
STATUTORY INSTRUMENTS

1997 No. 1830

The Prescription Only Medicines (Human Use) Order 1997

Citation, commencement and interpretation

1.—(1) This Order may be cited as the Prescription Only Medicines (Human Use) Order 1997 and shall come into force on 18th August 1997.

(2) In this Order, unless the context otherwise requires—

“the Act” means the Medicines Act 1968;

[^{F1}“additional supply optometrist” means a person who is registered as an optometrist, and against whose name particulars of the additional supply speciality have been entered in the relevant register;]

“aerosol” means a product which is dispersed from its container by a propellant gas or liquid;

[^{F2}“clinical management plan” means a written plan (which may be amended from time to time) relating to the treatment of an individual patient agreed by—

- (a) the patient to whom the plan relates,
- (b) the doctor or dentist who is a party to the plan, and
- (c) any supplementary prescriber who is to prescribe, give directions for administration or administer under the plan;]

[^{F3}“clinical trial” has the meaning given by regulation 2(1) of the Medicines for Human Use (Clinical Trials) Regulations 2003;]

^{F4} ...

[^{F5}“Common Services Agency” means the Common Services Agency for the Scottish Health Service established under section 10 of the National Health Service (Scotland) Act 1978;]

[^{F5}“Community marketing authorization” means a marketing authorization granted by the European Community under Council Regulation (EEC) No. 2309/93^{F6} or Regulation (EC) No. 726/2004 of the European Parliament and of the Council laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency;]

[^{F7}“community practitioner nurse prescriber” means a person—

- (a) who is a registered nurse or a registered midwife, and
- (b) against whose name is recorded in the professional register an annotation signifying that he is qualified to order drugs, medicines and appliances from the Nurse Prescribers' Formulary for Community Practitioners in the current edition of the British National Formulary;]

“controlled drug” has the meaning assigned to it by section 2 of the Misuse of Drugs Act 1971(1);

“cyanogenetic substances” means preparations which—

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Changes to legislation: There are currently no known outstanding effects for the The Prescription Only Medicines (Human Use) Order 1997, Section 1. (See end of Document for details)

- (a) are presented for sale or supply under the name of, or as containing, amygdalin, laetrile or vitamin B17, or
- (b) contain more than 0.1 per cent by weight of any substance having the formula either α -Cyanobenzyl-6-O- β -d-glucopyranosyl- β -d-glucopyranoside or α -Cyanobenzyl- β -d-glucopy ranosiduronic acid;

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“dosage unit” means—

- (a) where a medicinal product is in the form of a tablet or capsule or is an article in some other similar pharmaceutical form, that tablet, capsule or other article, or
- (b) where a medicinal product is not in any such form, the unit of measurement which is used as the unit by reference to which the dose of the medicinal product is measured;

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“external use” means application to the skin, hair, teeth, mucosa of the mouth, throat, nose, ear, eye, vagina or anal canal when a local action only is intended and extensive systemic absorption is unlikely to occur; and references to medicinal products for external use shall be read accordingly except that such references shall not include throat sprays, throat pastilles, throat lozenges, throat tablets, nasal drops, nasal sprays, nasal inhalations or teething preparations;

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[^{F5}“Health Authority”—

- (a) in relation to England and Wales, has the same meaning as in the National Health Service Act 1977;
- (b) in relation to Scotland, means a Health Board constituted under section 2 of the National Health Service (Scotland) Act 1978; and
- (c) in relation to Northern Ireland, means a Health and Social Services Board established under Article 16 of the Health and Personal Social Services (Northern Ireland) Order 1972;]

[^{F12}“health care” means services for or in connection with the prevention, diagnosis or treatment of disease;]

“health prescription” means a prescription issued by a doctor, dentist [^{F13}, supplementary prescriber] , [^{F14}a community practitioner nurse prescriber, a nurse independent prescriber or a pharmacist independent prescriber] under or by virtue of—

- (a) in England and Wales, the National Health Service Act 1977(2),
- (b) in Scotland, the National Health Service (Scotland) Act 1978(3), and
- (c) in Northern Ireland, the Health and Personal Social Services (Northern Ireland) Order 1972(4);

[^{F15}“health record” has the meaning given by section 68(2) of the Data Protection Act 1998(4);]

[^{F5}“homoeopathic certificate of registration” means a certificate of registration for the purposes of the Medicines (Homoeopathic Medicinal Products For Human Use) Regulations 1994;]

[^{F16}“independent clinic”—

(2) 1977 c. 49.
(3) 1978 c. 29.
(4) S.I. 1972/1265 (N.I. 14).
(4) S.I. 1972/1265 (N.I. 14).

- (a) in relation to England and Wales, has the meaning given by section 2(4) of the Care Standards Act 2000⁽⁶⁾, and
- (b) in relation to Scotland, has the meaning given by section 77(1) of the Regulation of Care (Scotland) Act 2001;]

[^{F16}“independent hospital”—

- (a) in relation to England and Wales, shall be construed in accordance with section 2(2), (3) and (6) of the Care Standards Act 2000, and
- (b) in relation to Scotland, means—
 - (i) an independent hospital, or
 - (ii) a private psychiatric hospital,as defined by section 77(1) of the Regulation of Care (Scotland) Act 2001;]

[^{F16}“independent medical agency”—

- (a) in relation to England and Wales, has the meaning given by section 2(5) of the Care Standards Act 2000, and
- (b) in relation to Scotland, has the meaning given by section 77(1) of the Regulation of Care (Scotland) Act 2001;]

“inhaler” does not include an aerosol;

[^{F17}“IRME practitioner” means, in relation to a medical exposure, a practitioner for the purposes of the Ionising Radiation (Medical Exposure) Regulations 2000⁽²⁾;]

[^{F18}“Local Health Board” has the same meaning as in the National Health Service Act 1977;]

[^{F5}“marketing authorization” includes a reference both to a United Kingdom marketing authorization and to a Community marketing authorization;]

“master” has the same meaning as in section 313(1) of the Merchant Shipping Act 1995⁽⁵⁾;

“maximum daily dose” or “MDD” means the maximum quantity of a substance contained in the amount of a medicinal product which it is recommended should be taken or administered in a period of 24 hours;

“maximum dose” or “MD” means the maximum quantity of a substance contained in the amount of a medicinal product which it is recommended should be taken or administered at any one time;

“maximum strength” means—

- (a) the maximum quantity of a substance by weight or volume contained in a dosage unit of a medicinal product;
- (b) the maximum percentage of a substance contained in a medicinal product calculated in any of the following ways—
 - (i) weight in weight,
 - (ii) weight in volume,
 - (iii) volume in weight, or
 - (iv) volume in volume,

and if the maximum percentage calculated in those ways differs, the higher or highest such percentage;

(6) S.I. 1985/2066.
(2) 1977 c. 49.
(5) 1995 c. 21.

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[^{F19}“medical exposure” has the same meaning as in the Ionising Radiation (Medical Exposure) Regulations 2000;]

“medicinal product” includes any article or substance in respect of which section 58 of the Act has effect by virtue of an order made under section 104 of the Act^{F20} ...

“the Misuse of Drugs Regulations” means, in relation to England, Wales and Scotland, the Misuse of Drugs Regulations 1985(6) and in relation to Northern Ireland, the Misuse of Drugs (Northern Ireland) Regulations 1986(7);

[^{F5}“NHS trust”—

- (a) in relation to England and Wales, has the same meaning as in the National Health Service and Community Care Act 1990;
- (b) in relation to Scotland, has the same meaning as in the National Health Service (Scotland) Act 1978; and
- (c) in relation to Northern Ireland, means a Health and Social Services trust established under article 10 of the Health and Social Services (Northern Ireland) Order 1991;]

[^{F21}“nurse independent prescriber” means a person—

- (a) who is a registered nurse or a registered midwife, and
- (b) against whose name is recorded in the professional register an annotation signifying that he is qualified to order drugs, medicines and appliances as a nurse independent prescriber or a nurse independent/supplementary prescriber;]

[^{F22}“NHS foundation trust” has the same meaning as in section 1(1) of the Health and Social Care (Community Health and Standards) Act 2003;]

[^{F23}“nursing home” has the meaning given by article 16 of the Registered Homes (Northern Ireland) Order 1992;]

“occupational health scheme” means a scheme in which a person, in the course of a business carried on by him, provides facilities for his employees for the treatment or prevention of disease;

“offshore installation” means an offshore installation within the meaning of the Mineral Workings (Offshore Installations) Act 1971(8) which is within—

- (a) tidal waters and parts of the sea in or adjacent to the United Kingdom up to the seaward limits of territorial waters;
- (b) waters in any area designated under section 1(7) of the Continental Shelf Act 1964(9);

[^{F24}“operator”—

- (a) in relation to an aircraft, means the person for the time being having management of the aircraft, and
- (b) for the purposes of article 7B, has the same meaning as in the Ionising Radiation (Medical Exposure) Regulations 2000;]

“parenteral administration” means administration by breach of the skin or mucous membrane;

[^{F5}“Patient Group Direction” means—

(6) S.I. 1985/2066.

(7) SR 1986 No. 52.

(8) 1971 c. 61; section 1 was substituted by section 24 of the Oil and Gas (Enterprise) Act 1982 (c. 23).

(9) 1964 c. 29.

- (a) in connection with the supply of a prescription only medicine as referred to in article 12A(2), [F25 12B, 12C, 12D or 12E], a written direction relating to the supply and administration of a description or class of prescription only medicine; or
- (b) in connection with the administration of a prescription only medicine as referred to in article 12A(2), 12B or 12C, a written direction relating to the administration of a description or class of prescription only medicine, and which, in the case of either (a) or (b)—
 - (i) is signed by a doctor or dentist, and by a pharmacist; and
 - (ii) relates to supply and administration, or to administration, to persons generally (subject to any exclusions which may be set out in the Direction);]

[F26 “pharmacist independent prescriber” means a person—

- (a) who is a pharmacist, and
- (b) against whose name is recorded in the relevant register an annotation signifying that he is qualified to order drugs, medicines and appliances as a pharmacist independent prescriber;]

“prescription only medicine” means a medicinal product of a description or falling within a class specified in article 3 of this Order;

[F5 “Primary Care Trust” has the same meaning as in the National Health Service Act 1977;]

[F27 “prison service” means—

- (a) in relation to England and Wales, a Minister of the Crown exercising functions in relation to prisons (within the meaning of the Prison Act 1952),
- (b) in relation to Scotland, the Scottish Ministers exercising functions in relation to prisons (within the meaning of the Prisons (Scotland) Act 1989), and
- (c) in relation to Northern Ireland, the Northern Ireland Department exercising functions in relation to prisons (within the meaning of the Prison Act (Northern Ireland) 1953);]

[F28 “professional register” means the register maintained by the Nursing and Midwifery Council [F29 under article 5 of] the Nursing and Midwifery Order 2001;]

“prolonged release” in relation to a medicinal product means a formulation of that product which—

- (a) is used to reduce the rate at which the active ingredient in that product is released after administration, and
- (b) is sold or supplied as a prolonged, controlled or sustained release medicinal product;

[F30 “radioactive medicinal product” means a medicinal product which is, which contains or which generates a radioactive substance and which is, contains or generates that substance in order, when administered, to utilize the radiation emitted therefrom;]

[F31 “registered chiropodist” means a person who is registered in Part 2 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;];

[F32 “registered dietitian” means a person who is registered in Part 4 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;]

[F33 “registered midwife” means a person registered in the Midwives' Part of the professional register;]

[F34 “registered nurse” means a person registered in the Nurses' Part [F35 or Specialist Community Public Health Nurses' Part] of the professional register;]

[^{F32}“registered occupational therapist” means a person who is registered in Part 6 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;]

[^{F36}“registered optometrist” means a person whose name is registered in the register of optometrists maintained under section 7(a) of the Opticians Act 1989;]

[^{F31}“registered orthoptist” means a person who is registered in Part 7 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;]

[^{F32}“registered orthotist and prosthetist” means a person who is registered in Part 10 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;]

[^{F31}“registered paramedic” means a person who is registered in Part 8 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;]

[^{F31}“registered physiotherapist” means a person who is registered in Part 9 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;]

[^{F37}“registered provider” means—

- (a) in relation to an independent hospital, an independent clinic or an independent medical agency—
 - (i) in relation to England and Wales, the person who is registered under Part II of the Care Standards Act 2000 as the person carrying on the establishment or agency,
 - (ii) in relation to Scotland, the person who is registered under Part 1 of the Regulation of Care (Scotland) Act 2001 as the person providing the establishment or agency, and
- (b) in relation to a nursing home, the person registered under Part III of the Registered Homes (Northern Ireland) Order 1992 as the person carrying on the nursing home, other than a manager who is to be treated as carrying on the home by virtue of article 17(2) of that order;]

[^{F31}“registered radiographer” means a person who is registered in Part 11 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;]

[^{F32}“registered speech and language therapist” means a person who is registered in Part 12 of the register maintained by the Health Professions Council under article 5 of the Health Professions Order 2001;]

[^{F37}“relevant manager” means—

- (a) in relation to an independent hospital, an independent clinic or an independent medical agency—
 - (i) in relation to England and Wales—
 - (aa) a person who is registered under Part II of the Care Standards Act 2000 as the manager of the establishment or agency, but who is not the registered provider for that establishment or agency, or
 - (bb) if there is no such person, but the registered provider has appointed a person to manage the establishment or agency, that appointed person,
 - (ii) in relation to Scotland, a person, other than the registered provider, who was identified as an individual who is to manage the establishment or agency on

the application for registration of that establishment or agency under Part 1 of Regulation of Care (Scotland) Act 2001, and

- (b) in relation to a nursing home, the manager of the nursing home, unless they are the registered provider for that home;]

[^{F37c}“relevant register” means—

- (a) in relation to a [^{F38}registered] nurse [^{F39}or registered midwife], the professional register,
^{F40} ...
- (b) in relation to a pharmacist, [^{F41}Part 1 of the register maintained under article 10(1) of the Pharmacists and Pharmacy Technicians Order 2007] or the register maintained in pursuance of Articles 6 and 9 of the Pharmacy (Northern Ireland) Order 1976; ^{F42} ...
- (c) [^{F43}in relation to a person whose name is registered in the part of the register maintained by the Health Professions Council in pursuance of article 5 of the Health Professions Order 2001 relating to—
- (i) chiropodists and podiatrists;
 - (ii) physiotherapists; or
 - (iii) radiographers diagnostic or therapeutic
- that register [^{F44}; and]
- (d) [^{F45}in relation to a registered optometrist, the register of optometrists maintained under section 7(a) of the Opticians Act 1989;]

“repeatable prescription” means a prescription which contains a direction that it may be dispensed more than once;

“sell” means sell by retail as defined in section 131 of the Act and “sale” has a corresponding meaning;

“soap” means any compound of a fatty acid with an alkali or amine;

[^{F5}“Special Health Authority”—

- (a) in relation to England and Wales, has the same meaning as in the National Health Service Act 1977;
- (b) in relation to Scotland, means a Special Health Board constituted under section 2 of the National Health Service (Scotland) Act 1978; and
- (c) in relation to Northern Ireland, means a Special Health and Social Services Agency established under the Health and Personal Social Services (Special Agencies) (Northern Ireland) Order 1990;]

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[^{F47c}“Strategic Health Authority” means a Strategic Health Authority established under section 8 of the National Health Service Act 1977;]

[^{F48c}“supplementary prescriber” means—

- (a) a [^{F49}registered] nurse, ^{F50} ...
- (b) a pharmacist, ^{F51} [^{F52} ...
- (c) a registered midwife,]^{F53} ...

- (d) [^{F54}a person whose name is registered in the part of the register maintained by the Health Professions Council in pursuance of article 5 of the Health Professions Order 2001 relating to—
- (i) chiropodists and podiatrists;
 - (ii) physiotherapists; or
 - (iii) radiographers diagnostic or therapeutic [^{F55}or]]

- (e) [^{F56}a registered optometrist,]

against whose name is recorded in the relevant register an annotation [^{F57}or entry] signifying that he is qualified to order drugs, medicines and appliances as a supplementary prescriber [^{F58}or, in the case of a nurse or midwife, as a nurse independent/supplementary prescriber];]

“supply” means supply in circumstances corresponding to retail sale as defined in section 131 of the Act;

“unit preparation” means a preparation, including a mother tincture, prepared by a process of solution, extraction or trituration with a view to being diluted tenfold or one hundredfold, either once or repeatedly, in an inert diluent, and then used either in this diluted form or, where applicable, by impregnating tablets, granules, powders or other inert substances.

[^{F59}“United Kingdom marketing authorization” means a marketing authorization granted by the licensing authority under the Medicines For Human Use (Marketing Authorisations Etc.) Regulations 1994 (including a product licence having effect as such an authorization by virtue of Schedule 6 to those Regulations [^{F59}or an authorization granted by the licensing authority in accordance with Article 126a of Directive 2001/83/EC].]

(3) For the purposes of this Order, the equivalence of a substance to a reference material shall be determined by calculating the amount of that reference material which is contained in that substance either by weight or, where the amount of the reference material is specified in terms of international units of activity, those units.

- (4) In this Order, unless the context otherwise requires, a reference—

- (a) to a numbered section is to the section of the Act which bears that number,
- (b) to a numbered article or Schedule is to the article of, or Schedule to, this Order which bears that number,
- (c) in an article or in a Part of a Schedule to a numbered paragraph is to the paragraph of that article or Part of that Schedule which bears that number, and
- (d) in a paragraph to a lettered sub-paragraph is to the sub-paragraph of that paragraph which bears that letter.

- (5) In [^{F60}Schedules 1, 2, 3A and 5]—

- (a) entries specified in columns 2 to 5 relate to the substances listed in column 1 against which they appear and where, in relation to a particular substance listed in column 1, an entry in columns 2 to 5 bears a number or letter it relates only to such entries in the other of those columns as bear the same number or letter;

- (b) the following abbreviations are used:

“g” for gram,
“iu” for international unit of activity,
“mcg” for microgram,
“mg” for milligram,
“ml” for millilitre.

(6) In Schedule 3, the abbreviation “NPF” means the Nurse Prescribers' Formulary Appendix in the British National Formulary.

[
F61(7) In articles 12 to [F6212E], a reference to a prescription only medicine being sold or supplied for the purpose of being administered in accordance with the written directions of a doctor or dentist relating to a particular person, or in accordance with a Patient Group Direction, includes a reference to it being sold or supplied in accordance with such directions or such a Direction.

(8) In articles 12A and 12C, a reference to an arrangement for the supply or administration of prescription only medicines includes a reference to an arrangement which covers such supply or administration and other matters.

(9) In Schedule 7, Part I, a reference to treatment of a clinical situation includes a reference to any form of management of that situation.]]

Textual Amendments

- F1 Words in art. 1(2) inserted (30.6.2005) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2005 (S.I. 2005/1507), arts. 1(1), **3(a)**
- F2 Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(a)**
- F3 Words in art. 1(2) substituted (1.5.2004) by The Medicines for Human Use (Clinical Trials) Regulations 2004 (S.I. 2004/1031), reg. 1, **Sch. 10 Pt. 2 para. 13(a)**
- F4 Words in art. 1 omitted (1.4.2002) by virtue of The Prescription Only Medicines (Human Use) Amendment Order 2002 (S.I. 2002/549), arts. 1(1), **2(2)(a)**
- F5 Words in art. 1(2) inserted (9.8.2000) by The Prescription Only Medicines (Human Use) Amendment Order 2000 (S.I. 2000/1917), arts. 1(1), **2(a)**
- F6 Words in art. 1(2) inserted (20.11.2005) by The Medicines (Marketing Authorisations Etc.) Amendment Regulations 2005 (S.I. 2005/2759), reg. 1(b), **Sch. para. 5(a)**
- F7 Words in art. 1(2) inserted (1.5.2006) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(a)**
- F8 Words in art. 1(2) omitted (1.5.2006) by virtue of The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(b)**
- F9 Words in art. 1(2) omitted (1.5.2006) by virtue of The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(c)**
- F10 Words in art. 1(2) omitted (1.5.2006) by virtue of The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(d)**
- F11 Words in art. 1(2) omitted (1.5.2006) by virtue of The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(e)**
- F12 Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(d)**
- F13 Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(e)**
- F14 Words in art. 1(2) substituted (1.5.2006) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(f)**
- F15 Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(f)**
- F16 Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(g)**
- F17 Words in art. 1(2) inserted (17.11.2006) by The Medicines for Human Use (Administration and Sale or Supply) (Miscellaneous Amendments) Order 2006 (S.I. 2006/2807), arts. 1(1), **2(a)**
- F18 Words in art. 1(2) inserted (19.11.2004) by The Prescription Only Medicines (Human Use) Amendment (No. 3) Order 2004 (S.I. 2004/2693), arts. 1(1), **2**

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- F19** Words in art. 1(2) inserted (17.11.2006) by The Medicines for Human Use (Administration and Sale or Supply) (Miscellaneous Amendments) Order 2006 (S.I. 2006/2807), arts. 1(1), **2(b)**
- F20** Words in art. 1(2) omitted (17.11.2006) by virtue of The Medicines for Human Use (Administration and Sale or Supply) (Miscellaneous Amendments) Order 2006 (S.I. 2006/2807), arts. 1(1), **2(c)**
- F21** Words in art. 1(2) inserted (1.5.2006) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(g)**
- F22** Words in art. 1(2) inserted (1.4.2004) by The Health and Social Care (Community Health and Standards) Act 2003 (Supplementary and Consequential Provision) (NHS Foundation Trusts) Order 2004 (S.I. 2004/696), arts. 1(1)(b), 3(8), **Sch. 8**
- F23** Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(h)**
- F24** Words in art. 1(2) substituted (17.11.2006) by The Medicines for Human Use (Administration and Sale or Supply) (Miscellaneous Amendments) Order 2006 (S.I. 2006/2807), arts. 1(1), **2(d)**
- F25** Words in art. 1 substituted (10.12.2003) by The Prescription Only Medicines (Human Use) Amendment (No. 2) Order 2003 (S.I. 2003/2915), arts. 1(1), **2**
- F26** Words in art. 1(2) inserted (1.5.2006) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(h)**
- F27** Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(i)**
- F28** Words in art. 1 inserted (1.4.2002) by The Prescription Only Medicines (Human Use) Amendment Order 2002 (S.I. 2002/549), arts. 1(1), **2(2)(e)**
- F29** Words in art. 1(2) substituted (1.8.2004) by The Health Act 1999 (Consequential Amendments) (Nursing and Midwifery) Order 2004 (S.I. 2004/1771), art. 1(1), **Sch. para. 39(a)(iv)**
- F30** Words in art. 1(2) inserted (17.11.2006) by The Medicines for Human Use (Administration and Sale or Supply) (Miscellaneous Amendments) Order 2006 (S.I. 2006/2807), arts. 1(1), **2(e)**
- F31** Words in art. 1 inserted (9.7.2003) by The Health Professions Order 2001 (Consequential Amendments) Order 2003 (S.I. 2003/1590), art. 1, **Sch. para. 21(2)(a)**
- F32** Words in art. 1(2) inserted (18.5.2004) by The Prescription Only Medicines (Human Use) Amendment (No. 2) Order 2004 (S.I. 2004/1189), arts. 1(1), **2**
- F33** Words in art. 1(2) substituted (1.8.2004) by The Health Act 1999 (Consequential Amendments) (Nursing and Midwifery) Order 2004 (S.I. 2004/1771), art. 1(1), **Sch. para. 39(a)(v)**
- F34** Words in art. 1(2) substituted (1.8.2004) by The Health Act 1999 (Consequential Amendments) (Nursing and Midwifery) Order 2004 (S.I. 2004/1771), art. 1(1), **Sch. para. 39(a)(vi)**
- F35** Words in art. 1(2) inserted (1.5.2006) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(i)**
- F36** Words in reg. 1(2) substituted (30.6.2005 as notified in the London Gazette dated 3.6.2005) by The Opticians Act 1989 (Amendment) Order 2005 (S.I. 2005/848), **Sch. 1 para. 27(a)**
- F37** Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(j)**
- F38** Word in art. 1(2) substituted (1.5.2006) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(j)**
- F39** Words in art. 1(2) inserted (1.8.2004) by The Health Act 1999 (Consequential Amendments) (Nursing and Midwifery) Order 2004 (S.I. 2004/1771), art. 1(1), **Sch. para. 39(a)(vii)**
- F40** Word in art. 1(2) omitted (7.4.2005) by virtue of The Medicines for Human Use (Prescribing) Order 2005 (S.I. 2005/765), arts. 1(1), **3(a)(i)**
- F41** Words in art. 1(2) substituted (coming into force in accordance with art. 1(2)(3) of the amending S.I.) by The Pharmacists and Pharmacy Technicians Order 2007 (S.I. 2007/289), **Sch. 1 para. 20**
- F42** Word in art. 1(b) omitted (30.6.2005) by virtue of The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2005 (S.I. 2005/1507), arts. 1(1), **3(b)(i)**
- F43** Words in art. 1(2) inserted (7.4.2005) by The Medicines for Human Use (Prescribing) Order 2005 (S.I. 2005/765), arts. 1(1), **3(a)(iii)**
- F44** Word in art. 1(c) inserted (30.6.2005) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2005 (S.I. 2005/1507), arts. 1(1), **3(b)(ii)**

- F45** Words inserted (30.6.2005) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2005 (S.I. 2005/1507), arts. 1(1), **3(b)(iii)**
- F46** Words in art. 1 omitted (9.7.2003) by virtue of The Health Professions Order 2001 (Consequential Amendments) Order 2003 (S.I. 2003/1590), art. 1, **Sch. para. 21(2)(b)**
- F47** Words in art. 1(2) inserted (1.10.2002) by The National Health Service Reform and Health Care Professions Act 2002 (Supplementary, Consequential etc. Provisions) Regulations 2002 (S.I. 2002/2469), regs. 1, 7, **Sch. 4**
- F48** Words in art. 1(2) inserted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(2)(k)**
- F49** Word in art. 1(2) substituted (1.5.2006) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(k)(i)**
- F50** Word in art. 1(2) omitted (1.8.2004) by virtue of The Health Act 1999 (Consequential Amendments) (Nursing and Midwifery) Order 2004 (S.I. 2004/1771), art. 1(1), **Sch. para. 39(a)(viii)**
- F51** Word in art. 1(2) omitted (7.4.2005) by virtue of The Medicines for Human Use (Prescribing) Order 2005 (S.I. 2005/765), arts. 1(1), **3(b)(i)**
- F52** Words in art. 1(2) inserted (1.8.2004) by The Health Act 1999 (Consequential Amendments) (Nursing and Midwifery) Order 2004 (S.I. 2004/1771), art. 1(1), **Sch. para. 39(a)(viii)**
- F53** Word in art. 1(2)(c) omitted (30.6.2005) by virtue of The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2005 (S.I. 2005/1507), arts. 1(1), **3(c)(i)**
- F54** Words in art. 1(2) inserted (7.4.2005) by The Medicines for Human Use (Prescribing) Order 2005 (S.I. 2005/765), arts. 1(1), **3(b)(iii)**
- F55** Word in art. 1(2)(d) inserted (30.6.2005) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2005 (S.I. 2005/1507), arts. 1(1), **3(c)(ii)**
- F56** Art. 1(2)(e) inserted (30.6.2005) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2005 (S.I. 2005/1507), arts. 1(1), **3(c)(iii)**
- F57** Words in art. 1(2) inserted (7.4.2005) by The Medicines for Human Use (Prescribing) Order 2005 (S.I. 2005/765), arts. 1(1), **3(c)**
- F58** Words in art. 1(2) inserted (1.5.2006) by The Medicines for Human Use (Prescribing) (Miscellaneous Amendments) Order 2006 (S.I. 2006/915), arts. 1(1), **2(k)(ii)**
- F59** Words in art. 1(2) inserted (20.11.2005) by The Medicines (Marketing Authorisations Etc.) Amendment Regulations 2005 (S.I. 2005/2759), reg. 1(b), **Sch. para. 5(b)**
- F60** Words in art. 1(5) substituted (1.4.2002) by The Prescription Only Medicines (Human Use) Amendment Order 2002 (S.I. 2002/549), arts. 1(1), **2(3)**
- F61** Words in art. 1(7)-(9) inserted (9.8.2000) by The Prescription Only Medicines (Human Use) Amendment Order 2000 (S.I. 2000/1917), arts. 1(1), **2(b)**
- F62** Word in art. 1(7) substituted (4.4.2003) by The Prescription Only Medicines (Human Use) Amendment Order 2003 (S.I. 2003/696), arts. 1(1), **2(3)**

Status:

Point in time view as at 07/02/2007. This version of this provision has been superseded.

Changes to legislation:

There are currently no known outstanding effects for the The Prescription Only Medicines (Human Use) Order 1997, Section 1.