituted in the phenyl or naphthyl ring to any extent.

(e) 1-Benzylpiperazine or any substance structurally derived from 1-benzylpiperazine or 1-phenylpiperazine by modification in any of the following ways:

(i) by substitution at the second nitrogen atom of the piperazine ring with alkyl, benzyl, haloalkyl or phenyl groups;

(ii) by substitution in the aromatic ring to any extent with alkyl, alkoxy, alkylenedioxy, halide or haloalkyl groups.
(f) Any substance structurally derived from fentanyl by modification in one or more of the following ways, that is to say:

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

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

(iii) by substitution in the piperidine ring with alkyl or alkenyl groups;

(iv) by substitution in the aniline ring with alkyl, alkoxy, alkylenedioxy, halogeno or haloalkyl groups;

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

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

(g) Any substance structurally derived from 2-(3-hydroxycyclohexyl)phenol by substitution at the 5-position of the phenolic ring by alkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl or 2-(4-morpholino)ethyl, whether or not further substituted in the cyclohexyl ring to any extent.

(h) Any substance structurally derived from 3-(1-naphthoyl)indole or 1H-indol-3-yl-(1-naphthyl) methane by substitution at the nitrogen atom of the indole ring by alkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl or 2-(4-morpholino)ethyl, whether or not further substituted in the indole ring to any extent and whether or not substituted in the naphthyl ring to any extent.

(i) Any substance structurally derived from 3-(1-naphthoyl)pyrrole by substitution at the nitrogen atom of the pyrrole ring by alkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl or 2-(4-morpholino)ethyl, whether or not further substituted in the pyrrole ring to any extent and whether or not substituted in the naphthyl ring to any extent.

(j) Any substance structurally derived from 1-(1-naphthylmethyl)indene by substitution at the 3-position of the indene ring by alkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl or 2-(4-morpholino)ethyl, whether or not further substituted in the indene ring to any extent and whether or not substituted in the naphthyl ring to any extent.

(k) Any substance structurally derived from pethidine by modification in one or more of the following ways, that is to say:

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

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

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

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

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

(l) Any substance (not being methoxyphenamine) structurally derived from phenethylamine, an N-alkyl-phenethylamine, α-methylphenethylamine, an
N-alkyl-α-methylphenethylamine, α-ethylphenethylamine, or an N-alkyl-α-ethylphenethylamine by substitution in the ring to any extent with alkyl, alkoxy, alkylenedioxy or halo substituents, whether or not further substituted in the ring by one or more other univalent substituents.

(m) Any substance structurally derived from 3-phenylacetylindole by substitution at the nitrogen atom of the indole ring with alkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl or 2-(4-morpholinyl)ethyl, whether or not further substituted in the indole ring to any extent and whether or not substituted in the phenyl ring to any extent.

(n) Any fungus containing any proportion of psilocin or of an ester of psilocin.

(o) 1,2,3,4-Tetrahydronaphthalen-2-amine, 1,2-dihydropthalen-2-amine or 2,3-dihydro-1H-inden-2-amine or any substance structurally derived from 1,2,3,4-tetrahydronaphthalen-2-amine, 1,2-dihydropthalen-2-amine or 2,3-dihydro-1H-inden-2-amine by modification in any of the following ways:

(i) by substitution in the phenyl ring to any extent with alkyl, alkoxy, alkenyl, alkynyl, alkylthio, alkylenedioxy, haloalkyl, hydroxy or halo substituents, whether or not further substituted by one or more other univalent substituents;

(ii) by mono - or di-substitution at the nitrogen atom with alkyl, alkenyl, alkynyl or haloalkyl groups or by inclusion of the nitrogen atom in a cyclic structure.

(p) Any substance structurally derived from tryptamine or from a ring-hydroxy tryptamine by substitution at the nitrogen atom of the side-chain with one or more alkyl substituents but no other substituent.

[[1B. (a) Alprazolam
Amineptine
Aminorex
Amphetaminil
2-Benzhydrylpiperidine
Bromazepam
Brotizolam
Buprenorphine
Butan-1,4-diol
Butorphanol
Camazepam
Cathine
Chlordiazepoxide
Clobazam
Clonazepam
Clorazepic acid]]
Clotiazepam
Cloxazolam
Delorazepam
Dextropropoxyphene
Diazepam
Diethylpropion
Dihydrofuran-2(3H)-one
Estazolam
Ethchlorvynol
Ethinamate
N-Ethylamphetamine
Ethyl lofazepate
Fencamfamin
Fenethylline
Fenproporex
Fludiazepam
Flunitrazepam
Flurazepam
Glutethimide
Halazepam
Haloxazolam
4-Hydroxybutanoic acid
Ketamine
Ketazolam
Lefetamine
Loprazolam
Lorazepam
Lormetazepam
Mazindol
Mecloqualone
Medazepam
Mefenorex
Meprobamate
Mesocarb
Methyprylon
Midazolam
Nalbuphine
Nimetazepam
Nitrazepam
Nordazepam
Oxazepam
Oxazolam
Pemoline
Pentazocine
Phenazepam
Phentermine
Pinazepam
Prazepam
Propylhexedrine
Pyrovalerone
Selegiline
Temazepam
Tetrazepam
Triazolam
Zaleplon
Zipepranol
Zolpidem
Zopiclone.

(b) Any substance structurally derived from barbituric acid by di-substitution at the 5-position, whether or not there is also substitution at the 1-position by a methyl substituent.

2. Any stereoisomeric form of a substance or product specified in paragraph 1 [or 1A] of this Schedule not being dextromethorphan or dextrorphan.

[2A. Any stereoisomeric form of a substance specified in paragraph 1B.]

3. Any ester or ether of a substance or product specified in paragraph 1 [or 1A] or 2 of this Schedule.

4. Any salt of a substance or product specified in [paragraph 1, 1A, 1B, 2, 2A or 3] of this Schedule.
5. Any preparation or product containing any proportion of a substance or product specified in [paragraph 1, 1A, 1B, 2, 2A, 3 or 4] of this Schedule.

6. In this Schedule—

“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 Irish 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;

“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.