Number 12 of 1977

MISUSE OF DRUGS ACT 1977

REVISED

Updated to 2 April 2020

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 the Emergency Measures in the Public Interest (Covid-19) Act 2020 (2/2020), enacted 27 March 2020, and all statutory instruments up to and including the Misuse of Drugs (Amendment) Regulations 2020 (S.I. No. 99 of 2020), made 2 April 2020, were considered in the preparation of this Revised Act.

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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 2019: this Act is one of a group of Acts included in this collective citation, to be construed together as one (Health Service Executive (Governance) Act 2019 (17/2019), s. 1(3)). 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), s. 36 and s. 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), except 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), the amendment to the Health (Fluoridation of Water Supplies) Act 1960 provided for in section 5 (6) 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)
• Health (General Practitioner Service) Act 2018 (13/2018)
• Health Service Executive (Governance) Act 2019 (17/2019), other than Part 3

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 annotated and includes textual and non-textual amendments, statutory instruments made pursuant to the Act and previous affecting provisions.

An explanation of how to read annotations is available at www.lawreform.ie/annotations.

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.

Where legislation or a fragment of legislation is referred to in annotations, changes to this legislation or fragment may not be reflected in this revision but will be reflected in a revision of the legislation referred to if one is available.

A list of legislative changes to any Act, and to statutory instruments from 1972, may be found linked from the page of the Act or statutory instrument at www.irishstatutebook.ie.

Acts which affect or previously affected this revision

- Parole Act 2019 (28/2019)
- Data Protection Act 2018 (7/2018)
- Misuse of Drugs (Supervised Injecting Facilities) Act 2017 (7/2017)
- Misuse of Drugs (Amendment) Act 2016 (9/2016)
- Criminal Justice (Spent Convictions and Certain Disclosures) Act 2016 (4/2016)
- Misuse Of Drugs (Amendment) Act 2015 (6/2015)
- Criminal Justice (Search Warrants) Act 2012 (33/2012)
- Nurses and Midwives Act 2011 (41/2011)
- Criminal Procedure Act 2010 (27/2010)
- Criminal Justice (Psychoactive Substances) Act 2010 (22/2010)
- Fines Act 2010 (8/2010)
- Criminal Justice Act 2007 (29/2007)
- Pharmacy Act 2007 (20/2007)
- Health Act (42/2004)
- Euro Changeover (Amounts) Act 2001 (16/2001)
- Criminal Justice Act 1999 (10/1999)
- Licensing (Combating Drug Abuse) Act 1997 (33/1997)
- Criminal Justice (Drug Trafficking) Act 1996 (29/1996)
- Misuse of Drugs Act 1984 (18/1984)
- Postal and Telecommunications Services Act 1983 (24/1983)
- Transport Act 1950 (12/1950)

All Acts up to and including Emergency Measures in the Public Interest (Covid-19) Act 2020 (2/2020), enacted 27 March 2020, were considered in the preparation of this revision.

Statutory instruments which affect or previously affected this revision

- Misuse of Drugs (Amendment) Regulations 2020 (S.I. No. 99 of 2020)
- Misuse of Drugs (Prescription and Control of Supply of Cannabis for Medical Use) (Amendment) Regulations 2019 (S.I. No. 649 of 2019)
<table>
<thead>
<tr>
<th>Regulation Title</th>
<th>Order/Regulation No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Misuse of Drugs (Prescription and Control of Supply of Cannabis for Medical Use) (Amendment) Regulations 2019</td>
<td>S.I. No. 583 of 2019</td>
</tr>
<tr>
<td>Misuse Of Drugs (Prescription And Control Of Supply Of Cannabis For Medical Use) Regulations 2019</td>
<td>S.I. No. 282 of 2019</td>
</tr>
<tr>
<td>Misuse Of Drugs (Designation) (Amendment) Order 2019</td>
<td>S.I. No. 281 of 2019</td>
</tr>
<tr>
<td>Misuse Of Drugs (Amendment) Regulations 2019</td>
<td>S.I. No. 262 of 2019</td>
</tr>
<tr>
<td>European Union (Freezing and Confiscation of Instrumentalities and Proceeds of Crime) Regulations 2017</td>
<td>S.I. No. 540 of 2017</td>
</tr>
<tr>
<td>Misuse of Drugs (Designation) (Amendment) Order 2017</td>
<td>S.I. No. 533 of 2017</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) Regulations 2017</td>
<td>S.I. No. 532 of 2017</td>
</tr>
<tr>
<td>Misuse of Drugs Act 1977 (Controlled Drugs) (Declaration) Order 2017</td>
<td>S.I. No. 531 of 2017</td>
</tr>
<tr>
<td>Misuse of Drugs (Supervision of Prescription and Supply of Methadone and Medicinal Products containing Buprenorphine authorised for Opioid Substitution Treatment) Regulations 2017</td>
<td>S.I. No. 522 of 2017</td>
</tr>
<tr>
<td>Misuse of Drugs (Exemption) Order 2017</td>
<td>S.I. No. 175 of 2017</td>
</tr>
<tr>
<td>Misuse of Drugs (Designation) Order 2017</td>
<td>S.I. No. 174 of 2017</td>
</tr>
<tr>
<td>Misuse of Drugs Regulations 2017</td>
<td>S.I. No. 173 of 2017</td>
</tr>
<tr>
<td>Misuse of Drugs (Designation) (Amendment) (No. 2) Order 2014</td>
<td>S.I. No. 584 of 2014</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) (No. 2) Regulations 2014</td>
<td>S.I. No. 583 of 2014</td>
</tr>
<tr>
<td>Misuse of Drugs Act 1977 (Controlled Drugs) (Declaration) Order 2014</td>
<td>S.I. No. 571 of 2014</td>
</tr>
<tr>
<td>Misuse of Drugs (Designation) (Amendment) Order 2014</td>
<td>S.I. No. 324 of 2014</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) Regulations 2014</td>
<td>S.I. No. 323 of 2014</td>
</tr>
<tr>
<td>Misuse of Drugs (Licence Fees) (Amendment) Regulations 2012</td>
<td>S.I. No. 544 of 2012</td>
</tr>
<tr>
<td>Misuse of Drugs (Exemption) (Amendment) Order 2011</td>
<td>S.I. No. 554 of 2011</td>
</tr>
<tr>
<td>Misuse of Drugs (Designation) (Amendment) Order 2011</td>
<td>S.I. No. 553 of 2011</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) Regulations 2011</td>
<td>S.I. No. 552 of 2011</td>
</tr>
<tr>
<td>Misuse of Drugs Act 1977 (Controlled Drugs) (Declaration) Order 2011</td>
<td>S.I. No. 551 of 2011</td>
</tr>
<tr>
<td>Finance (Transfer of Departmental Administration and Ministerial Functions) Order 2011</td>
<td>S.I. No. 418 of 2011</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) (No. 2) Regulations 2010</td>
<td>S.I. No. 607 of 2010</td>
</tr>
<tr>
<td>Misuse of Drugs (Exemption) (Amendment) Order 2010</td>
<td>S.I. No. 202 of 2010</td>
</tr>
<tr>
<td>Misuse of Drugs (Designation) (Amendment) Order 2010</td>
<td>S.I. No. 201 of 2010</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) Regulations 2010</td>
<td>S.I. No. 200 of 2010</td>
</tr>
<tr>
<td>Misuse of Drugs Act 1977 (Controlled Drugs) (Declaration) Order 2010</td>
<td>S.I. No. 199 of 2010</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) (No. 2) Regulations 2009</td>
<td>S.I. No. 122 of 2009</td>
</tr>
<tr>
<td>Misuse of Drugs Act 1977 (Controlled Drugs) (Declaration) Order 2009</td>
<td>S.I. No. 121 of 2009</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) Regulations 2009</td>
<td>S.I. No. 63 of 2009</td>
</tr>
<tr>
<td>Pharmaceutical Society of Ireland (Fees) Rules 2008</td>
<td>S.I. No. 496 of 2008</td>
</tr>
<tr>
<td>Medicinal Products (Control of Advertising) Regulations 2007</td>
<td>S.I. No. 541 of 2007</td>
</tr>
<tr>
<td>Misuse of Drugs Act 1977 (Controlled Drugs) (Declaration) Order 2006</td>
<td>S.I. No. 55 of 2006</td>
</tr>
<tr>
<td>Misuse of Drugs (Exemption) (Amendment) Order 2006</td>
<td>S.I. No. 54 of 2006</td>
</tr>
<tr>
<td>Misuse of Drugs (Amendment) Regulations 2006</td>
<td>S.I. No. 53 of 2006</td>
</tr>
<tr>
<td>Misuse of Drugs (Scheduled Substances) (Amendment) Regulations 2004</td>
<td>S.I. No. 92 of 2004</td>
</tr>
<tr>
<td>Misuse of Drugs (Scheduled Substances) (Exemption) Order 2004</td>
<td>S.I. No. 91 of 2004</td>
</tr>
<tr>
<td>Misuse of Drugs Act, 1977 (Controlled Drugs) (Declaration No. 4) Order 2004</td>
<td>S.I. No. 78 of 2004</td>
</tr>
<tr>
<td>Misuse of Drugs (Scheduled Substances) (Exemption) (Amendment) Order 2003</td>
<td>S.I. No. 169 of 2003</td>
</tr>
<tr>
<td>Misuse of Drugs Act, 1977 (Controlled Drugs) (Declaration No 3) Order 2003</td>
<td>S.I. No. 43 of 2003</td>
</tr>
<tr>
<td>Misuse of Drugs (Scheduled Substances) (Exemption) (Amendment) Order 2003</td>
<td>S.I. No. 42 of 2003</td>
</tr>
</tbody>
</table>
• Misuse of Drugs (Scheduled Substances) (Amendment) Regulations 2003 (S.I. No. 41 of 2003)
• Customs-free Airport (Extension of Laws) Regulations 2000 (S.I. No. 169 of 2000)
• Misuse of Drugs (Amendment No. 1) Regulations 1999 (S.I. No. 273 of 1999)
• Misuse of Drugs (Supervision of Prescription and Supply of Methadone) Regulations 1998 (S.I. No. 225 of 1998)
• Misuse of Drugs (Designation) Order 1998 (S.I. No. 69 of 1998)
• Misuse of Drugs (Amendment) Regulations 1993 (S.I. No. 342 of 1993)
• Misuse of Drugs (Scheduled Substances) (Exemption) Order 1993 (S.I. No. 341 of 1993)
• Misuse of Drugs (Designation) Order 1993 (S.I. No. 340 of 1993)
• Misuse of Drugs (Exemption) (Amendment) Order 1993 (S.I. No. 339 of 1993)
• Misuse of Drugs (Scheduled Substances) Regulations 1993 (S.I. No. 338 of 1993)
• Misuse of Drugs Act, 1977 (Controlled Drugs) (Declaration) Order 1993 (S.I. No. 328 of 1993)
• Medical Preparations (Advertising) Regulations 1993 (S.I. No. 76 of 1993)
• European Communities (Merchandise Road Transport) Regulations 1991 (S.I. No. 60 of 1991)
• European Communities (Road Passenger Transport) Regulations 1991 (S.I. No. 59 of 1991)
• Misuse of Drugs Regulations 1988 (S.I. No. 328 of 1988)
• Misuse of Drugs (Designation) Order 1988 (S.I. No. 327 of 1988)
• Misuse of Drugs (Exemption) Order 1988 (S.I. No. 326 of 1988)
• European Communities (Merchandise and Road Transport) Regulations 1988 (S.I. No. 180 of 1988)
• Misuse of Drugs (Licence Fees) (Amendment) Regulations 1988 (S.I. No. 11 of 1988)
• Misuse of Drugs (Exemption) (Amendment) Order 1987 (S.I. No. 264 of 1987)
• Misuse of Drugs (Amendment) Regulations 1987 (S.I. No. 263 of 1987)
• Misuse of Drugs Act, 1977 (Controlled Drugs) (Declaration) Order 1987 (S.I. No. 251 of 1987)
• European Communities (Recognition of Qualifications in Pharmacy) Regulations 1987 (S.I. No. 239 of 1987)
• Misuse of Drugs (Licence Fees) (Amendment) Regulations 1986 (S.I. No. 172 of 1986)
• Misuse of Drugs (Licence Fees) (Amendment) Regulations 1985 (S.I. No. 29 of 1985)
• Misuse of Drugs (Committees of Inquiry) Regulations 1984 (S.I. No. 264 of 1984)
• Misuse of Drugs (Safe Custody) Regulations 1982 (S.I. No. 321 of 1982)
• European Communities (Recognition of Dental Qualifications) Regulations 1980 (S.I. No. 90 of 1980)
• Misuse of Drugs (Custodial Treatment Centre) Order 1980 (S.I. No. 30 of 1980)
• Misuse of Drugs (Licence Fees) Regulations 1979 (S.I. No. 164 of 1979)
• Misuse of Drugs Regulations 1979 (S.I. No. 32 of 1979)
• Misuse of Drugs (Committees of Inquiry, Advisory Committees and Advisory Panels) Regulations 1979 (S.I. No. 31 of 1979)
• Misuse of Drugs (Designation) Order 1979 (S.I. No. 30 of 1979)
• Misuse of Drugs (Exemption) Order 1979 (S.I. No. 29 of 1979)
• Misuse of Drugs Act, 1977 (Commencement) Order 1979 (S.I. No. 28 of 1979)

All statutory instruments up to and including Misuse of Drugs (Amendment) Regulations 2020 (S.I. No. 99 of 2020), made 2 April 2020, were considered in the preparation of this revision.
Number 12 of 1977

MISUSE OF DRUGS ACT 1977
REVISED
Updated to 2 April 2020

ARRANGEMENT OF SECTIONS

Section
1. Interpretation.
2. Controlled drugs.
3. Restriction on possession of controlled drugs.
4. Regulations permitting possession of controlled drugs.
5. Regulations to prevent misuse of controlled drugs.
6. Directions prohibiting prescribing, supply etc. of controlled drugs by practitioners or pharmacists convicted of offences.
7. Special directions prohibiting prescribing etc. of controlled drug in certain cases.
8. Investigation of cases where Minister considers there are grounds for special direction.
9. Prohibition of prescribing etc. in cases of urgency.
10. Investigation on initiative of Dental Board, Medical Registration Council or Veterinary Council. (repealed)
11. Appeals.
12. Regulations (committees and panels).
13. Additional powers in relation to certain controlled drugs.
14. Licences etc.
15. Possession of controlled drugs for unlawful sale or supply.
15A. Offence relating to possession of drugs with value of €13,000 or more.
15B. Importation of controlled drugs in excess of certain value.
15C. Supply of controlled drugs into prisons and places of detention.
16. Prohibition of certain activities etc. relating to opium.
17. Prohibition of cultivation of opium poppy or cannabis plant.
18. Forged or fraudulently altered prescriptions.
19. Occupiers etc. permitting certain activities to take place on land, vehicle or vessel to be guilty of an offence.
20. Offences relating to acts outside the State.
Section
21. Attempts etc. and miscellaneous other offences.
22. Onus of proof.
23. Power of Garda Síochána to search persons, vehicles, vessels or aircraft.
24. Powers to inspect and demand production of drugs, books or documents.
26. Search warrants.
27. Penalties.
28. 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.
29. Defences generally.
30. Forfeiture.
32. Poisons for purposes of Pharmacy Acts, 1875 to 1962.
34. Amendment of section 2 of Pharmacy Act, 1962.
35. Amendment of Pharmacopoeia Act, 1931.
36. Amendment of section 65 of Health Act, 1947.
37. Service etc. of notices.
38. Regulations generally; laying of orders.
39. Expenses.
41. Repeal of Dangerous Drugs Act, 1934, and transitional provision.
42. Miscellaneous repeals and transitional provisions.
43. Short title, commencement and collective citations.

SCHEDULE
ACTS REFERRED TO

<table>
<thead>
<tr>
<th>Act</th>
<th>Year</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dentists Act, 1928</td>
<td>1928, No. 25</td>
</tr>
<tr>
<td>Pharmacy Act, 1951</td>
<td>1951, No. 30</td>
</tr>
<tr>
<td>Amendment Act, 1890</td>
<td>1890, c. 48</td>
</tr>
<tr>
<td>Medical Practitioners Act, 1927</td>
<td>1927, No. 25</td>
</tr>
<tr>
<td>Pharmacy Act (Ireland), 1875</td>
<td>1875, c. 57</td>
</tr>
<tr>
<td>Veterinary Surgeons Act, 1931</td>
<td>1931, No. 36</td>
</tr>
<tr>
<td>Pharmacy Act, 1962</td>
<td>1962, No. 14</td>
</tr>
<tr>
<td>Pharmacopoeia Act, 1931</td>
<td>1931, No. 22</td>
</tr>
<tr>
<td>Health Act, 1947</td>
<td>1947, No. 28</td>
</tr>
<tr>
<td>Health Act, 1953</td>
<td>1953, No. 26</td>
</tr>
<tr>
<td>Act</td>
<td>Year</td>
</tr>
<tr>
<td>------------------------------------------</td>
<td>---------</td>
</tr>
<tr>
<td>Health Act, 1970</td>
<td>1970, No. 1</td>
</tr>
<tr>
<td>Companies Act, 1963</td>
<td>1963, No. 33</td>
</tr>
<tr>
<td>Public Offices Fees Act, 1879</td>
<td>1879, c. 58</td>
</tr>
<tr>
<td>Dangerous Drugs Act, 1934</td>
<td>1934, No. 1</td>
</tr>
</tbody>
</table>

BE IT ENACTED BY THE OIREACHTAS AS FOLLOWS:

Annotations

Modifications (not altering text):


Dissolution of health boards and other specified bodies.

58.—The specified bodies are, by this Act, dissolved on the establishment day.

Transfer of functions of specified bodies to Executive.

59.—(1) The functions that, immediately before the establishment day, were the functions of a specified body under or in connection with the enactments referred to in Schedule 3 are, by this Act, transferred to the Executive on that day.

(2) If a provision of an enactment referred to in Schedule 3 , or a provision of an instrument made under such enactment, does not come into effect until on or after the establishment day, a function that on the passing of that enactment or the making of that instrument was assigned under or in connection with that provision to a specified body is, by this Act, transferred to the Executive on the commencement of that provision.

(3) The functions transferred by this Act to the Executive include the functions specified in any enactment referred to in Schedule 3 as a function of the following:

(a) the chief executive officer of a health board;

(b) the Regional Chief Executive of the Eastern Regional Health Authority;

(c) the area chief executive of an Area Health Board.
SCHEDULE THREE

Transfer of Functions and References to Functional Areas

Section 59 and 67.


2. The provisions of the Misuse of Drugs Acts 1977 and 1984 (Nos. 12 and 18 of 1977 and 1984 respectively), together with all Orders and Regulations made thereunder and for the time being in force, are hereby extended to the Customs-free Airport at Shannon.

C3 Application of collectively cited Health Acts restricted (7.05.1986) by Health (Amendment) Act 1986 (10/1986), s. 2(1), commenced on enactment. Note: references to health board to be read as references to the Health Service Executive as provided (1.01.2005) by Health Act 2004 (42/2004), s. 66, S.I. No. 887 of 2004.

Charges by health boards for provision of in-patient services and out-patient services in respect of certain injuries caused by mechanically propelled vehicles.

2.—(1) Where—

(a) injury is caused to a person by the negligent use of a mechanically propelled vehicle in a public place, and

(b) in-patient services or out-patient services have been, are being or will be provided by or on behalf of a health board in respect of the injury, and

(c) any one of the following, that is to say, the person aforesaid, his personal representative or dependant, has received, or is entitled to receive damages or compensation in respect of the negligent use aforesaid from the person liable to pay such damages or compensation in respect of that injury, or any loss, damage or expense (or mental distress in the case of a dependant) arising therefrom,

the health board shall, notwithstanding anything in the Health Acts, 1947 to 1985, make a charge upon the person who received or is entitled to receive such damages or compensation in respect of the said in-patient services or out-patient services.

Editorial Notes:


E3 Offence under Act (other than a first offence under s. 3) designated an excluded offence by National Vetting Bureau (Children and Vulnerable Persons) Act 2012 (47/2012), s. 14A and sch. 3 item 16, as inserted (29.04.2016) by Criminal Justice (Spent Convictions and Certain Disclosures) Act 2016 (4/2016), s. 29, S.I. No. 215 of 2016.

E5 Procedure for complaints in relation to registered retail pharmacy businesses, where certain persons have been convicted of an offence under Misuse of Drugs Acts 1977 to 2006 or Poisons Acts 1961 and 1977 prescribed (1.08.2009) by Pharmacy Act 2007 (20/2007), s. 36(1)(a), S.I. No. 281 of 2009.


Interpretation.

1.—(1) In this Act—

F1‘business’ includes a profession.

F2‘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;
F3[‘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;
F4[‘opium poppy’ means a plant of the species Papaver somniferum L or Papaver
bracteatum Lind.;]
“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;
F5[“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;
F6[“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;]
F5[“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;

F7 [‘temporary direction’ means a direction under section 9 of this Act;]

F8 [‘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.

Annotations

Amendments:


Modifications (not altering text):

C4 References to a pharmaceutical chemist registered under the Pharmacy Act (Ireland) 1875 or a dispensing chemist and druggist registered under the Pharmacy Act 1951 construed (29.11.2008) by Pharmacy Act 2007 (20/2007), s. 75(1), S.I. No. 487 of 2008.

Interpretation of references to pharmaceutical chemists, persons keeping open shop, etc.

75.— (1) Any reference (however expressed) in a prior enactment to a pharmaceutical chemist registered under the Pharmacy Act (Ireland) 1875 or a dispensing chemist and druggist registered under the Pharmacy Act 1951 shall be construed as a reference to a registered pharmacist.

...
Construction of references to registered medical practitioner and Medical Council, etc.

108.—(1) Every reference to a registered medical practitioner contained in any enactment or any statutory instrument shall be construed as a reference to a registered medical practitioner within the meaning of section 2.

(3) Every reference to—

(b) the Medical Registration Council,

contained in any other enactment or any statutory instrument shall be construed as the Council within the meaning of section 2.

Editorial Notes:

E14 Previous affecting provision: definition of “practitioner” amended (1.05.2007) by Irish Medicines Board (Miscellaneous Provisions) Act 2006 (3/2006), s. 3(c), S.I. No. 194 of 2007; substituted as per F-note above.

E15 Previous affecting provision: definition of “registered nurse” inserted (1.05.2007) by Irish Medicines Board (Miscellaneous Provisions) Act 2006 (3/2006), s. 3(d), S.I. No. 194 of 2007; substituted as per F-note above.

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).

Annotations

Editorial Notes:


E18 Power pursuant to subs. (2) exercised (11.11.2011) by Misuse of Drugs Act 1977 (Controlled Drugs) (Declaration) Order 2011 (S.I. No. 551 of 2011), in effect as per art. 2.

E19 Power pursuant to subs. (2) exercised (11.05.2010) by Misuse of Drugs Act 1977 (Controlled Drugs) (Declaration) Order 2010 (S.I. No. 199 of 2010), in effect as per art. 2.
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).

Annotations

Modifications (not altering text):

C6 Application of subss. (1) and (2) restricted (30.11.2017) by Misuse of Drugs (Supervised Injecting Facilities) Act 2017 (7/2017), s. 10(1), S.I. No. 517 of 2017.

Disapplication of sections 3, 19 and 21 of Act of 1977

10. (1) Subsections (1) and (2) of section 3 of the Act of 1977 do not apply to an authorised user.

...
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.
Annotations

Editorial Notes:


E43 Power pursuant to section exercised (26.06.2019) by Misuse Of Drugs (Prescription And Control Of Supply Of Cannabis For Medical Use) Regulations 2019 (S.I. No. 262 of 2019).


E45 Power pursuant to section exercised (4.05.2017) by Misuse of Drugs Regulations 2017 (S.I. No. 173 of 2017), in effect as per reg. 1(2).


E48 Power pursuant to section exercised (1.11.2011) by Misuse of Drugs (Amendment) Regulations 2011 (S.I. No. 552 of 2011).

E49 Power pursuant to section exercised (11.05.2010) by Misuse of Drugs (Amendment) Regulations 2010 (S.I. No. 200 of 2010).

E50 Power pursuant to section exercised (31.03.2009) by Misuse of Drugs (Amendment) (No. 2) Regulations 2009 (S.I. No. 122 of 2009).


E52 Power pursuant to section exercised (1.05.2007) by Misuse of Drugs (Amendment) Regulations 2007 (S.I. No. 200 of 2007).

E53 Power pursuant to section exercised (31.01.2006) by Misuse of Drugs (Amendment) Regulations 2006 (S.I. No. 53 of 2006), in effect as per reg. 2.


E55 Power pursuant to section exercised (1.03.1983) by Misuse of Drugs (Safe Custody) Regulations 1982 (S.I. No. 321 of 1982). Note that this SI was given statutory status (12.03.2015) by Misuse Of Drugs (Amendment) Act 2015 (6/2015), s. 2 and sch. 2, commenced as per s. 1(3). It was revoked by Misuse Of Drugs (Amendment) Act 2016 (9/2016), s. 7(b), not commenced as of date of revision.


E57 Previous affecting provision: power pursuant to section exercised (1.03.2004) by Misuse of Drugs (Scheduled Substances) (Amendment) Regulations 2004 (S.I. No. 92 of 2004); revoked (22.12.2009) by European Communities (Control of Drug Precursors) Regulations 2009 (S.I. No. 558 of 2009), reg. 37(1)(c).
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.

F9[(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 F10[[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 F11[...] manu-
ufacture, 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.

F9[(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.]

F12[(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.]
Annotations

Amendments:

F9 Substituted (4.05.2017) by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 3(a), (b), (d), S.I. No. 172 of 2017.

F10 Substituted (4.05.2017) by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 3(c), S.I. No. 172 of 2017.


F12 Inserted (4.05.2017) by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 3(e), S.I. No. 172 of 2017.

Modifications (not altering text):


Construction of references to registered medical practitioner and Medical Council, etc.

108. — (1) Every reference to a registered medical practitioner contained in any enactment or any statutory instrument shall be construed as a reference to a registered medical practitioner within the meaning of section 2.

C8 References to a person dispensing prescriptions for controlled drugs under Pharmacy Act (Ireland) 1875 or a dispensing chemist and druggist registered under the Pharmacy Act 1951 construed (29.11.2008) by Pharmacy Act 2007 (20/2007), s. 75(2), S.I. No. 487 of 2008.

Interpretation of references to pharmaceutical chemists, persons keeping open shop, etc.

75. — ...  
(2) Any reference (however expressed) in a prior enactment to a person who is keeping open shop for the dispensing or compounding of medical prescriptions under the Pharmacy Acts 1875 to 1977—

(a) where that person is, in relation to a pharmacist, a representative within the meaning given by section 25(2), shall be construed as a reference to such a representative, and

(b) in any other case, shall be construed as a reference to a registered retail pharmacy business.

Editorial Notes:


E66 Power pursuant to section exercised (26.06.2019) by Misuse Of Drugs (Prescription And Control Of Supply Of Cannabis For Medical Use) Regulations 2019 (S.I. No. 262 of 2019).


E68 Power pursuant to section exercised (22.11.2017) by Misuse of Drugs (Supervision of Prescription and Supply of Methadone and Medicinal Products containing Buprenorphine authorised for Opioid Substitution Treatment) Regulations 2017 (S.I. No. 522 of 2017), in effect as per reg. 1(2).

E69 Power pursuant to section exercised (4.05.2017) by Misuse of Drugs Regulations 2017 (S.I. No. 173 of 2017), in effect as per reg. 1(2).
E70  Power pursuant to section exercised (17.12.2014) by Misuse of Drugs (Amendment) (No. 2) Regulations 2014 (S.I. No. 583 of 2014).


E72  Power pursuant to section exercised (1.11.2011) by Misuse of Drugs (Amendment) Regulations 2011 (S.I. No. 552 of 2011).


E74  Power pursuant to section exercised (11.06.2010) by Misuse of Drugs (Amendment) Regulations 2010 (S.I. No. 200 of 2010).

E75  Power pursuant to section exercised (31.03.2009) by Misuse of Drugs (Amendment) Regulations 2009 (S.I. No. 122 of 2009).

E76  Power pursuant to section exercised (1.05.2007) by Misuse of Drugs (Amendment) Regulations 2007 (S.I. No. 200 of 2007).

E77  Power pursuant to section exercised (31.01.2006) by Misuse of Drugs (Amendment) Regulations 2006 (S.I. No. 53 of 2006), in effect as per reg. 2.

E78  Power pursuant to section exercised (27.08.1999) by Misuse of Drugs (Amendment No. 1) Regulations 1999 (S.I. No. 273 of 1999).


E83  Power pursuant to section exercised (1.03.1983) by Misuse of Drugs (Safe Custody) Regulations 1982 (S.I. No. 321 of 1982), in effect as per reg. 2. Note that this SI was given statutory status (12.03.2015) by Misuse Of Drugs (Amendment) Act 2015 (6/2015), s. 2 and sch. 2, commenced as per s. 1(3). It was revoked by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 7(13), not commenced as of date of revision.


E85  Previous affecting provision: power pursuant to section exercised (16.07.1998) by Misuse of Drugs (Supervision of Prescription and Supply of Methadone) Regulations 1998 (S.I. No. 225 of 1998), in effect as per reg. 1(2); given statutory effect (12.03.2015) by Misuse Of Drugs (Amendment) Act 2015 (6/2015), s. 2 and sch. 2, commenced as per s. 1(3); revoked (22.11.2017) by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 7(f), S.I. No. 521 of 2017.

E86  Previous affecting provision: subs. (1)(f) substituted (1.05.2007) by Irish Medicines Board (Miscellaneous Provisions) Act 2006 (3/2006), s. 4(a), S.I. No. 194 of 2007; substituted as per F-note above.

E87  Previous affecting provision: subs. (1A) inserted (1.05.2007) by Irish Medicines Board (Miscellaneous Provisions) Act 2006 (3/2006), s. 4(b), S.I. No. 194 of 2007; substituted as per F-note above.

Directions prohibiting prescribing, supply etc. of controlled drugs by practitioners or pharmacists convicted of offences.

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 F14[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.

F15[(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.]

Annotatıons
Amendments:

5(b), S.I. No. 453 of 2013.


Modifications (not altering text):
C9 References to a person dispensing prescriptions for controlled drugs under the Pharmacy Acts
1875 to 1977 construed (29.11.2008) by Pharmacy Acts 2007 (20/2007), s. 75(2), S.I. No. 487 of
2008.

Interpretation of references to pharmaceutical chemists, persons keeping open shop, etc.
75. — …

(2) Any reference (however expressed) in a prior enactment to a person who is keeping open
shop for the dispensing or compounding of medical prescriptions under the Pharmacy Acts 1875
to 1977—

(a) where that person is, in relation to a pharmacist, a representative within the meaning
given by section 25(2), shall be construed as a reference to such a representative, and

(b) in any other case, shall be construed as a reference to a registered retail pharmacy business.
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 administration 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 practitioner 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 F16[...] suspend the operation of a special direction.

(4) The Minister may F16[...] 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.

Annotations

Amendments:


F17[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 investigated, 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.

Annotations

Amendments:


Modifications (not altering text):


Construction of references to registered medical practitioner and Medical Council, etc.

108. — (1) Every reference to a registered medical practitioner contained in any enactment or any statutory instrument shall be construed as a reference to a registered medical practitioner within the meaning of section 2.

...
(3) Every reference to—
(a) the Medical Council, or

... contained in any other enactment or any statutory instrument shall be construed as the Council within the meaning of section 2.

Editorial Notes:

E98 Previous affecting provision: power pursuant to section exercised (1.05.1979) by Misuse of Drugs (Committees of Inquiry, Advisory Committees and Advisory Panels) Regulations 1979 (S.I. No. 31 of 1979), in effect as per reg. 2; revoked (23.10.1984) by Misuse of Drugs (Committees of Inquiry) Regulations 1984 (S.I. No. 264 of 1984), reg. 8.

Prohibition of prescribing etc. in cases of urgency.

F20[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 authorising the administration or supply of such controlled drugs as may 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 so 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 representations 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.]

Annotations

Amendments:

Editorial Notes:
E99 Previous affecting provision: power pursuant to section exercised (1.05.1979) by Misuse of Drugs (Committees of Inquiry, Advisory Committees and Advisory Panels) Regulations 1979 (S.I. No. 31 of 1979), in effect as per reg. 2; revoked (23.10.1984) by Misuse of Drugs (Committees of Inquiry) Regulations 1984 (S.I. No. 264 of 1984), reg. 8.
Investigation on
initiative of
Dental Board,
Medical Registration Council or
Veterinary Council.

10.—F21[...]

Annotations

Amendments:


Editorial Notes:

E100 Previous affecting provision: power pursuant to section exercised (1.05.1979) by Misuse of Drugs (Committees of Inquiry, Advisory Committees and Advisory Panels) Regulations 1979 (S.I. No. 31 of 1979), in effect as per reg. 2; revoked (23.10.1984) by Misuse of Drugs (Committees of Inquiry) Regulations 1984 (S.I. No. 264 of 1984), reg. 8.

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.

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 F22[...] established pursuant to section 8 F22[...] of this Act.

(2) 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 F23[...] referred to in subsection (1) of this section may regulate its procedure and business.

Annotations

Amendments:


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).

Annotations

Amendments:

F24 Substituted by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 4, not commenced as of date of revision.

Modifications (not altering text):

C11 Prospective affecting provision: subs. (1)(b) amended by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 4, not commenced as of date of revision.

13.—(1) ...

(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 Health Products Regulatory Authority],

...

Editorial Notes:


E105 Power pursuant to section exercised (4.05.2017) by Misuse of Drugs (Designation) Order 2017 (S.I. No. 174 of 2017), in effect as per art. 2.
Licences etc.

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.

Annotations

Amendments:

F25 Substituted by Irish Medicines Board (Miscellaneous Provisions) Act 2006 (3/2006), s. 7(a), not commenced as of date of revision.

F26 Inserted by Irish Medicines Board (Miscellaneous Provisions) Act 2006 (3/2006), s. 7(b), not commenced as of date of revision.

Modifications (not altering text):


14.—(1) The F25[Irish Medicines Board] 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.

F26(3) A licence, permit or authorisation—

(o) granted or issued by the Minister under subsection (1) (including granted or issued by way of being renewed) at any time before the commencement of this subsection, and
(b) in force immediately before that commencement,
shall, on and after that commencement but subject to the conditions, if any, attached under subsection (1) to it and in force immediately before that commencement, continue in force, unless sooner revoked under subsection (1), for the unexpired portion of the period of validity, if any, which it had left to run immediately before that commencement as if, on that commencement, the Irish Medicines Board had, under subsection (1)—
(c) granted or issued that licence, permit or authorisation, and
(d) attached to that licence, permit or authorisation those conditions, if any,
and the provisions of this Act shall apply to the licence, permit or authorisation accordingly.

Editorial Notes:

E114 Prospective affecting provision: in carrying out its duties under subs. (1) the Irish Medicines Board is required to comply with any directive or guideline issued by Minister for Health by Irish Medicines Board Act 1995 (29/1995), s. 4(5), as substituted by Irish Medicines Board (Miscellaneous Provisions) Act 2006 (3/2006), s. 11(b); amendment transferring duties to Irish Medicines Board not commenced as of date of revision.

E115 Power pursuant to section exercised (1.01.2013) by Misuse of Drugs (Licence Fees) (Amendment) Regulations 2012 (S.I. No. 544 of 2012), in effect as per reg. 3.

E116 Power pursuant to section exercised (1.02.1988) by Misuse of Drugs (Licence Fees) (Amendment) Regulations 1988 (S.I. No. 11 of 1988), in effect as per reg. 3.

E117 Power pursuant to section exercised (1.06.1986) by Misuse of Drugs (Licence Fees) (Amendment) Regulations 1986 (S.I. No. 172 of 1986), in effect as per reg. 3.

E118 Power pursuant to section exercised (1.03.1985) by Misuse of Drugs (Licence Fees) (Amendment) Regulations 1985 (S.I. No. 29 of 1985), in effect as per reg. 3.

E119 Power pursuant to section exercised (15.05.1979) by Misuse of Drugs (Licence Fees) Regulations 1979 (S.I. No. 164 of 1979), in effect as per reg. 2. Note that this SI was given statutory status (12.03.2015) by Misuse Of Drugs (Amendment) Act 2015 (6/2015), s. 2 and sch. 2, commenced as per s. 1(3). It was revoked by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 7(a), not commenced as of date of revision.

Possession of controlled drugs for unlawful sale or supply.

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.

Annotations

Editorial Notes:

E120 Subs. (1) designated as relevant offence for purposes of retrial and double jeopardy procedure (1.09.2010) by Criminal Procedure Act 2010 (27/2010), ss. 7-18 and sch. para. 16, S.I. No. 414 of 2010.
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.

F29[(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.
Annotatons

Amendments:


F28 Substituted (1.01.2002) by Euro Changeover (Amounts) Act 2001 (16/2001), s. 1(3) and (4) and schs. 3 ref. no. 6 and 4 ref. no. 9, commenced as per s. 1(3).


Editorial Notes:


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.

(5) In this section ‘market value’ and ‘an officer of customs and excise’ have the meanings they have in section 15A of this Act.]
F31 Supply of controlled drugs into prisons and places of detention.

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.

Annotations

Amendments:


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, 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) [F34].

(F32) [(2) Every person who cultivates opium poppy, 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.]

Annotatons

Amendments:


Editorial Notes:

E128 Offences relating to the sale and supply of objects in the use of cultivation of plants in contravention of section prescribed and powers and procedures established (23.08.2010) by Criminal Justice (Psychoactive Substances) Act 2010 (22/2010), ss. 4, 5, 7-10, 12-14 and 21, S.I. No. 401 of 2010.

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.

Annotations

Editorial Notes:


Occupiers etc. permitting certain activities to take place on land, vehicle or vessel to 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.
Annot ations

Modific ations (not alt ering text):


Disapplic ation of sections 3, 19 and 21 of Act of 1977

10. ...

(2) Section 19(1)(e) of the Act of 1977, in so far as it relates to the preparation or production for immediate personal consumption by injection of a controlled drug by an authorised user, and section 19(1)(i) of the Act of 1977, do not apply to a licence holder who knowingly permits or suffers the preparation or production for immediate personal consumption by injection or the possession of a controlled drug, in a supervised injecting facility by an authorised user.

...


Amendmen t of sections 17 and 19 of Principal Act.

11. —...

(2) Section 19 of the Principal Act shall be construed and have effect as if the reference in subsection (1) (a) thereof to the cultivation contrary to section 17 of that Act of opium poppy included a reference to the cultivation contrary to the said section 17, as amended by subsection (1) of this section, of any plant of the genus Erythroxylon.

Editorial Not es:

E139 Power to apply for revoc ation or forfeitur e of licence and disqualific ation fr om holding licence on conviction for an offence under subs. (1)(g) prescribed (21.06.1997) by Licensing (Combating Drug Abuse) Act 1997 (33/1997), ss. 3 and 17, commenced as per s. 22(4).


E141 Previous affecting provision: offences under section designated relevant offences (21.03.1991) for the purposes of European Communities (Road Passenger Transport) Regulations 1991 (S.I. No. 59 of 1991), reg. 2(1) and sch. 1 para. (g); revoked (4.12.2011) by European Union (Occupation of Road Transport Operator) Regulations 2011 (S.I. No. 697 of 2011), reg. 11(a).

E142 Previous affecting provision: offences under section designated relevant offences (30.09.1988) for the purposes of European Communities (Merchandise Road Transport) Regulations 1988 (S.I. No. 180 of 1988), reg. 2 and sch. 1 para. (k), in effect as per reg. 1(3); revoked (21.03.1991) by European Communities (Merchandise Road Transport) Regulations 1991 (S.I. No. 60 of 1991), reg. 21(1), subject to transitional provisions in regs. 14(1) and 21(2), (3).

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.

Annotations

Editorial Notes:


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) Any 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.

Annotations

Amendments:

F36 Substituted by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 5, not commenced as of date of revision.
Modifications (not altering text):

C15 Prospective affecting provision: subs. (6) amended by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 5, not commenced as of date of revision.

(6) Any person who contravenes a condition attached to a licence, permit or authorisation granted or [issued under section 14 by the Health Products Regulatory Authority] under this Act (other than section 24) or under regulations made under this Act shall be guilty of an offence.

...


Disapplication of sections 3, 19 and 21 of Act of 1977

10. ...

(3) Section 21(2) of the Act of 1977, in so far as it relates to the possession, preparation or production of a controlled drug for immediate personal consumption by injection does not apply to an authorised user.

Onus of proof.

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 F37[ 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.

Annotations

Amendments:


Power of Garda Síochána to search persons, vehicles, vessels or aircraft.

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 F38[(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) F38[(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.

F38[(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.

Annotations

Amendments:


24.—(1) For the purpose of enforcing this Act and regulations made thereunder, a member of the Garda Síochána or a person authorised in F39[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 F40[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 F41[[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.

F42[(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 F41[[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 2016 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 F44[or if, F45(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

F46[(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,

F47[such Justice, Commissioner or, as the case may be, member] may issue a search warrant mentioned in subsection (2) of this section.

F48[(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.]

Power of Garda Síochána to enter supervised injecting facility

11. Notwithstanding section 26 of the Act of 1977, a member of An Garda Síochána, whether in uniform or not, or accompanied by such other person as may be necessary, may at any time enter the premises of a supervised injecting facility without a warrant, and there make such inspection, examination, observation and enquiry as he or she may think proper for the prevention or detection of offences under the Misuse of Drugs Acts 1977 to 2016, other than offences, which pursuant to section 10, do not apply to an authorised user.

C19 Application of section restricted (9.09.1996) by Criminal Justice (Drug Trafficking) Act 1996 (29/1996), s. 8(2)-(3), S.I. No. 257 of 1996, as amended (25.07.2012) by Criminal Justice (Search Warrants) Act 2012 (33/2012), s. 3(1), commenced as per s. 4(2), subject to transitional provision in s. 3(2).

Search warrants.

8.— ...

(2) A member of the Garda Síochána not below the rank of superintendent shall not issue a search warrant under the said section 26 unless he or she is satisfied—

(a) that the warrant is necessary for the proper investigation of a drug trafficking offence, and

(b) that circumstances of urgency giving rise to the need for the immediate issue of the search warrant would render it impracticable to apply to a judge of the District Court or a Peace Commissioner under the said section 26 for the issue of the warrant.

(2A) (a) A member of the Garda Síochána not below the rank of superintendent may issue a search warrant under the said section 26 only if he or she is independent of the investigation of the offence in relation to which the search warrant is being sought.

(b) In this subsection ‘independent of’, in relation to the investigation of an offence, means not being in charge of, or involved in, that investigation.

(2B) A member of the Garda Síochána not below the rank of superintendent who issues a search warrant under the said section 26 shall, either at the time the warrant is issued or as soon as reasonably practicable thereafter, record in writing the grounds on which the warrant was issued, including how he or she was satisfied as to the matters referred to in subsection (2).

(3) Notwithstanding subsection (2) of section 26 of the Act of 1977, a search warrant issued by a member of the Garda Síochána not below the rank of superintendent under subsection (1) of that section shall cease to have effect after a period of 24 hours has elapsed from the time of the issue of the warrant.

Penalties.

27.— F49[(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 impris-
onment;

(b) in any other case—
   (i) on summary conviction, to a fine not exceeding £1,000 or, at the discr-
etion 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 imprison-
ment.

(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 discr-
etion 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 discr-
etion 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.

F50[(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

F50 41
(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—

(II) the circumstances in which the indication was given,

(ii) 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).]
(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) F51[...]

(11) F51[...]

(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.
Annotations

Amendments:


Modifications (not altering text):

C20 References to fines in subss. (1)-(4) and (6)-(9) construed (25.06.2001) as per conversion in Euro Changeover (Amounts) Act 2001 (16/2001), s. 1 and sch. 1, commenced on enactment.

The Euro values of fines in this section are as follows:

£200 = €253.95 (Class E fine, see below)
£300 = €380.92 (Class D fine, see below)
£400 = €507.90 (Class D fine, see below)
£500 = €634.87 (Class C fine, see below)
£1,000 = €1,269.74 (Class C fine, see below)

References to fines in subss. (1)-(4) and (6)-(9) construed (4.01.2011) by indexation in Fines Act 2010 (8/2010), ss. 3, 6, 7 and 8, S.I. No. 662 of 2010.

Definitions.

3. — In this Part— ... 

"class C fine" means a fine not exceeding €2,500;
"class D fine" means a fine not exceeding €1,000;
"class E fine" means a fine not exceeding €500;
...

"commencement date" means the date of the coming into operation of this Part;

"enactment" means—

(a) an Act of the Oireachtas,

(b) a statute that was in force in Saorstát Éireann immediately before the date of the coming into operation of the Constitution and that continues in force by virtue of Article 50 of the Constitution, or

(c) an instrument made under—

(i) an Act of the Oireachtas, or

(ii) such a statute.

Class C fines.

6. — ... 

(3) Where an enactment enacted before the commencement date provides that a person who commits an offence under the enactment shall be liable, upon summary conviction, to a fine not exceeding an amount that—

(a) was provided for by virtue of a subsequent enactment enacted during a period specified in column (2) of the Table opposite a particular reference number specified in column (1) of the Table, and

(b) falls within the range of amounts specified in column (3) of the Table opposite the same reference number,
a person who commits that offence after the commencement date shall, upon summary conviction, not be liable to that fine but shall instead be liable to a class C fine.

### Class D fines.

(3) Where an enactment enacted before the commencement date provides that a person who commits an offence under the enactment shall be liable, upon summary conviction, to a fine not exceeding an amount that—

(a) was provided for by virtue of a subsequent enactment enacted during a period specified in column (2) of the Table opposite a particular reference number specified in column (1) of the Table, and

(b) falls within the range of amounts specified in column (3) of the Table opposite the same reference number,

a person who commits that offence after the commencement date shall, upon summary conviction, not be liable to that fine but shall instead be liable to a class D fine.

### Class E fines.

(3) Where an enactment enacted before the commencement date provides that a person who commits an offence under the enactment shall be liable, upon summary conviction, to a fine not exceeding an amount that—

(a) was provided for by virtue of a subsequent enactment enacted during a period specified in column (2) of the Table opposite a particular reference number specified in column (1) of the Table, and

(b) falls within the range of amounts specified in column (3) of the Table opposite the same reference number,

a person who commits that offence after the commencement date shall, upon summary conviction, not be liable to that fine but shall instead be liable to a class E fine.
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, F52 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 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 15 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, F52 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 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) a condition requiring such person to undergo medical treatment recommended in the report,

(iii) 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,

(iv) a condition requiring the person to attend a specified course of education, instruction or training, being a course which, if undergone by such person, would, in the opinion of the court, improve his vocational opportunities or social circumstances, facilitate his social rehabilitation or reduce the likelihood of his committing a further offence under this Act, or

(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.

(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 F58[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 F59[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 F59[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 F59[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;

F60[...]
Defences generally.

29.—F62[(1) In any proceedings for an offence under this Act or an offence under section 34 of the Criminal Justice Act, 1994 in which it is proved that the defendant had in his possession or supplied a controlled drug, the defendant shall not be acquitted of the offence charged by reason only of proving that he neither knew nor suspected nor had reason to suspect that the substance, product or preparation in question was the particular controlled drug alleged.]

(2) In any such proceedings in which it is proved that the defendant had in his possession a controlled drug, or a forged prescription, or a duly issued prescription altered with intent to deceive, it shall be a defence to prove that—

(a) he did not know and had no reasonable grounds for suspecting—

(i) that what he had in his possession was a controlled drug or such a prescription, as may be appropriate, or

(ii) that he was in possession of a controlled drug or such a prescription, as may be appropriate, or

(b) he believed the substance, product or preparation to be a controlled drug, or a controlled drug of a particular class or description, and that, if the substance, product or preparation had in fact been that controlled drug or a controlled drug of that class or description, he would not at the material time have been committing an offence under this Act, or

(c) knowing or suspecting it to be such a drug or prescription, he took or retained possession of it for the purpose of—

(i) preventing another from committing or continuing to commit an offence in relation to the drug or document, as may be appropriate, or

(ii) delivering it into the custody of a person lawfully entitled to take custody of it,

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.

F63[(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.

Annotations
Amendments:

Editorial Notes:
E155 Previous affecting provision: subs. (3) substituted (26.05.1999) by Criminal Justice Act 1999 (10/1999), s. 6, S.I. No. 154 of 1999; substituted as per F-note above.

Forfeiture.

30.—F64[(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.

Annotations
Amendments:

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.

Annotations
Amendments:
F65...

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.”.
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.”.

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.”, and

(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.”.

Service etc. of notices.

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.

Regulations generally; laying of orders.

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.

Annotations

Modifications (not altering text):

C22 Functions transferred and reference to “Minister for Finance” construed (29.07.2011) by Finance (Transfer of Departmental Administration and Ministerial Functions) Order 2011 (S.I. No. 418 of 2011), arts. 3 and 5 and sch. 1 part 2, in effect as per art. 1(2), subject to transitional provisions in arts. 6-9.

3. The functions conferred on the Minister for Finance by or under the provisions of —

(a) the enactments specified in Schedule 1, and

(b) the statutory instruments specified in Schedule 2,

are transferred to the Minister for Public Expenditure and Reform.

...
## Editorial Notes:

- **E159** Power pursuant to section exercised (4.05.2017) by *Misuse of Drugs Regulations 2017* (S.I. No. 173 of 2017), in effect as per reg. 1(2).
- **E162** Power pursuant to section exercised (1.01.2013) by *Misuse of Drugs (Licence Fees) (Amendment) Regulations 2012* (S.I. No. 544 of 2012), in effect as per reg. 3.
- **E163** Power pursuant to section exercised (1.11.2011) by *Misuse of Drugs (Amendment) Regulations 2011* (S.I. No. 552 of 2011).
- **E164** Power pursuant to section exercised (11.05.2010) by *Misuse of Drugs (Amendment) Regulations 2010* (S.I. No. 200 of 2010).
- **E165** Power pursuant to section exercised (31.03.2009) by *Misuse of Drugs (Amendment) (No. 2) Regulations 2009* (S.I. No. 122 of 2009).
- **E166** Power pursuant to section exercised (1.05.2007) by *Misuse of Drugs (Amendment) Regulations 2007* (S.I. No. 200 of 2007).
- **E167** Power pursuant to section exercised (31.01.2006) by *Misuse of Drugs (Amendment) Regulations 2006* (S.I. No. 53 of 2006), in effect as per reg. 3.
- **E169** Power pursuant to section exercised (1.02.1988) by *Misuse of Drugs (Licence Fees) (Amendment) Regulations 1988* (S.I. No. 11 of 1988), in effect as per reg. 3.
- **E170** Power pursuant to section exercised (1.06.1986) by *Misuse of Drugs (Licence Fees) (Amendment) Regulations 1986* (S.I. No. 172 of 1986), in effect as per reg. 3.
- **E171** Power pursuant to section exercised (1.03.1985) by *Misuse of Drugs (Licence Fees) (Amendment) Regulations 1985* (S.I. No. 29 of 1985), in effect as per reg. 3.
E172 Power pursuant to section exercised (23.10.1984) by Misuse of Drugs (Committees of Inquiry) Regulations 1984 (S.I. No. 264 of 1984), in effect as per reg. 3.

E173 Power pursuant to section exercised (1.03.1983) by Misuse of Drugs (Safe Custody) Regulations 1982 (S.I. No. 321 of 1982), in effect as per reg. 2. Note that this SI was given statutory status (12.03.2015) by Misuse Of Drugs (Amendment) Act 2015 (6/2015), s. 2 and sch. 2, commenced as per s. 1(3). It was revoked by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 7(b), not commenced as of date of revision.

E174 Power pursuant to section exercised (15.05.1979) by Misuse of Drugs (Licence Fees) Regulations 1979 (S.I. No. 164 of 1979), in effect as per reg. 2. Note that this SI was given statutory status (12.03.2015) by Misuse Of Drugs (Amendment) Act 2015 (6/2015), s. 2 and sch. 2, commenced as per s. 1(3).

E175 Previous affecting provision: power pursuant to section exercised (1.03.2004) by Misuse of Drugs (Scheduled Substances) (Amendment) Regulations 2004 (S.I. No. 92 of 2004); revoked (22.12.2009) by European Communities (Control of Drug Precursors) Regulations 2009 (S.I. No. 558 of 2009), reg. 37(1)(c).

E176 Previous affecting provision: power pursuant to section exercised (4.02.2003) by Misuse of Drugs (Scheduled Substances (Amendment) Regulations 2003 (S.I. No. 41 of 2003); revoked (1.03.2004) by Misuse of Drugs (Scheduled Substances) (Amendment) Regulations 2004 (S.I. No. 92 of 2004), reg. 2.


E178 Previous affecting provision: power pursuant to section exercised (8.12.1988) by Misuse of Drugs Regulations 1988 (S.I. No. 328 of 1988). Note that this SI was given statutory status (12.03.2015) by Misuse Of Drugs (Amendment) Act 2015 (6/2015), s. 2 and sch. 2, commenced as per s. 1(3). It was revoked (4.05.2017) by Misuse of Drugs (Amendment) Act 2016 (9/2016), s. 7(d), S.I. No. 172 of 2017.


E180 Previous affecting provision: power pursuant to section exercised (1.05.1979) by Misuse of Drugs Regulations 1979 (S.I. No. 32 of 1979), in effect as per reg. 2; revoked (8.12.1988) by Misuse of Drugs Regulations 1988 (S.I. No. 328 of 1988), reg. 2.

E181 Previous affecting provision: power pursuant to section exercised (1.05.1979) by Misuse of Drugs (Committees of Inquiry, Advisory Committees and Advisory Panels) Regulations 1979 (S.I. No. 31 of 1979), in effect as per reg. 2; revoked (23.10.1984) by Misuse of Drugs (Committees of Inquiry) Regulations 1984 (S.I. No. 264 of 1984), reg. 8.

Expenses.

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.

Collection and disposal of moneys payable under Act.

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.
3. The functions conferred on the Minister for Finance by or under the provisions of —
   (a) the enactments specified in Schedule 1, and
   (b) the statutory instruments specified in Schedule 2,
   are transferred to the Minister for Public Expenditure and Reform.

5. References to the Minister for Finance contained in any Act or instrument under an Act and
   relating to any functions transferred by this Order shall, from the commencement of this Order,
   be construed as references to the Minister for Public Expenditure and Reform.

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### Schedule 1

#### Enactments

<table>
<thead>
<tr>
<th>Number and Year</th>
<th>Short Title</th>
<th>Provision</th>
</tr>
</thead>
<tbody>
<tr>
<td>(1)</td>
<td>(2)</td>
<td>(3)</td>
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<tr>
<th>No. 12 of 1977</th>
<th>Misuse of Drugs Act 1977</th>
<th>Sections 38 and 40</th>
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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.

Annotations

Editorial Notes:


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.

Power pursuant to section exercised (1.03.1979 and 1.05.1979) by Misuse of Drugs Act, 1977 (Commencement) Order 1979 (S.I. No. 28 of 1979).

3. Sections 1, 2 and 4 to 14 (inclusive), section 21 (1), section 27 (2), subsections (1), (2), (6) and (7) of section 29, sections 32 to 40 (inclusive) and subsections (2), (3), (4), (5) and (6) of section 43 of the Act, together with the Schedule to the Act, shall come into operation generally on the 1st day of March, 1979.

4. Subsection (1) of section 42 of the Act shall come into operation—

(a) for the purpose of effecting the repeal of the matter referred to in paragraph (a) of that subsection, on the 1st day of March, 1979, and

(b) for the purpose of effecting the repeal of the matter referred to in paragraph (b) of that subsection, on the 1st day of May, 1979.

5. The provisions of the Act, other than those referred to in articles 3 and 4 of this Order, shall come into operation generally on the 1st day of May, 1979.
SCHEDULE

Controlled Drugs

1 Acetorphine.
Acetyldihydrocodeine.
Acetylmethadone.
Allylprodine.
Alphacetylmethadone.
Alphameprodine.
Alphamethadone.
Alphaprodine.
Amphetamine.
Amylobarbitone.
Anileridine.
Benzethidone.
Benzphetamine.
Benzylmorphine (3-benzylmorphine).
Betacetylmethadone.
Betameprodine.
Betamethadone.
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.
Diampromide.
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.
Ethylmethylthiambutene.
Ethylmorphine (3-ethylmorphine).
Etonitazene.
Etorphine.
Etoxeridine.
Fentanyl.
Furethidine.
Hydrocodone.
Hydromorphinol.
Hydromorphone.
Hydroxypethidine.
Isomethadone.
Ketobemidone.
Levomethorphan.
Levoramide.
Levophaenacylmorphan.
Levorphanol.
Lysergamide.
Lysergide and other N-alkyl derivatives of lysergamide.
Mephentermine.
Mescaline.
Metazocine.
Methadone.
Methaqualone.
Methylamphetamine.
Methyldesorphine.
Methyldihydromorphine (6-methylidihydromorphine).
Methylphenidate.
Metopon.
Morpheridine.
Morphine.
Morphine methobromide, morphine N-oxide and other pentavalent nitrogen morphine derivatives.
Myrophine.
Nicocodine.
Nicodicide (6-nicotinoyldihydrocodeine).
Nicomorphine.
Noracymethadol.
Norcodeine.
Norlevorphanol.
Normethadone.
Normorphine.
Norpipanone.
Opium, whether raw, prepared or medicinal.
Oxycodone.
Oxymorphone.
Pentobarbitone.
Pethidine.
Phenadoxone.
Phenampromide.
Phenzocine.
Phendimetrazine.
Phenmetrazine.
Phenobarbitone.
Phenomorph.
Phenoperidine.
Pholcodine.
Pimnidine.
Pipradrol.
Piritramide.
Poppy straw and concentrate of poppy straw.
Proheptazine.
Properidine.
Propiram.
Psilocin.
Quinalbarbitone.
Racemethorphan.
Racemoramide.
Racemorph.
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.

F66[F67][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-][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-
benzoxazin-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
MetH[QS.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-(cyclohexylmethyl) 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-enolate 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-1,2,3,4,6,7,12b-octahydroindolo[2,3-a]quinolizin-2-yl]-3-methoxyprop-2-enolate 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-Phenylclohexylamine
4-(1-Phenylclohexyl) morpholine
1-Piperidinoclohexanecarbonitrile
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-benzoyl)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-morpholiny1)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.
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.

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-morpholinyl)ethyl, whether or not further substituted in the cyclohexyl ring to any extent.

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-morpholinyl)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.

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-morpholinyl)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.

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-morpholinyl)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.

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.

Any substance (not being methoxyphenamine) structurally derived from phenethylamine, an \( N \)-alkyl-phenethylamine, \( \alpha \)-methylphenethylamine, an
$N$-alkyl-$\alpha$-methylphenethylamine, $\alpha$-ethylphenethylamine, or an $N$-alkyl-$\alpha$-ethylphenethylamine by substitution in the ring to any extent with alkyl, alkoxy, alklyenedioxy 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-dihydronaphthalen-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-dihydronaphthalen-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, alklyenedioxy, 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.]
Clotiazepam
Cloxazolam
Delorazepam
Dextropropoxyphene
Diazepam
Diethylpropion
Dihydrofuran-2(3H)-one
Estazolam
Ethchlorvynol
Ethinamate
N-Ethylamphetamine
Ethyl loflazepate
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
Nimetrazepam
Nitrazepam
Nordazepam
Oxazepam
Oxazolam
Pemoline
Pentazocine
Phenazepam
Phentermine
Pinazepam
Prazepam
Propylhexedrine
Pyrovalerone
Selegiline
Temazepam
Tetrazepam
Triazolam
Zaleplon
Zipeprrol
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 of this Schedule not being dextromethorphan or dextorphan.

F70[2A. Any stereoisomeric form of a substance specified in paragraph 1B of this Schedule.

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

4. Any salt of a substance or product specified in paragraph 1 of this Schedule.
5. Any preparation or product containing any proportion of a substance or product specified in F73[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” 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.

**Annotations**

**Amendments:**

- **F66** Inserted (12.03.2015) by *Misuse Of Drugs (Amendment) Act 2015* (6/2015), s. 1(a) and sch. 1, commenced as per s. 3(3).

- **F67** Substituted (4.05.2017) by *Misuse of Drugs (Amendment) Act 2016* (9/2016), s. 6(b) and sch., S.I. No. 172 of 2017.

- **F68** Inserted (12.05.2015) by *Misuse of Drugs (Amendment) Act 2015* (6/2015), s. 1(a) and sch. 1, commenced on enactment.

- **F69** Inserted (12.03.2015) by *Misuse Of Drugs (Amendment) Act 2015* (6/2015), s. 1(b), commenced as per s. 3(3).

- **F70** Inserted (12.03.2015) by *Misuse Of Drugs (Amendment) Act 2015* (6/2015), s. 1(c), commenced as per s. 3(3).

- **F71** Inserted (12.03.2015) by *Misuse Of Drugs (Amendment) Act 2015* (6/2015), s. 1(d), commenced as per s. 3(3).

- **F72** Inserted (12.03.2015) by *Misuse Of Drugs (Amendment) Act 2015* (6/2015), s. 1(e), commenced as per s. 3(3).

- **F73** Inserted (12.03.2015) by *Misuse Of Drugs (Amendment) Act 2015* (6/2015), s. 1(f), commenced as per s. 3(3).