MEDICINES (PRESCRIPTION ONLY) (JERSEY) ORDER 1997

Official Consolidated Version

This is an official version of consolidated legislation compiled and issued under the authority of the Legislation (Jersey) Law 2021.

Showing the law from 12 August 2022 to Current

MEDICINES (PRESCRIPTION ONLY) (JERSEY) ORDER 1997

Contents

Article

Jersey

MEDICINES (PRESCRIPTION ONLY) (JERSEY) ORDER 1997[1]

THE HEALTH AND SOCIAL SERVICES COMMITTEE in pursuance of Articles 57 and 110 of the Medicines (Jersey) Law 1995, after consultation with the Medicines Advisory Council and having otherwise complied with Article 110 of the Law, orders as follows –

Commencement [see endnotes]

1 Interpretation

(1) In this Order, unless the context otherwise requires –

"aerosol" means a product that is dispersed from its container by a propellent gas or liquid;

"controlled drug" has the same meaning as it has in Article 3 of the Misuse of Drugs (Jersey) Law 1978;

"cyanogenetic substances" means preparations –

(a) that are presented for sale or supply under the name of, or as containing, amygdalin, laetrile or vitamin B17; or

(b) that contain more than 0.1% by weight of any substance having the formula α-cyanobenzyl-6-*O*-ß-D-glucopyranosyl-ß-D-glucopyranoside or α-cyanobenzyl-ß-D-glucopyranosiduronic acid;

"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; and

(b) where a medicinal product is not in any such form, the quantity of the product that is used as the unit by reference to which the dose is measured;

"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; but does not mean application by means of a throat spray, throat pastille, throat lozenge, throat tablet, nasal drop, nasal spray, nasal inhalation or teething preparation;

"health prescription" means a prescription described in Article 15(2) of the Health Insurance (Jersey) Law 1967;

"health record" has the same meaning as in the Data Protection (Jersey) Law 2018;

"inhaler" does not include an aerosol;

"Law" means the Medicines (Jersey) Law 1995;

"master" has the same meaning as it has in the Merchant Shipping Act 1894 of the United Kingdom;

"maximum daily dose" or "MDD" means, in relation to a substance contained in the amount of a medicinal product for internal use, the recommended maximum quantity to be taken or administered in a period of 24 hours;

"maximum dose" or "MD" means, in relation to a substance contained in the amount of a medicinal product for internal use, the recommended maximum quantity to be taken or administered at any one time;

"maximum strength" means such of the following as may be specified –

(a) the maximum quantity of a substance by weight or volume contained in a dosage unit of a medicinal product;

(b) the maximum number of units of activity contained in a dosage unit or a weight of a medicinal product; and

(c) the maximum percentage of a substance contained in a medicinal product calculated in terms of weight in weight, weight in volume, volume in weight or volume in volume, as appropriate;

"medicinal product" does not include a veterinary drug;

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

"operator", in relation to an aircraft, means the person for the time being having the management of the aircraft;

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

"prescription only medicine" means a medicinal product that is specified by this Order as a prescription only medicine;

"registered optometrist" has the same meaning as it has in Article 1(1) of the Opticians (Registration) (Jersey) Law 1962;

"repeatable prescription" means a prescription containing a direction that it shall or may be dispensed more than once;

"soaps" means any compounds of a fatty acid with an alkali or amine;

"state registered paramedic" means a person who is registered in the register established and maintained under section 60 and paragraph 1(a) of Schedule 3 to the Health Act 1999 of the United Kingdom;

"supplementary prescriber" means any of the following –

(a) a midwife or nurse, whose entry on the register established and maintained under article 5 of the Nursing and Midwifery Order 2001 of the United

Kingdom indicates that he or she is, or may act as, only a supplementary prescriber;

(b) an optometrist, whose entry on the register established and maintained under section 7(a) of the Opticians Act 1989 of the United Kingdom indicates that he or she is, or may act as, only a supplementary prescriber;

(c) a pharmacist, whose entry on the register established and maintained under article 19 of the Pharmacy Order 2010 of the United Kingdom indicates that he or she is, or may act as, only a supplementary prescriber;

(d) a physiotherapist, podiatrist or radiographer, whose entry on the register established and maintained under article 5 of the Health and Social Work Professions Order 2001 of the United Kingdom indicates that he or she is, or may act as, only a supplementary prescriber.

"unit preparation" means a preparation (including a mother tincture) that is 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 that diluted form or (where applicable) by impregnating tablets, granules, powders or other inert substances.[2]

(2) In this Order –

(a) in Schedules 1 and 2, the following abbreviations are used –

"g" for gram;

"iu" for international unit of activity;

"mcg" for microgram;

"mg" for milligram; and

"ml" for millilitre; and

(b) in Schedule 1 –

(i) entries in any of columns 2, 3 and 4 of Parts 1 and 2 relate only to the substances specified in column 1 against which they appear,

(ii) where, in relation to a particular substance specified in column 1, an entry in any of columns 2, 3 and 4 bears a number or letter, that entry relates only to entries in the other of columns 2, 3 and 4 that bear the same number or letter, and

(iii) the entries in column 4 of Part 1 shall be read subject to the note at the end of that Part.

(3) Without prejudice to Article 10 of the Interpretation (Jersey) Law 1954, every provision in the Medicines (Jersey) Law 1995 that relates in any other way to its interpretation shall also apply in the same way to this Order, unless the context otherwise requires.

2 Prescription only medicines

The following descriptions and classes of medicinal products are specified for the purposes of Article 57(1)(a) of the Law, and are accordingly prescription only medicines, namely –

(a) medicinal products that consist of or contain a substance specified in column 1 of Part 1 of Schedule 1 to this Order;

(b) medicinal products that are controlled drugs;

(c) medicinal products that are for parenteral administration, whether or not they fall within sub-paragraph (a) or (b) of this paragraph;

(d) medicinal products that –

(i) are not of a description and do not fall within a class specified in any of sub-paragraphs (a), (b) and (c) of this paragraph,

(ii) are of a description in respect of which the conditions in Article 58(1) of the Law are fulfilled, and

(iii) are products in respect of which a product licence is granted, after the commencement of this Order, containing a provision to the effect that the method of sale or supply of the medicinal product is to be only in accordance with a prescription given by a person who is an appropriate practitioner; and

(e) cyanogenetic substances, other than preparations for external use.[3]

3 Medicinal products that are not prescription only medicines

(1) Notwithstanding Article 2, a medicinal product shall not be a prescription only medicine by reason that it consists of or contains a substance specified in column 1 of Part 1 of Schedule 1, where –

(a) in relation to that substance there is an entry in any of columns 2, 3 and 4;

(b) the maximum strength in the product of that substance does not exceed the maximum strength (if any) specified in column 2; and

(c) the medicinal product is sold or supplied –

(i) if a pharmaceutical form or a route of administration is specified in column 3, in such pharmaceutical form, and for administration only by such route, as may be so specified,

(ii) if a use is specified in column 3, in a container or package labelled (in either case) to show a use so specified to which the medicinal product is to be put but no use not so specified,

(iii) if a maximum dose is specified in column 4, in a container or package labelled (in either case) to show a maximum dose not exceeding that specified, and

(iv) if a maximum daily dose is specified in column 4, in a container or package labelled (in either case) to show a maximum daily dose not exceeding that specified.

(2) Notwithstanding Article 2 of this Order, a medicinal product shall not be a prescription only medicine by reason that it is a controlled drug, where –

(a) it contains not more than one of the substances specified in column 1 of Part 2 of Schedule 1 to this Order and no other controlled drug;

(b) it contains that substance at a strength that does not exceed the maximum strength specified in column 2; and

(c) it is sold or supplied –

(i) in such pharmaceutical form as may be specified in column 3, and

(ii) in or from a container or package labelled (in either case) to show a maximum dose not exceeding that specified in column 4.

(4) Notwithstanding Article 2, a medicinal product specified in Part 3 or 4 of Schedule 1 shall not be a prescription only medicine.

4 New medicinal products

For the purposes of Article 58(2)(a) of the Law their duration shall be a period of 5 years.

5 Appropriate practitioner[4]

(1) For the purposes of Article 57(1), the following are appropriate practitioners –

(a) doctors;

(b) dentists;

(c) veterinary surgeons;

(d) nurse independent prescribers;

(e) optometrist independent prescribers, but only in relation to a medicinal product that is not a controlled drug and that is not for parenteral administration;

(f) paramedic independent prescribers, but only in relation to a medicinal product that does not contain a controlled drug other than –

(i) Codeine,

(ii) Fentanyl,

(iii) Midazolam, or

(iv) Morphine; and

(g) pharmacist independent prescribers;

(h) physiotherapist independent prescribers, but only in relation to a medicinal product that does not contain a controlled drug other than –

(i) Dihydrocodeine,

(ii) Fentanyl,

(iii) Morphine,

(iv) Oxycodone, or

(v) Temazepam;

(i) podiatrist independent prescribers, but only in relation to a medicinal product that does not contain a controlled drug other than Dihydrocodeine or Temazepam;

(j) supplementary prescribers;

(k) therapeutic radiographer independent prescribers, but only in relation to a medicinal product that does not contain a controlled drug other than –

(i) Codeine,

(ii) Fentanyl,

(iii) Midazolam,

(iv) Morphine,

(v) Oxycodone,

(vi) Temazepam, or

(vii) Tramadol.

(2) In this Article, a reference to the following occupations is taken to be a reference to a person registered in respect of that occupation under the Health Care (Registration) (Jersey) Law 1995 –

(a) nurse independent prescriber;

(b) optometrist independent prescriber;

(c) paramedic independent prescriber;

(d) pharmacist independent prescriber;

(e) physiotherapist independent prescriber;

(f) podiatrist independent prescriber;

(g) therapeutic radiographer independent prescriber.

6 Conditions for prescriptions relating to sale and supply[5]

(1) For the purposes of Article 57(2)(a) of the Law (read with paragraph (4) of that Article), a prescription only medicine shall not be taken to be sold or supplied in accordance with a prescription given by an appropriate practitioner unless the conditions in paragraph (2) of this Article are fulfilled.

(2) The conditions to which paragraph (1) refers are –

(a) the prescription shall be written in ink or otherwise so as to be indelible, unless it is a health prescription that is not for a controlled drug specified in any of Schedules 1, 2 and 3 to the Misuse of Drugs (General Provisions) (Jersey) Order 1989, in which case it may be written using carbon paper or similar material;

(b) the prescription shall be signed in ink, with his or her own name, by the practitioner giving it;

(c) the prescription shall contain the following information –

(i) the address of the practitioner giving it,

(ii) the appropriate date,

(iii) by virtue of which of the paragraphs of Article 5 the practitioner giving it is an appropriate practitioner,

(iv) where the practitioner giving it is an appropriate practitioner other than a veterinary surgeon, the name and address of the person for whose treatment it is given and (if that person is under 12) his or her age, and

(v) where the practitioner giving it is a veterinary surgeon, the name and address of the person to whom the prescription only medicine is to be delivered, and a declaration by that veterinary surgeon that the prescription only medicine is prescribed for an animal or herd under his or her care;

(ca) in the case of a prescription given by a supplementary prescriber, the supplementary prescriber –

(i) has given the prescription in accordance with the terms of a clinical management plan containing the information specified in Schedule 4, such plan relating to an individual patient and to which the following are parties –

(A) the patient,

(B) the patient's doctor or dentist, and

(C) the supplementary prescriber, and

(ii) has access to the health records of the patient to whom the clinical management plan relates to the extent that such records are used by the doctor or dentist who is a party to the plan;

(d) the prescription shall not be dispensed after the end of the period of 6 months from the appropriate date unless it is a repeatable prescription, in which case it shall not be dispensed for the first time after the end of that period or otherwise than in accordance with the direction contained in the repeatable prescription; and

(e) in the case of a repeatable prescription that does not specify the number of times that it may be dispensed, the prescription shall not be dispensed on more than 2 occasions unless it is a prescription for oral contraceptives, in which case it may be dispensed 6 times before the end of the period of 6 months from the appropriate date.[6]

(3) The restrictions in Article 57(2)(a) of the Law shall not apply to a sale or supply of a prescription only medicine that, by reason only that a condition in paragraph (2) of this Article is not fulfilled, is not in accordance with a prescription given by an appropriate practitioner, where the person selling or supplying the prescription only medicine, having exercised all due diligence, believes on reasonable grounds that that condition is fulfilled in relation to that sale or supply.

(4) In paragraph (2), the "appropriate date" means –

(a) in the case of a health prescription –

(i) the date on which it was signed by the practitioner by whom it was given, or

(ii) if it also contains a date indicated by him or her as being the date before which it shall not be dispensed, the later of the 2 dates; and

(b) in every other case, the date on which the prescription was signed by the practitioner by whom it was given.

6A Conditions for prescriptions – administration[7]

(1) For the purposes of Article 57(2)(b) of the Law (read with paragraph (4) of that Article), a prescription only medicine shall not be taken to be administered by a supplementary prescriber or by a person acting in accordance with the directions of a supplementary prescriber unless the conditions in paragraph (2) are met.

(2) Those conditions are that –

(a) the supplementary prescriber is acting in accordance with the terms of a clinical management plan containing the information specified in

Schedule 4, such plan relating to an individual patient to which the following are parties –

(i) the patient,

(ii) the patient's doctor or dentist, and

(iii) the supplementary prescriber; and

(b) the supplementary prescriber has access to the health records of the patient to whom the clinical management plan relates to the extent that such records are used by the doctor or dentist who is a party to the plan.

7 Exemption for highly diluted medicinal products

The restrictions in Article 57(2) of the Law shall not apply to the sale, supply or administration of a medicinal product that is not for parenteral administration and only consists of or only contains one or more of the substances specified in column 1 of Part 1 or 2 of Schedule 1 to this Order, where –

(a) each unit preparation has been diluted to at least one part in a million (6x), and the person selling, supplying or administering the medicinal product has been requested by or on behalf of a particular person and in that person's presence to use his or her own judgment as to the treatment required; or

(b) each such unit preparation has been diluted to at least one part in a million million (6c).

8 Exemptions for specified categories of persons

(1) The restrictions in Article 57(2)(a) of the Law shall not apply –

(a) to the sale or supply by a person specified in column 1 of Part 1 of Schedule 2 to this Order; or

(b) to the supply by a person specified in column 1 of Part 2 of Schedule 2 to this Order,

of a prescription only medicine specified in column 2 of Part 1 or 2 of that Schedule in relation to that person, where the conditions in the corresponding paragraph in column 3 of that Part are fulfilled.

(2) The restriction in Article 57(2)(b) of the Law shall not apply to the administration by a person specified in column 1 of Part 3 of Schedule 2 to this Order of a prescription only medicine for parenteral administration specified in column 2 of that Part in relation to that person, where the conditions in the corresponding paragraph in column 3 of that Part are fulfilled.

9 Exemption for emergency sale or supply

(1) The restrictions in Article 57(2)(a) of the Law shall not apply to the sale or supply of a prescription only medicine by a person who is lawfully conducting a retail pharmacy business, where the conditions in paragraph (2) of this Article or the alternative conditions in paragraph (3) of this Article are fulfilled.

(2) The conditions to which paragraph (1) of this Article refers are –

(a) the pharmacist by or under whose supervision the prescription only medicine is to be sold or supplied is satisfied that the sale or supply has been requested by a doctor who, by reason of any emergency, is unable to furnish a prescription immediately;

(b) the doctor has undertaken to furnish the person lawfully conducting the retail pharmacy business with a prescription within 72 hours;

(c) the prescription only medicine is sold or supplied in accordance with the directions of the doctor requesting it;

(d) the prescription only medicine is not a controlled drug specified in any of Schedules 1, 2 and 3 to the Misuse of Drugs (General Provisions) (Jersey) Order 1989; and

(e) an entry is made in the register to be kept under Article 3(1) of the Medicines (Sale and Supply) (Miscellaneous Provisions) (Jersey) Order 1997, within the appropriate time specified in that Article, of the information in paragraph 1 of Schedule 2 to that Order.

(3) The alternative conditions to which paragraph (1) of this Article refers are –

(a) the pharmacist by or under whose supervision the prescription only medicine is to be sold or supplied has interviewed the person requesting a prescription only medicine and is satisfied –

(i) that there is an immediate need for that prescription only medicine to be sold or supplied and that it is impracticable in the circumstances to obtain a prescription without undue delay,

(ii) that treatment with that prescription only medicine has been prescribed on a previous occasion by a doctor for the person requesting it from the pharmacist, or (as far as the pharmacist is reasonably able to ascertain) has been lawfully prescribed on a previous occasion by a medical practitioner outside Jersey for the person requesting it from the pharmacist, and

(iii) as to the dose which, in the circumstances, it would be appropriate for that person to take;

(b) the prescription only medicine –

(i) will be sold or supplied in no greater quantity than will provide 5 days' treatment,

(ii) is a preparation of insulin, an aerosol for the relief of asthma, an ointment or a cream, and has been made up for sale in a container elsewhere than at the place of sale or supply, and is the smallest pack that the pharmacist has available for sale or supply,

(iii) is an oral contraceptive and is sufficient, but no more than sufficient, for a full cycle, or

(iv) is an antibiotic for oral administration in liquid form, and is the smallest quantity that will provide a full course of treatment;

(c) the prescription only medicine does not consist of or contain a substance specified in Schedule 3 to this Order and is not a controlled drug specified in any of the Schedules to the Misuse of Drugs (General Provisions) (Jersey) Order 1989;

(d) an entry is made in the register to be kept under Article 3(1) of the Medicines (Sale and Supply) (Miscellaneous Provisions) (Jersey) Order 1997 within the appropriate time specified in that Article, of the information in paragraph 3 of Schedule 2 to that Order; and

(e) the container or package of the prescription only medicine is labelled so as to show –

(i) the date on which the prescription only medicine is sold or supplied,

(ii) the name, quantity and (unless it is apparent from the name) the pharmaceutical form and strength of the prescription only medicine,

(iii) the name of the person requesting the prescription only medicine,

(iv) the name and address of the registered pharmacy from which the prescription only medicine is sold or supplied, and

(v) the words "Emergency Supply".[8]

(4) The conditions in paragraph (2)(d) of this Article and in paragraph (3)(c) of this Article shall not apply where the prescription only medicine –

(a) consists of or contains phenobarbitone or phenobarbitone sodium (but no other substance specified in Schedule 3 to this Order or in any of the Schedules to the Misuse of Drugs (General Provisions) (Jersey) Order 1989); and

(b) is sold or supplied for use in the treatment of epilepsy.

10 Exemption for sale or supply in hospitals or the prison[9]

(1) The restrictions in Article 57(2)(a) of the Law shall not apply to the sale or supply of any prescription only medicine –

(a) in the course of the business of a hospital; or

(b) by a pharmacy to the prison under the terms of a contract to supply medicinal products for the benefit of prisoners,

in accordance with the written directions of an appropriate practitioner, other than a supplementary prescriber, even though those directions do not fulfil the conditions in Article 6(2) of this Order.

(2) In the case of directions given by a supplementary prescriber, paragraph (1) applies except that the condition in Article 6(2)(ca) must be fulfilled as if the references to a prescription in that sub-paragraph were references to the directions given by the supplementary prescriber.

11 Exemption for authorised needle supply services[10]

The restrictions of Article 57(2)(a) of the Law shall not apply to the supply by a person, for parenteral administration, of ampoules of sterile water, if the supply is made by the person in the course of acting on behalf of a service provided by or on behalf of the States for the purpose of enabling the supply of syringes, and associated articles, so as to reduce the spread of disease.

12 Exemption for sale or supply in cases involving another's default

The restrictions in Article 57(2)(a) of the Law shall not apply to the sale or supply of a prescription only medicine by a person who, having exercised all due diligence, believes on reasonable grounds that the product sold or supplied is not a prescription only medicine, and it is because of an act or default of another person that the product is a product to which that sub-paragraph applies.

13 Exemption in the case of forged prescription

The restrictions in Article 57(2)(a) of the Law shall not apply to the sale or supply of a prescription only medicine by a pharmacist in accordance with a forged prescription where the pharmacist, having exercised all due diligence, believes on reasonable grounds that the prescription is genuine.

14 Exemption for parenteral administration to human beings

The restriction in Article 57(2)(b) of the Law shall not apply to the administration to human beings of any of the following medicinal products for parenteral administration, namely –

adrenaline injection BP;

atropine sulphate injection;

chlorpheniramine injection;

cobalt edetate injection;

dextrose injection strong B.P.C.;

diphenhydramine injection;

glucagon injection;

hydrocortisone injection;

mepyramine injection;

naloxone injection;

promethazine hydrochloride injection;

snake venom antiserum;

sodium nitrite injection;

sodium thiosulphate injection; and

sterile pralidoxime injection,

where it is administered for the purpose of saving life in an emergency.

15 Exemption for non-parenteral administration to human beings

The restriction in Article 57(2)(b) of the Law shall not apply to the administration to human beings of a prescription only medicine that is not for parenteral administration.

16 Citation

This Order may be cited as the Medicines (Prescription Only) (Jersey) Order 1997.

SCHEDULE 1

(Articles 1(2), 2(a), 3(1) and 7)

PART 1[11]

PRESCRIPTION ONLY MEDICINES

[**Note –**

(x) indicates that the entry is to be read subject to paragraph 1 of the note at the end of Part 1 of Schedule 1

(y) indicates that the entry is to be read subject to paragraph 2 of the note at the end of Part 1 of Schedule 1]

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Column 1	*Column 2*	*Column 3*	*Column 4*
Substance	Maximum strength	Use, pharmaceutical form or route of administration	Maximum dose and maximum daily dose
Acamprosate			
Acarbose			
Acebutolol Hydrochloride			
Aceclofenac			
Acemetacin			
Acetarsol			
Acetazolamide			
Acetazolamide Sodium			
Acetohexamide			
Acetylcholine Chloride	0.2%	External	
Acetylcysteine			
Aciclovir			
Acipimox			
Acitretin			
Aclarubicin Hydrochloride			
Aconite	1.3%	External	
Acrivastine			
Acrosoxacin			
Actinomycin C			
Actinomycin D			
Adapalene			
Adenosine			
Adrenaline		(1) By inhaler (2) External	

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Adrenaline Acid Tartrate		(1) By inhaler (2) External	
Adrenaline Hydrochloride		(1) By inhaler (2) External	
Adrenocortical Extract			
Aclofenac			
Albendazole			
Alclometasone Dipropionate			
Alcuronium Chloride			
Aldesleukin			
Aldosterone			
Alendronate Sodium			
Alfacalcidol			
Alfuzosin Hydrochloride			
Allergen Extracts			
Allopurinol			
Allyloestrenol			
Aloxiprin			
Alphadolone Acetate			
Alphaxalone			
Alprenolol			
Alprenolol Hydrochloride			
Alprostadil			
Alseroxylon			
Altretamine			
Amantadine Hydrochloride			
Ambenonium Chloride			
Ambutonium Bromide			
Amcinonide			
Ametazole Hydrochloride			
Amethocaine		Any use (except local ophthalmic use)	
Amethocaine Gentisate		Any use (except local ophthalmic use)	
Amethocaine Hydrochloride		Any use (except local ophthalmic use)	
Amikacin Sulphate			
Amiloride Hydrochloride			
Aminocaproic Acid			
Aminoglutethimide			
Aminopterin Sodium			
Amiodarone Hydrochloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Amiphenazole Hydrochloride			
Amisulpride			
Amitriptyline			
Amitriptyline Embonate			
Amitriptyline Hydrochloride			
Amlodipine Besylate			
Ammonium Bromide			
Amodiaquine Hydrochloride			
Amorolfine Hydrochloride			
Amoxapine			
Amoxycillin			
Amoxycillin Sodium			
Amoxycillin Trihydrate			
Amphomycin Calcium			
Amphotericin			
Ampicillin			
Ampicillin Sodium			
Ampicillin Trihydrate			
Amsacrine			
Amygdalin			
Amyl Nitrite			
Amylocaine Hydrochloride		Any use (except local ophthalmic use)	
Anastrazole			
Ancrod			
Androsterone			
Angiotensin Amide			
Anistreplase			
Anterior Pituitary Extract			
Antimony Barium Tartrate			
Antimony Dimercaptosuccinate			
Antimony Lithium Thiomalate			
Antimony Pentasulphide			
Antimony Potassium Tartrate			
Antimony Sodium Tartrate			
Antimony Sodium Thiogycollate			
Antimony Sulphate			
Antimony Trichloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Antimony Trioxide			
Antimony Trisulphide			
Apiol			
Apomorphine			
Apomorphine Hydrochloride			
Apraclonidine Hydrochloride			
Aprotinin			
Arecoline Hydrobromide			
Argipressin			
Aristolochia			
Aristolochia Clematitis			
Aristolochia Contorta			
Aristolochia Debelis			
Aristolochia Fang-chi			
Aristolochia Manshuriensis			
Aristolochia Serpentaria			
Arsenic			
Arsenic Triiodide			
Arsenic Trioxide			
Arsphenamine			
Aspirin		Any form (except non-effervescent tablets or capsules)	
Astemizole			
Atenolol			
Atorvastatin			
Atorvastatin Calcium			
Atovaquone			
Atracurium Besylate			
Atropine		(1) Internal: (a) by inhaler (b) otherwise than by inhaler	(b) 300 mcg (MD) 1 mg (MDD)(x)
		(2) External (except local ophthalmic use)	
Atropine Methobromide		(1) Internal: (a) by inhaler (b) otherwise than by inhaler	(b) 400 mcg (MD) 1.3mg (MDD)(x)
		(2) External (except local ophthalmic use)	

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Atropine Methonitrate		Internal: (a) by inhaler (b) otherwise than by inhaler	(b) 400 mcg (MD) 1.3 mg (MDD)(x)
Atropine Oxide Hydrochloride		(1) Internal: (a) by inhaler (b) otherwise than by inhaler	(b) 360mcg (MD) 1.2mg (MDD)(x)
		(2) External (except local opthalmic use)	
Atropine Sulphate		(1) Internal: (a) by inhaler (b) otherwise than by inhaler	(b) 360 mcg (MD) 1.2 mg (MD)(x)
		(2) External (except local ophthalmic use)	
Auranofin			
Azapropazone			
Azathioprine			
Azathioprine Sodium			
Azelaic Acid			
Azelastine Hydrochloride			
Azidocillin Potassium			
Azithromycin			
Azlocillin Sodium			
Aztreonam			
Bacampicillin Hydrochloride			
Bacitracin			
Bacitracin Methylene Disalicylate			
Bacitracin Zinc			
Baclofen			
Balsalazide Sodium			
Bambuterol Hydrochloride			
Barium Carbonate			
Barium Chloride			
Barium Sulphide			
Beclamide			
Beclomethasone			
Beclomethasone Diproprionate			
Belladonna Herb		(1) Internal (2) External	(1) 1 mg of the alkaloids (MDD)

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Belladonna Root		(1) Internal (2) External	(1) 1 mg of the alkaloids (MDD)
Bemegride			
Bemegride Sodium			
Benapryzine Hydrochloride			
Bendrofluazide			
Benethamine Penicillin			
Benoxaprofen			
Benperidol			
Benserazide			
Benserazide Hydrochloride			
Bentiromide			
Benzathine Penicillin			
Benzbromarone			
Benzhexol Hydrochloride			
Benzilonium Bromide			
Benzocaine		Any use (except local ophthalmic use)	
Benzoctamine Hydrochloride			
Benzoyl Peroxide	10.0%	External	
N-Benzoyl Sulphanilamide			
Benzquinamide			
Benzquinamide Hydrochloride			
Benzthiazide			
Benztropine Mesylate			
Benzylpenicillin Calcium			
Benzylpenicillin Potassium			
Benzylpenicillin Sodium			
Beractant			
Betahistine Hydrochloride			
Betamethasone			
Betamethasone Adamantoate			
Betamethasone Benzoate			
Betamethasone Dipropionate			
Betamethasone Sodium Phosphate			
Betamethasone Valerate			
Betaxolol Hydrochloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Bethanechol Chloride			
Bethanidine Sulphate			
Bezafibrate			
Bicalutamide			
Biperiden Hydrochloride			
Biperiden Lactate			
Bismuth Glycollylarsanilate			
Bisoprolol Fumarate			
Bleomycin			
Bleomycin Sulphate			
Bretylium Tosylate			
Brimonidine Tartrate			
Bromhexine Hydrochloride			
Bromocriptine Mesylate			
Bromperidol			
Bromvaletone			
Brotizolam			
Budesonide			
Bufexamac			
Bumetanide			
Buphenine Hydrochloride			6 mg (MD) 18 mg (MDD)
Bupivacaine		Any use (except local ophthalmic use)	
Bupivacaine Hydrochloride		Any use (except local ophthalmic use)	
Buserelin Acetate			
Buspirone Hydrochloride			
Busulphan			
Butacaine Sulphate		Any use (except local ophthalmic use)	
Butorphenol Tartrate			
Butriptyline Hydrochloride			
Cabergoline			
Calcipotriol			
Calcipotriol Hydrate			
Calcitonin			
Calcitriol			
Calcium Amphomycin			
Calcium Benzamidosalicylate			
Calcium Bromide			

Prescription Only Medicine	Circumstances In Which Substances Are Not Prescription Only Medicines		
Calcium Bromidolactobionate			
Calcium Carbimide			
Calcium Folinate			
Calcium Metrizoate			
Calcium Sulphaloxate			
Candesartan Cilexetil			
Candicidin			
Canrenoic Acid			
Cantharidin	0.01%	External	
Capreomycin Sulphate			
Captopril			
Carbachol			
Carbamazepine			
Carbaryl			
Carbasalate Calcium			
Carbenicillin Sodium			
Carbenoxolone Sodium		(1) Pellet	(1) 5 mg (MD) 25 mg (MDD)
	(2) 2.0%	(2) Gel	
Carbidopa			
Carbimazole			
Carbocisteine			
Carbon Tetrachloride			
Carboplatin			
Carboprost Trometamol			
Carbuterol Hydrochloride			
Carfecillin Sodium			
Carindacillin Sodium			
Carisoprodol			
Carmustine			
Carperidine			
Carteolol Hydrochloride			
Cefaclor			
Cefadroxil			
Cefazedone Sodium			
Cefdinir			
Cefixime			
Cefodizime Sodium			
Cefotaxime Sodium			
Cefoxitin Sodium			
Cefpodoxime Proxetil			
Cefprozil			
Cefsulodin Sodium			
Ceftazidime			
Ceftizoxime Sodium			
Ceftriaxone Sodium			
Cefuroxime Axetil			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Cefuroxime Sodium			
Celiprolol Hydrochloride			
Cephalexin			
Cephalexin Sodium			
Cephaloridine			
Cephalothin Sodium			
Cephamandole Nafate			
Cephazolin Sodium			
Cephradine			
Cerium Oxalate			
Cerivastatin			
Cerivastatin Sodium			
Ceruletide Diethylamine			
Cetirizine			
Chenodeoxycholic Acid			
Chloral Hydrate		External	
Chlorambucil			
Chloramphenicol			
Chloramphenicol Cinnamate			
Chloramphenicol Palmitate			
Chloramphenicol Sodium Succinate			
Chlorhexadol			
Chlormadinone Acetate			
Chlormerodrin			
Chlormethiazole			
Chlormethiazole Edisylate			
Chlormezanone			
Chloroform	(1) 5.0%	(1) Internal (2) External	
Chloroquine Phosphate		Prophylaxis of malaria	
Chloroquine Sulphate		Prophylaxis of malaria	
Chlorothiazide			
Chlorotrianisene			
Chlorphenoxamine Hydrochloride			
Chlorpromazine			
Chlorpromazine Embonate			
Chlorpromazine Hydrochloride			
Chlorpropamide			
Chlorprothixene			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Chlorprothixene Hydrochloride			
Chlortetracycline			
Chlortetracycline Calcium			
Chlortetracycline Hydrochloride			
Chlorthalidone			
Chlorzoxazone			
Cholestyramine			
Chorionic Gonadotrophin			
Ciclacillin			
Ciclobendazole			
Cidofovir			
Cilastatin Sodium			
Cilazapril			
Cimetidine			
Cimetidine Hydrochloride			
Cinchocaine	3.0%	Any use (except local ophthalmic use)	
Cinchocaine Hydrochloride	Equivalent of 3.0% of Cinchocaine	Any use (except local ophthalmic use)	
Cinchophen			
Cinoxacin			
Ciprofibrate			
Ciprofloxacin			
Ciprofloxacin Hydrochloride			
Cisapride			
Cisplatin			
Citalopram Hydrobromide			
Clarithromycin			
Clavulanic Acid			
Clenbuterol Hydrochloride			
Clidinium Bromide			
Clindamycin			
Clindamycin Hydrochloride			
Clindamycin Palmitate Hydrochloride			
Clindamycin Phosphate			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Clioquinol	(1) 35 mg	(1) Treatment of mouth ulcers (2) External (except treatment of mouth ulcers)	(1) 350 mg (MDD)
Clobetasol Propionate			
Clobetasone Butyrate			
Clofazimine			
Clofibrate			
Clomiphene Citrate			
Clomipramine			
Clomipramine Hydrochloride			
Clomocycline			
Clomocycline Sodium			
Clonidine			
Clonidine Hydrochloride			
Clopamide			
Clopenthixol Decanoate			
Clopenthixol Hydrochloride			
Clorexolone			
Clostebol Acetate			
Clotrimazole		External but, in the case of vaginal use, only for the treatment of vaginal candidiasis	
Cloxacillin Benzathine			
Cloxacillin Sodium			
Clozapine			
Cocculus Indicus			
Co-dergocrine Mesylate			
Colaspase			
Colchicine			
Colestipol Hydrochloride			
Colfosceril Palmitate			
Colistin Sulphate			
Colistin Sulphomethate			
Colistin Sulphomethate Sodium			
Coniine			
Conium Leaf	7.0%	External	
Corticotrophin			
Cortisone			
Cortisone Acetate			
Co-tetroxazine			
Co-Trimoxazole			

Prescription Only Medicine	Circumstances In Which Substances Are Not Prescription Only Medicines		
Copropamide			
Crotethamide			
Croton Oil			
Croton Seed			
Curare			
Cyclofenil			
Cyclopenthiazide			
Cyclopentolate Hydrochloride			
Cyclophosphamide			
Cycloserine			
Cyclosporin			
Cyclothiazide			
Cyproterone Acetate			
Cytarabine			
Cytarabine Hydrochloride			
Dacarbazine			
Dalteparin Sodium			
Danazol			
Danthron			
Dantrolene Sodium			
Dapsone			
Dapsone Ethane Ortho Sulphonate			
Daunorubicin Hydrochloride			
Deanol Bitartrate			26 mg (MDD)
Debrisoquine Sulphate			
Demecarium Bromide			
Demeclocycline			
Demeclocycline Calcium			
Demeclocycline Hydrochloride			
Deoxycortone Acetate			
Deoxycortone Pivalate			
Deptotropine Citrate			
Dequalinium Chloride	(1) 0.25 mg	(1) Internal: throat lozenges or throat pastilles	
	(2) 1.0%	(2) External: paint	
Deserpidine			
Desferrioxamine Mesylate			
Desflurane			
Desfluorotriamcinolone			
Desipramine Hydrochloride			
Deslanoside			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Desmopressin			
Desogestrel			
Desonide			
Desoxymethasone			
Dexamethasone			
Dexamethasone Acetate			
Dexamethasone Isonicotinate			
Dexamethasone Phenylpropionate			
Dexamethaone Pivalate			
Dexamethasone Sodium *m*-Sulphobenzoate			
Dexamethasone Sodium Phosphate			
Dexamethasone Troxundate			
Dexfenfluramine Hydrochloride			
Dextromethorphan Hydrobromide		Internal	In the case of a controlled release preparation: equivalent of 30 mg of Dextromethor-phan (MD) equivalent of 75 mg of Dextromethor-phan (MDD)
			In any other case: equivalent of 15 mg of Dextromethor-phan (MD) equivalent of 75 mg of Dextromethor-phan (MDD)
Dextrothyroxine Sodium			
Diazoxide			
Dibenzepin Hydrochloride			
Dichloralphenazone			
Dichlorphenamide			
Diclofenac Diethylammonium			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Diclofenac Potassium			
Diclofenac Sodium			
Dicyclomine Hydrochloride			10 mg (MD) 60 mg (MDD)
Didanosine			
Dienoestrol			
Diethanolamine Fusidate			
Diflucortolone Valerate			
Diflunisal			
Digitalin			
Digitalis Leaf			
Digitalis, Prepared			
Digitoxin			
Digoxin			
Dihydralazine Sulphate			
Dihydroergotamine Mesylate			
Dihydrostreptomycin			
Dihydrostreptomycin Sulphate			
Diloxanide Furoate			
Diltiazem Hydrochloride			
Dimercaprol			
Dimethisoquin Hydrochloride		Any use (except local ophthalmic use)	
Dimethisterone			
Dimethothiazine Mesylate			
Dimethyl Sulphoxide			
Dimethyltubocurarine Bromide			
Dimethyltubocurarine Chloride			
Dimethyltubocurarine Iodide			
Dinoprost			
Dinoprost Trometamol			
Dinoprostone			
Diphenhydramine Hydrochloride		All preparations except liquid-filled capsules	
Dipivefrin Hydrochloride			
Dipyridamole			
Disodium Etidronate			
Disopyramide			
Disopyramide Phosphate			
Distigmine Bromide			
Disulfiram			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Dithranol	1.00%		
Dobutamine Hydrochloride			
Dolasetron Mesilate			
Domperidone			
Domperidone Maleate			
Donepezil			
Donepezil Hydrochloride			
Dopamine Hydrochloride			
Dopexamine Hydrochloride			
Dorzolamide Hydrochloride			
Dothiepin			
Dothiepin Hydrochloride			
Doxapram Hydrochloride			
Doxazosin Mesylate			
Doxepin Hydrochloride			
Doxorubicin			
Doxorubicin Hydrochloride			
Doxycycline			
Doxycycline Calcium Chelate			
Doxycycline Hydrochloride			
Droperidol			
Drostanolone			
Drostanolone Propionate			
Dydrogesterone			
Dyflos			
Econazole		External, but in the case of vaginal use, only for the treatment of vaginal candidiasis	
Econazole Nitrate		External, but in the case of vaginal use, only for the treatment of vaginal candidiasis	
Ecothiopate Iodide			
Edrophonium Chloride			
Eflornithine Hydrochloride			
Eformoterol Fumarate			
Embutramide			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Emepronium Bromide			
Emetine	1.0%		
Emetine Bismuth			
Iodide			
Emetine Hydrochloride	Equivalent of 1.0% of Emetine		
Enalapril Maleate			
Encephalitis Virus, Tick-borne, Central European			
Enoxacin			
Enoxaparin Sodium			
Enoximone			
Ephedrine		(1) Internal (other than nasal sprays or nasal drops)	(1) 30 mg (MD) 60 mg (MDD)
	(2) 2.0%	(2) Nasal sprays or nasal drops	
		(3) External	
Ephedrine Hydrochloride		(1) Internal (other than nasal sprays or nasal drops)	(1) Equivalent of 30 mg of Ephedrine (MD) Equivalent of 60 mg of Ephedrine (MDD)
	(2) Equivalent of 2.0% of Ephedrine	(2) Nasal sprays or nasal drops	
		(3) External	
Ephedrine Sulphate		(1) Internal (other than nasal sprays or nasal drops)	(1) Equivalent of 30 mg of Ephedrine (MD) Equivalent of 60 mg of Ephedrine (MDD)
	(2) Equivalent of 2.0% of Ephedrine	(2) Nasal sprays or nasal drops	
		(3) External	
Epicillin			
Epirubicin			
Epirubicin Hydrochloride			
Epithiazide			
Epoetin Alfa			
Epoetin Beta			
Epoprostenol Sodium			
Ergometrine Maleate			
Ergometrine Tartrate			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Ergot, Prepared			
Ergotamine Tartrate			
Erythromycin			
Erythromycin Estolate			
Erythromycin Ethyl Carbonate			
Erythromycin Ethyl Succinate			
Erythromycin Lactobionate			
Erythromycin Phosphate			
Erythromycin Stearate			
Erythromycin Thiocyanate			
Esmolol Hydrochloride			
Estramustine Phosphate			
Estramustine Sodium Phosphate			
Etafedrine Hydrochloride			
Ethacrynic Acid			
Ethambutol Hydrochloride			
Ethamivan			
Ethamsylate			
Ethiazide			
Ethinyl Androstenediol			
Ethinyloestradiol			
Ethionamide			
Ethisterone			
Ethoglucid			
Ethoheptazine Citrate			
Ethopropazine Hydrochloride			
Ethosuximide			
Ethotoin			
Ethyl Biscoumacetate			
Ethyloestrenol			
Ethynodiol Diacetate			
Etodolac			
Etomidate			
Etomidate Hydrochloride			
Etoposide			
Etretinate			
Exemestane			
Famotidine			
Fazadinium Bromide			
Felbinac			
Felodipine			
Felypressin			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Fenbufen			
Fencamfamin Hydrochloride			
Fenclofenac			
Fenfluramine Hydrochloride			
Fenofibrate			
Fenoprofen			
Fenoprofen Calcium			
Fenoterol Hydrobromide			
Fenticonazole Nitrate		External use (but, in the case of vaginal use, only for the treatment of vulvovaginal candidiasis)	
Feprazone			
Ferrous Arsenate			
Ferumoxsil			
Fexofenadine Hydrochloride			
Filgrastim			
Finasteride			
Flavoxate Hydrochloride			
Flecainide Acetate			
Flosequinan			
Fluanisone			
Flubendazole			
Fluclorolone Acetonide			
Flucloxacillin Magnesium			
Flucloxacillin Sodium			
Fluconazole			
Flucylosine			
Fludrocortisone Acetate			
Flufenamic Acid			
Flumazenil			
Flumethasone			
Flumethasone Pivalate			
Flunisolide			
Fluocinolone Acetonide			
Fluocinonide			
Fluocortin Butyl			
Fluocortolone			
Fluocortolone Hexanoate			
Fluocortolone Pivalate			
Fluorescein Dilaurate			
Fluorometholone			
Fluorouracil			
Fluorouracil Trometamol			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Fluoxetine Hydrochloride			
Fluoxymesterone			
Flupenthixol Decanoate			
Flupenthixol Hydrochloride			
Fluperolone Acetate			
Fluphenazine Decanoate			
Fluphenazine Enantate			
Fluphenazine Hydrochloride			
Fluprednidene Acetate			
Fluprednisolone			
Fluprostenol Sodium			
Flurandrenolone			
Flurbiprofen	8.75 mg	throat lozenges	43.75 mg (MDD)
Flurbiprofen Sodium			
Fluspirilene			
Flutamide			
Fluticasone Propionate		Aqueous nasal sprays for the treatment of allergic rhinitis in persons not less than 18 years	
Flutrimazole			
Fluvastatin Sodium			
Fluvoxamine Maleate			
Folic Acid			500 mcg (MDD)
Formestane			
Formocortal			
Formoterol Fumarate			
Foscarnet Sodium			
Fosfestrol Sodium			
Fosfomycin Trometamol			
Fosinopril Sodium			
Framycetin Sulphate			
Frusemide			
Furazolidone			
Fusafungine			
Fusidic Acid			
Gabapentin			
Gadolinium			
Gadoteridol			
Gallamine Triethiodide			
Ganciclovir			
Ganciclovir Sodium			
Gelsemine	0.1%		

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Gelsemium			25 mg (MD) 75 mg (MDD)
Gemeprost			
Gemfibrozil			
Gentamicin			
Gentamicin Sulphate			
Gestodene			
Gestrinone			
Gestronol			
Gestronol Hexanoate			
Glibenclamide			
Glibornuride			
Gliclazide			
Glimepiride			
Glipizide			
Gliquidone			
Glisoxepide			
Glucagon			
Glycopyrronium Bromide			1 mg (MD) 2 mg (MDD)
Glymidine			
Gonadorelin			
Goserelin Acetate			
Gramicidin	0.2%	External	
Granisetron Hydrochloride			
Griseofulvin			
Growth Hormone			
Guanethidine Monosulphate			
Guanfacine Hydrochloride			
Guanoclor Sulphate			
Guanoxan Sulphate			
Halcinonide			
Halofantrine Hydrochloride			
Haloperidol			
Haloperidol Decanoate			
Heparin		External	
Heparin Calcium		External	
Hexachlorophane		External:	
	(a) 2.0% (b) 0.1% (c) 0.75%	(a) soaps (b) aerosols (c) preparations other than soaps and aerosols	
Hexamine Phenylcinchoninate			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Hexobarbitone			
Hexobarbitone Sodium			
Hexoestrol			
Hexoestrol Dipropionate			
L-Histidine Hydrochloride		Dietary or nutritive use	
Homatropine		(1) Internal	(1) 0.15 mg (MD) 0.45 mg (MDD)
		(2) External (except local ophthalmic use)	
Homatropine Hydrobromide			0.2 mg (MD) 0.6 mg (MDD)
Homatropine Methylbromide			2 mg (MD) 6 mg (MDD)
Hydralazine Hydrochloride			
Hydrargaphen		Local application to skin	
Hydrobromic Acid			
Hydrochlorothiazide			
Hydrocortisone			
Hydrocortisone Acetate			
Hydrocortisone Butyrate			
Hydrocortisone Caprylate			
Hydrocortisone Hydrogen Succinate			
Hydrocortisone Sodium Phosphate			
Hydrocortisone Sodium Succinnate			
Hydrocyanic Acid			
Hydroflumethiazide			
Hydroxychloroquine Sulphate		Prophylaxis of malaria	
Hydroxyprogesterone			
Hydroxyprogesterone Enanthate			
Hydroxyprogesterone Hexanoate			
Hydroxyurea			
Hydroxyzine Embonate			
Hydroxyzine Hydrochloride			
Hyoscine	(1) 0.15%	(1) Internal	

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
		(2) External (except local ophthalmic use)	
Hyoscine Butylbromide		(1) Internal: (a) by inhaler	(MDD) (x)
		(2) External	
Hyoscine Hydrobromide		(1) Internal: (a) by inhaler (b) otherwise than by inhaler	 (b) 300 mcg (MD) 900 mcg (MDD)(x)
		(2) External (except local ophthalmic use)	
Hyoscine Methobromide		(1) Internal: (a) by inhaler	
		(b) otherwise than by inhaler	(b) 2.5 mg (MD) 7.5 mg (MDD)(x)
		(2) External	
Hyoscine Methonitrate		(1) Internal: (a) by inhaler	
		(b) otherwise than by inhaler	(b) 2.5 mg (MD) 7.5 mg (MDD)(x)
		(2) External	
Hyoscyamine		(1) Internal: (a) by inhaler	
		(b) otherwise than by inhaler	(b) 300 mcg (MD) 1 mg (MDD)(x)
		(2) External	
		(3) Preparations for the relief of asthma in the form of cigarettes, smoking mixtures or fumigants which contain Hyoscyamine as an alkaloid of Stramonium	
Hyoscyamine Hydrobromide		(1) Internal: (a) by inhaler	

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
		(b) otherwise than by inhaler	(b) Equivalent of 300 mcg of Hyoscyamine (MD) Equivalent of 1 mg of Hyoscyamine (MDD)(x)
		(2) External	
Hyoscyamine Sulphate		(1) Internal: (a) by inhaler	
		(b) otherwise than by inhaler	(b) Equivalent of 300 mcg of Hyoscyamine (MD) Equivalent of 1 mg of Hyoscyamine (MDD)(x)
		(2) External	
Ibuprofen			
Ibuprofen Lysine		Rheumatic and muscular pain, pain of non-serious arthritic conditions, backache, neuralgia, migraine, headache, dental pain, dysmenorrhoea, feverishness, symptoms of colds and influenza	
		Internal	(a) In the case of a prolonged release preparation 600 mg (MD) 1,200 mg (MDD)
			(b) In any other case 400 mg (MD) 1,200 mg (MDD)

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Idarubicin Hydrochloride			
Idoxuridine			
Ifosfamide			
Ignatius Bean			
Imidapril Hydrochloride			
Imipenem Hydrochloride			
Imipramine			
Imipramine Hydrochloride			
Imipramine Ion Exchange Resin Bound Salt or Complex			
Indapamide Hemihydrate			
Indinavir			
Indomethacin			
Indomethacin Sodium			
Indoramin Hydrochloride			
Indoprofen			
Inosine Pranobex			
Insulin			
Iodamide			
Iodamide Meglumine			
Iodamide Sodium			
Iohexol			
Iomeprol			
Iopamidol			
Iopentol			
Iothalamic Acid			
Ioversol			
Ioxaglic Acid			
Ipratropium Bromide			
Iprindole Hydrochloride			
Iproniazid Phosphate			
Irbesartan			
Isoaminile			
Isoaminile Citrate			
Isocarboxazid			
Isoconazole Nitrate		External, but in the case of vaginal use, only for the treatment of vaginal candidiasis	
Isoetharine			
Isoetharine Hydrochloride			
Isoetharine Mesylate			
Isoniazid			
Isoprenaline Hydrochloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Isoprenaline Sulphate			
Isopropamide Iodide			Equivalent of 2.5 mg of Isoprop-amide ion (MD) Equivalent of 5.0 mg of Isoprop-amide ion (MDD)
Isotretinoin			
Isradipine			
Itraconazole			
Jaborandi		External	
Kanamycin Acid Sulphate			
Kanamycin Sulphate			
Ketamine Hydrochloride			
Ketoconazole			
Ketoprofen			
Ketorolac Trometamol			
Ketotifen Pumarate			
Labetalol Hydrochloride			
Lachesine Chloride			
Lacidipine			
Lamivudine			
Lamotrigine			
Lanatoside C			
Lanatoside Complex A, B and C			
Lanzoprazole			
Latamoxef Disodium			
Latanaprost			
Lercanidipine Hydrochloride			
Letrozole			
Levallorphan Tartrate			
Levobunolol Hydrochloride			
Levocabastine Hydrochloride			
Levocarnitine		For dietary supplementation	
Levodopa			
Levofloxacin Hemihydrate			
Levonorgestrel			
Lidoflazine			
Lignocaine		Any use (except local ophthalmic use)	

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Lignocaine Hydrochloride		Any use (except local ophthalmic use)	
Lincomycin			
Lincomycin Hydrochloride			
Liothyronine Sodium			
Lisinopril			
Lithium Carbonate			Equivalent of 5 mg of Lithium (MD) Equivalent of 15 mg of Lithium (MDD)
Lithium Citrate			
Lithium Sulphate			Equivalent of 5 mg of Lithium (MD) Equivalent of 5 mg of Lithium (MDD)
Lithium Succinate			
Lobeline		(1) Internal	(1) 3 mg (MD) 9 mg (MDD)
		(2) External	
Lobeline Hydrochloride		(1) Internal	(1) Equivalent of 3 mg of Lobeline (MD)
			Equivalent of 9 mg of Lobeline (MDD)
		(2) External	
Lobeline Sulphate		(1) Internal	(1) Equivalent of 3 mg of Lobeline (MD) Equivalent of 9 mg of Lobeline (MDD)
		(2) External	

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Lodaximide Trometamol	equivalent of 0.1% Lodoxamide	For the treatment of ocular signs and symptoms of allergic conjunctivitis, in adults and in children aged 4 years and over	
Lofepramine			
Lofepramine Hydrochloride			
Lofexidine Hydrochloride			
Lomefloxacin Hydrochloride			
Lornoxicam			
Lomustine			
Loperamide Hydrochloride		Treatment of acute diarrhoea	
Loratidine			
Losartan Potassium			
Loxapine Succinate			
Lung Surfactant Porcine			
Luteinising Hormone			
Lymecycline			
Lynoestrenol			
Lypressin			
Lysuride Maleate			
Mafenide			
Mafenide Acetate			
Mafenide Hydrochloride			
Mafenide Propionate	5.0%	Eye drops	
Magnesium Fluoride			
Magnesium Metrizoate			
Mandragora Autumnalis			
Mannomustine Hydrochloride			
Maprotiline Hydrochloride			
Mebanazine			
Mebendazole			
Mebeverine Hydrochloride		(a) For the symptomatic relief of irritable bowel syndrome (b) For uses other than the symptomatic relief of irritable bowel syndrome	(a) 135mg (MD) 405mg (MDD) (b) 100 mg (MD) 300mg (MDD)

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Mebeverine Pamoate			
Mebhydrolin			
Mebhydrolin Napadisylate			
Mecamylamine Hydrochloride			
Mecillinam			
Meclofenoxate Hydrochloride			
Medigoxin			
Medrogestone			
Medroxyprogesterone Acetate			
Mefenamic Acid			
Mefloquine Hydrochloride			
Mefruside			
Megestrol			
Megestrol Acetate			
Meglumine Gadopentetate			
Meglumine lodoxamate			
Meglumine loglycamate			
Meglumine lothalamate			
Meglumine lotroxate			
Meglumine loxaglate			
Meloxicam			
Melphalan			
Melphalan Hydrochloride			
Menotrophin			
Mepenzolate Bromide			25 mg (MD) 75 mg (MDD)
Mephenesin			
Mephenesin Carbamate			
Mepivacaine Hydrochloride		Any use (except local ophthalmic use)	
Meptazinol Hydrochloride			
Mequitazine			
Mercaptamine Bitartrate			
Mercaptopurine			
Mersalyl			
Mersalyl Acid			
Mesalazine			
Mesna			
Mesterolone			
Mestranol			
Metaraminol Tartrate			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Metergoline			
Metformin Hydrochloride			
Methacycline			
Methacycline Calcium			
Methacycline Hydrochloride			
Methallenoestril			
Methandienone			
Methicillin Sodium			
Methixene			
Methixene Hydrochloride			
Methocarbamol			
Methocidin		Throat lozenges and throat pastilles	
Methohexitone Sodium			
Methoin			
Methoserpidine			
Methotrexate			
Methotrexate Sodium			
Methotrimeprazine			
Methotrimeprazine Hydrochloride			
Methotrimeprazine Maleate			
Methoxamine Hydrochloride	0.25%	Nasal sprays, or nasal drops, not containing in either case liquid paraffin as a vehicle	
Methsuximide			
Methyclothiazide			
Methyldopa			
Methyldopa Hydrochloride			
Methylephedrine Hydrochloride			30 mg (MD) 60 mg (MDD)
Methylprednisolone			
Methylprednisolone Acetate			
Methylprednisolone Sodium Succinate			
Methyltestosterone			
Methylthiouracil			
Methysergide Maleate			
Metipranolol			
Metirosine			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Metoclopramide Hydrochloride			
Metolazone			
Metoloprolol Tartrate			
Metoprolol Fumarate			
Metoprolol Succinate			
Metronidazole			
Metronidazole Benzoate			
Metyrapone			
Mexiletine Hydrochloride			
Mezlocillin Sodium			
Mianserin Hydrochloride			
Mibefradil Dihydrochloride			
Miconazole		External	
Miconazole Nitrate		External, but in the case of vaginal use, only for the treatment of vaginal candidiasis	
Mifepristone			
Miglitol			
Milrinone			
Milrinone Lactate			
Minocycline			
Minocycline Hydrochloride			
Minoxidil	(1) 2.0%	External	
	(2) 5.0%	External, for the treatment of alopecia androgenetica in men who have attained the age of 18 years but have not attained the age of 65 years	
Mirtazapine			
Misoprostol			
Mitobronitol			
Mitomycin C			
Mitozantrone Hydrochloride			
Mivacurium Chloride			
Mizolastine			
Moclobemide			
Modafinil			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Moexipril Hydrochloride			
Molgramostim			
Molindone Hydrochloride			
Mometasone Furoate			
Moracizine Hydrochloride			
Morazone Hydrochloride			
Moxisylyte Hydrochloride			
Moxonidine			
Mupirocin			
Mupirocin Calcium			
Mustine Hydrochloride			
Mycophenolate Mofetil			
Nabilone			
Nabumetone			
Nadolol			
Nafarelin Acetate			
Naftidrofuryl Oxalate			
Naftifine Hydrochloride			
Nalbuphine Hydrochloride			
Nalidixic Acid			
Nalorphine Hydrobromide			
Naloxone Hydrochloride			
Naltrexone Hydrochloride			
Nandrolone Decanoate			
Nandrolone Laurate			
Nandrolone Phenylpropionate			
Naphazoline Hydrochloride	(1) 0.05%	(1) Nasal sprays, or nasal drops, not containing in either case liquid paraffin as a vehicle	
	(2) 0.015%	(2) Eye drops	
Naphazoline Nitrate	0.05%	Nasal sprays, or nasal drops, not containing in either case liquid paraffin as a vehicle	
Naproxen			
Naproxen Sodium			
Naratriptan Hydrochloride			
Natamycin			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Nebivolol Hydrochloride			
Nedocromil Sodium			
Nefazadone Hydrochloride			
Nefopam Hydrochloride			
Neomycin			
Neomycin Oleate			
Neomycin Palmitate			
Neomycin Sulphate			
Neomycin Undecanoate			
Neostigmine Bromide			
Neostigmine Methylsulphate			
Netilmicin Sulphate			
Nicardipine Hydrochloride			
Nicergoline			
Niceritrol			
Nicotinic Acid		Any use (except for the treatment of hyperlipid-aemia)	600 mg (MDD)
Nicoumalone			
Nifedipine			
Nifenazone			
Nikethamide			
Nilutamide			
Nimodipine			
Niridazole			
Nisoldipine			
Nitrendipine			
Nitrofurantoin			
Nitrofurazone			
Nizatidine			
Nomifensine Maleate			
Noradrenaline			
Noradrenaline Acid Tartrate			
Norethandrolone			
Norethisterone			
Norethisterone Acetate			
Norethisterone Heptanoate			
Norethynodrel			
Norfloxacin			
Norgestimate			
Norgestrel			
Nortriptyline Hydrochloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Noscapine			
Noscapine Hydrochloride			
Novobiocin Calcium			
Novobiocin Sodium			
Nux Vomica Seed			
Nystatin			
Octacosactrin			
Octreotide			
Oestradiol			
Oestradiol Benzoate			
Oestradiol Cypionate			
Oestradiol Dipropionate			
Oestradiol Diundecanoate			
Oestradiol Enanthate			
Oestradiol Phenylpropionate			
Oestradiol Undecanoate			
Oestradiol Valerate			
Oestriol			
Oestriol Di-Hemi Succinate			
Oestrogenic Substances, Conjugated			
Oestrone			
Ofloxacin			
Olanzapine			
Olsalazine Sodium			
Omeprazole			
Omeprazole Magnesium			
Ondansetron			
Ondansetron Hydrochloride			
Orciprenaline Sulphate			
Orphenadrine Citrate			
Orphenadrine Hydrochloride			
Ouabain			
Ovarian Gland, Dried			
Oxamniquine			
Oxandronolone			
Oxantel Pamoate			
Oxaprozin			
Oxatomide			
Oxedrine Tartrate			
Oxethazaine			
Oxidronate Sodium			
Oxitropium Bromide			
Oxolinic Acid			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Oxpentifylline			
Oxprenolol Hydrochloride		Any use (except local ophthalmic use)	
Oxybuprocaine Hydrochloride			
Oxybutynin Hydrochloride			
Oxymetholone			
Oxypertine			
Oxypertine Hydrochloride			
Oxyphenbutazone			
Oxyphencyclimine Hydrochloride			
Oxyphenonium Bromide			5 mg (MD) 15 mg (MDD)
Oxytetracycline			
Oxytetracycline Calcium			
Oxytetracycline Dihydrate			
Oxytetracycline Hydrochloride			
Oxytocin, Natural			
Oxytocin, Synthetic			
Pamidronate Disodium			
Pancreatin	(1) 21,000 European Pharmacopoeia units of lipase per capsule		(1) Capsules
	(2) 25,000 European Pharmacopoeia units of lipase per g		(2) Powder
Pancuronium Bromide			
Pantoprazole			
Pantoprazole Sodium			
Papaverine		(1) By inhaler	
		(2) Otherwise than by inhaler	(2) 50 mg (MD) 150 mg (MDD)
Papaverine Hydrochloride		(1) By inhaler	

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
		(2) Otherwise than by inhaler	(2) Equivalent of 50 mg of Papaverine (MD) Equivalent of 150 mg of Papaverine (MDD)
Paracetamol		Any form (except non-effervescent tablets and capsules)	
Paraldehyde			
Paramethadione			
Paramethasone Acetate			
Parathyroid Gland			
Pargyline Hydrochloride			
Paroxetine Hydrochloride			
Pecilocin			
Penamecillin			
Penbutolol Sulphate			
Penciclovir			
Penicillinamine			
Penicillinamine Hydrochloride			
Pentamidine Isethionate			
Pentamidronate Disodium			
Penthienate Methobromide			5 mg (MD) 15 mg (MDD)
Pentolinium Tartrate			
Perfluamine			
Pergolide Mesylate			
Perhexiline Maleate			
Perindopril			
Pericyazine			
Perindopril Erbumine			
Perphenazine			
Phenacetin	0.1%		
Phenazone		External	
Phenazone and Caffeine Citrate			
Phenazone Salicylate			
Phenbutrazate Hydrochloride			
Phenelzine Sulphate			
Phenethicillin Potassium			
Phenformin Hydrochloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Phenglutarimide Hydrochloride			
Phenindione			
Phenolphthalein			
Phenoxybenzamine Hydrochloride			
Phenoxymethylpenicillin			
Phenoxymethylpenicillin Calcium			
Phenoxymethylpenicillin Potassium			
Phenprocoumon			
Phensuximide			
Phentolamine Hydrochloride			
Phentolamine Mesylate			
Phenylbutazone			
Phenylbutazone Sodium			
Phenylpropanolamine Hydrochloride		Internal: (a) all preparations (except controlled release capsules, nasal sprays or nasal drops)	(a) 25 mg (MD) 100 mg (MDD)
		(b) controlled release capsules	(b) 50 mg (MD) 100 mg (MDD)
	(c) 2.0%	(c) nasal sprays or nasal drops	
Phenytoin			
Phenytoin Sodium			
Phthalylsulphathiazole			
Physostigmine			
Physostigmine Aminoxide Salicylate			
Physostigmine Salicylate			
Physostigmine Sulphate			
Phytomenadine		Any use except the prevention or treatment of haemorrhagic disorders	
Picrotoxin			
Pilocarpine			
Pilocarpine Hydrochloride			
Pilocarpine Nitrate			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Pimozide			
Pindolol			
Pipenzolate Bromide			5 mg (MD) 15 mg (MDD)
Piperacillin Sodium			
Piperazine Oestrone Sulphate			
Piperidolate Hydrochloride			50 mg (MD) 150 mg (MDD)
Pipothiazine Palmitate			
Piracetam			
Pirbuterol Acetate			
Pirbuterol Hydrochloride			
Pirenzepine Dihydrochloride Monohydrate			
Pirenzepine Hydrochloride			
Piretamide			
Piroxicam			
Piroxicam Beta-Cyclodextrin			
Pituitary Gland (Whole Dried)		By inhaler	
Pituitary, Powdered (Posterior Lobe)		By inhaler	
Pivampicillin Hydrochloride			
Pivmecillinam			
Pivmecillinam Hydrochloride			
Pizotifen			
Pizotifen Malate			
Plicamycin			
Podophyllotoxin			
Podophyllum			
Podophyllum Indian			
Podophyllum Resin	20.0%	External: ointment or impregnated plaster	
Poldine Methylsulphate			2 mg (MDD) 6 mg (MDD)
Polidexide			
Polymyxin B Sulphate			
Polyestradiol Phosphate			
Polythiazide			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Poppy Capsule			
Potassium Arsenite	0.0127%		
Potassium Bromide			
Potassium Canrenoate			
Potassium Clavulanate			
Potassium Perchlorate			
Practolol			
Pralidoxime Chloride			
Pralidoxime lodide			
Pralidoxime Mesylate			
Pramipexole Dihydrochloride			
Pravastatin Sodium			
Prazosin Hydrochloride			
Prednisolone			
Prednisolone Acetate			
Prednisolone Butylacetate			
Prednisolone Hexanoate			
Prednisolone Pivalate			
Prednisolone Sodium Phosphate			
Prednisolone Sodium m-Sulphobenzoate			
Prednisolone 21-Steaglate			
Prednisolone m-Sulphobenzoate			
Prednisone			
Prednisone Acetate			
Prenalterol Hydrochloride			
Prenylamine Lactate			
Prilocaine Hydrochloride		Any use (except local ophthalmic use)	
Primidone			
Probenecid			
Probucol			
Procainamide Hydrochloride			
Procaine Hydrochloride		Any use (except local ophthalmic use)	
Procaine Penicillin			
Procarbazine Hydrochloride			
Prochlorperazine			
Prochlorperazine Edisylate			
Prochlorperazine Maleate			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Prochlorperazine Mesylate			
Procyclidine Hydrochloride			
Progesterone			
Prolactin			
Proligestone			
Prolintane Hydrochloride			
Promazine Embonate			
Promazine Hydrochloride			
Propafenone			
Propafenone Hydrochloride			
Propanidid			
Propantheline Bromide			15 mg (MD) 45 mg (MDD)
Propiverine Hydrochloride			
Propofol			
Propranolol Hydrochloride			
Propylthiouracil			
Proquazone			
Protamine Sulphate			
Prothionamide			
Protirelin			
Protriptyline Hydrochloride			
Proxymetacaine Hydrochloride		Any use (except local ophthalmic use)	
Pseudoephedrine Hydrochloride		Internal	In the case of a controlled release preparation: 120 mg (MD) 180 mg (MDD) In any other case: 60 mg (MD) 180 mg (MDD)
Pseudoephedrine Sulphate			60 mg (MD) 180 mg (MDD)
Pyrantel Embonate			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Pyrantel Tartrate			
Pyrazinamide			
Pyridostigmine Bromide			
Pyrimethamine			
Quetiapine Fumarate			
Quinagolide Hydrochloride			
Quinapril			
Quinapril Hydrochloride			
Quinestradol			
Quinestrol			
Quinethazone			
Quinidine			
Quinidine Bisulphate			
Quinidine Polygalacturonate			
Quinidine Sulphate			
Quinine			100 mg (MD) 300 mg (MDD)
Quinine Bisulphate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Dihydrochloride			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Ethyl Carbonate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Glycero-phosphate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Quinine Hydrobromide			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Hydrocholride			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Iodobismuthate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Phenylcinchoninate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Phosphate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Salicylate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine Sulphate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Quinine Tannate			Equivalent of 100 mg of Quinine (MD) Equivalent of 300 mg of Quinine (MDD)
Quinine and Urea Hydrochloride			
Ramipril			
Ranitidine Bismuth Citrate			
Ranitidine Hydrochloride			
Rauwolfia Serpentina			
Rauwolfia Vomitoria			
Reboxetine			
Reboxetine Mesilate			
Remoxipride Hydrochloride			
Reproterol Hydrochloride			
Rescinnamine			
Reserpine			
Rifabutin			
Rifampicin			
Rifampicin Sodium			
Rifamycin			
Rimexolone			
Rimiterol Hydrobromide			
Risperidone			
Ritodrine Hydrochloride			
Ritonavir			
Rolitetracycline Nitrate			
Ropinorole Hydrochloride			
Sabadilla			
Salbutamol			
Salbutamol Sulphate			
Salcatonin			
Salcatonin Hydrated Polyacetate			
Salmefamol			
Salmeterol Hydroxynaphthoate			
Salsalate			
Saquinavir			
Saralasin Acetate			
Selegiline Hydrochloride			
Sera and Antisera –			
Botulin Antitoxin			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Diphtheria Antitoxin			
Gas-gangrene Antitoxin (Oedematiens)			
Gas-gangrene Antitoxin (Perfringens)			
Gas-gangrene Antitoxin (Septicum)			
Mixed Gas-gangrene Antitoxin			
Leptospira Antiserum			
Rabies Antiserum			
Scorpion Venom Antiserum			
Snake Venom Antiserum			
Tetanus Antitoxin			
Serum Gonadotrophin			
Sermorelin			
Sertindole			
Sertraline Hydrochloride			
Sevoflurane			
Sibutramine Hydrochloride			
Silver Sulphadiazine			
Simvastatin			
Sissomicin			
Sissomicin Sulphate			
Snake Venoms			
Sodium Acetrizoate			
Sodium Aminosalicylate			
Sodium Antimonylgluconate			
Sodium Arsanilate			
Sodium Arsenate			
Sodium Arsenite	0.013%		
Sodium Bromide			
Sodium Clodronate			
Sodium Cromoglycate		Administration through the nose	
Sodium Ethacrynate			
Sodium Fluoride	(1) 0.33% (y)	(1) Dentifrices	
		(2) Other preparations for use in the prevention of dental of dental caries in the form of – (a)tablets or drops;	(a) 2.2 mg (MDD)

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
	(b) 0.2%	(b) mouth-washes (other than those for daily use);	
	(c) 0.05%	(c) mouth-washes for daily use	
Sodium Fusidate			
Sodium Metrizoate			
Sodium Monofluorophosphate	1.14% (y)	Dentifrice	
Sodium Stibogluconate			
Sodium Valproate			
Somatorelin Acetate			
Somatrem			
Somatropin			
Sotalol Hydrochloride			
Sparfloxacin			
Spectinomycin			
Spectinomycin Hydrocloride			
Spiramycin			
Spiramycin Adipate			
Spironolactone			
Stannous Fluoride	(1) 0.62% (y)	(1) Dentifrice	
	(2) 0.4%	(2) Dental gels for use in the prevention and treatment of dental caries and decalcification of the teeth	
Stanolone			
Stanozolol			
Stavudine			
Stilboestrol			
Stilboestrol Dipropionate			
Streptodornase		External	
Streptokinase		External	
Streptomycin			
Streptomycin Sulphate			
Strychnine			
Strychnine Arsenate			
Strychnine Hydrochloride			
Strychnine Nitrate			
Styramate			
Succinylsulphathiazole			
Sucralfate			
Sulbactam			
Sodium			
Sulbenicillin			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Sulbenicillin Sodium			
Sulbenicillin Tosylate			
Sulconazole Nitrate		External (except vaginal use)	
Sulfacytine			
Sulfadoxine			
Sulfamonomethoxine			
Sulindac			
Sulphabenzamide			
Sulphacetamide			
Sulphacetamide Sodium			
Sulphadiazine			
Sulphadiazine Sodium			
Sulphadimethoxine			
Sulphadimidine			
Sulphadimidine Sodium			
Sulphafurazole			
Sulphafurazole Diethanolamine			
Sulphaguanidine			
Sulphaloxic Acid			
Sulphamerazine			
Sulphamerazine Sodium			
Sulphamethizole			
Sulphamethoxazole			
Sulphamethoxydazine			
Sulphamethoxypyridazine			
Sulphamethoxypyridazine Sodium			
Sulphametopyrazine			
Sulphamoxole			
Sulphanilamide			
Sulphaphenazole			
Sulphapyridine			
Sulphapyridine Sodium			
Sulphasalazine			
Sulphathiazole			
Sulphathiazole Sodium			
Sulphaurea			
Sulphinpyrazone			
Sulpiride			
Sultamicillin			
Sultamicillin Tosylate			
Sulthiame			
Sumatriptan			
Sumatriptan Succinate			
Suprofen			
Suxamethonium Bromide			
Suxamethonium Chloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Suxethonium Bromide			
Tacalcitol Monohydrate			
Tacrine Hydrochloride			
Talampicillin			
Talampicillin Hydrochloride			
Talampicillin Napsylate			
Tamoxifen			
Tamoxifen Citrate			
Tamsulosin Hydrochloride			
Tazarotene			
Tazobactam Sodium			
Teicoplanin			
Temocapril Hydrochloride			
Temocillin Sodium			
Tenoxicam			
Terazosan Hydrochloride			
Terbinafine			
Terbinafine Hydrochloride			
Terbutaline			
Terbutaline Sulphate			
Terfenadine			
Terodiline Hydrochloride			
Tertipressin			
Testosterone			
Testosterone Acetate			
Testosterone 17B Chloral Hemiacetal			
Testosterone Cyclohexyl-propionate			
Testosterone Cypionate			
Testosterone Decanoate			
Testosterone Enanthate			
Testosterone Isocaproate			
Testosterone Phenylpropionate			
Testosterone Propionate			
Testosterone Undecanoate			
Tetrabenazine			
Tetracosactrin			
Tetracosactrin Acetate			
Tetracycline			
Tetracycline Hydrochloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Tetracycline Phosphate Complex			
Tetroxoprim			
Thallium Acetate			
Thallous Chloride			
Thiabendazole			
Thiambutosine			
Thiethylperazine			
Thiethylperazine Maleate			
Thiocarlide			
Thioguanine			
Thiopentone Sodium			
Thiopropazate Hydrochloride			
Thioproperazine Mesylate			
Thioridazine			
Thioridazine Hydrochloride			
Thiosinamine			
Thiotepa			
Thiothixene			
Thiouracil			
Thymoxamine Hydrochloride			
Thyroid			
Thyrotrophin			
Thyroxine Sodium			
Tiamulin Fumarate			
Tiaprofenic Acid			
Tibolone			
Ticarcillin Sodium			
Ticlopidine Hydrochloride			
Tigloidine Hydrobromide			
Tiludronate Disodium			
Timolol Maleate			
Tinidazole			
Tinzaparin			
Tioconazole	2.0%	(1) External, but in the case of vaginal use only, external use for treatment of vaginal candidiasis	
Tizanidine Hydrochloride			
Tobramycin			
Tobramycin Sulphate			
Tocainide Hydrochloride			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Tofenacin Hydrochloride			
Tolazamide			
Tolazoline Hydrochloride		External	
Tolbutamide			
Tolbutamide Sodium			
Tolcapone			
Tolfenamic Acid			
Tolmetin Sodium			
Topiramate			
Torasemide			
Toremifene			
Tramadol Hydrochloride			
Trandolapril			
Tranexamic Acid			
Tranylepromine Sulphate			
Trazodone Hydrochloride			
Treosulfan			
Tretinoin			
Triamcinolone			
Triamcinolone Acetonide			
Triamcinolone Diacetate			
Triamcinolone Hexacetonide			
Triamterene			
Tribavirin			
Triclofos Sodium			
Trientine Dihydrochloride			
Trifluoperazine			
Trifluoperazine Hydrochloride			
Trifluperidol			
Trifluperidol Hydrochloride			
Trilostane			
Trimeprazine			
Trimeprazine Tartrate			
Trimetaphan Camsylate			
Trimetazidine			
Trimetazidine Hydrochloride			
Trimethoprim			
Trimipramine Maleate			
Trimipramine Mesylate			
Tropicamide			
Tropisetron Hydrochloride			
Troxidone			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
L-Tryptophan		(1) Dietary or nutritive use (2) Any external use	
Tubocurarine Chloride			
Tulobuterol			
Tulobuterol Hydrochloride			
Tyrothricin		Throat lozenges or throat pastilles	
Uramustine			
Urea Stibamine			
Urethane			
Uridine-5-Triphosphoric Acid			
Urofollitrophin			
Urokinase			
Ursodeoxychloric Acid			
Vaccines –			
Athrax Vaccine (Bacillus) Anthracis)			
Bacillus Calmette-Guerin Vaccine			
Bacillus Salmonella Typhi Vaccine			
Percutaneous Bacillus Calmette- Guerin Vaccine			
Cholera Vaccine			
Diphtheria Vaccine			
Adsorbed Diphtheria Vaccine			
Diphtheria and Tetanus Vaccine			
Adsorbed Diphtheria and Tetanus Vaccine			
Diphtheria, Tetanus and Pertussis Vaccine			
Adsorbed Diphtheria, Tetanus and Pertussis Vaccine			
Diphtheria, Tetanus and Poliomyelitis Vaccine			
Diphtheria, Tetanus, Pertussis and Poliomyelitis Vaccine			
Eltor Vaccine			
Influenza Vaccine			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Hepatitis B Vaccine			
Measles Vaccine (Live Attenuated)			
Meningococcal Polysaccharide Vaccine			
Mumps Vaccine			
Pertussis Vaccine			
Plague Vaccine			
Pneumococcal Vaccine (Bacterial Antigen)			
Poliomyelitis Vaccine (Inactivated)			
Poliomyelitis Vaccine (Live Oral)			
Rabies Vaccine			
Rubella Vaccine (Live Attenuated)			
Rubella, Mumps, Measles Vaccine			
Tetanus Vaccine			
Adsorbed Tetanus Vaccine			
Tetanus and Pertussis Vaccine			
Tuberculin Purified Protein Derivative			
Old Tuberculin			
Typhoid Vaccine			
Typhoid – Paratyphoid A and B Vaccine			
Typhoid – Paratyphoid A and B and Cholera Vaccine			
Typhoid – Paratyphoid A and B and Tetanus Vaccine			
Typhus Vaccine			
Yellow Fever Vaccine			
Valaciclovir			
Valaciclovir Hydrochloride			
Valproic Acid			
Valsartan			
Vancomycin Hydrochloride			
Vasopressin Injection			
Vasopressin Tannate			
Vecuronium Bromide			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Venlafaxine			
Venlafaxine Hydrochloride			
Verapamil Hydrochloride			
Veratrine			
Veratrum (Green and White)			
Vidarabine			
Vigabatrin			
Viloxazine Hydrochloride			
Vinblastine Sulphate			
Vincristine Sulphate			
Vindesine Sulphate			
Viomycin Pantothenate			
Viomycin Sulphate			
Vitamin A		(1) Internal	(1) 7500 iu (2250 mcg Retinol equivalent) (MDD)
		(2) External	
Vitamin A Acetate		(1) Internal	(1) Equivalent to 7500 iu Vitamin A (2250 mcg Retinol equivalent) (MDD)
		(2) External	
Vitamin A Palmitate		(1) Internal	(1) Equivalent to 7500 iu Vitamin A (2250 mcg Retinol equivalent) (MDD)
		(2) External	
Warfarin			
Warfarin Sodium			
Xamoterol Fumarate			
Xipamide			
Yohimbine Hydrochloride			
Zalcitabine			
Zidovudine			
Zimeldine Hydrochloride			
Zolmitriptan			
Zolpidem			
Zomepirac Sodium			

Prescription Only Medicine	**Circumstances In Which Substances Are Not Prescription Only Medicines**		
Zopiclone			
Zuclopenthixol Acetate			
Zuclopenthixol Decanoate			
Zuclopenthixol Hydrochloride			
Note –	1. In relation to a medicinal product that contains more than one of the substances Atropine, Atropine Methobromide, Atropine Methonitrate, Atropine Oxide Hydrochloride, Atropine Sulphate, Hyoscine, Hyoscine Butylbromide, Hyoscine Hydrobromide, Hyoscine Methobromide, Hyoscine Methonitrate, Hyoscyamine, Hyoscyamine Hydrobromide and Hyoscyamine Sulphate, the maximum daily dose for the purposes of column 4 is 1 mg of the total alkaloids contained in the product that are derived from Belladonna, Hyoscyamus, Stramonium or other solanaceous plant, and there is no maximum dose.		
	2. In relation to a medicinal product that contains more than one of the substances Sodium Fluoride, Sodium Monofluorophosphate and Stannous Fluoride combined in a dentifrice, the maximum strength of the combination for the purposes of column 2 shall not exceed 0.15% calculated as Fluorine.		

PART 2

(Articles 1(2), 3(2) and 7)

Controlled Drugs	**Circumstances In Which Controlled Drugs Are Not Prescription Only Medicines**		
Column 1	*Column 2*	*Column 3*	*Column 4*
Substance	**Maximum strength**	**Pharmaceutical form**	**Maximum dose**
Codeine; its salts	Equivalent of 1.5% of Codeine Monohydrate		Equivalent of 20 mg of Codeine Monohydrate
Dihydrocodeine; its salts	Equivalent of 1.5%of Dihydrocodeine		Equivalent of 10 mg of Dihydroco-deine
Ethylmorphine; its salts	Equivalent of 0.2% of Ethylmorphine		Equivalent of 7.5 mg of Ethyl-morphine
Morphine; its salts	(1) Equivalent of 0.02% of anhydrous Morphine	(1) Liquid	(1) Equivalent of 3 mg of anhydrous Morphine
	(2) Equivalent of 0.04% of anhydrous Morphine; equivalent of 300 mcg of anhydrous Morphine	(2) Solid	(2) Equivalent of 3 mg of anhydrous Morphine
Medicinal Opium	(1) Equivalent of 0.02%of anhydrous Morphine	(1) Liquid	(1) Equivalent of 3 mg of anhydrous Morphine
	(2) Equivalent of 0.04% of anhydrous Morphine	(2) Solid	(2) Equivalent of 3 mg of anhydrous Morphine
Pholcodine; its salts	Equivalent of 1.5% of Pholcodine		Equivalent of 20 mg of Pholcodine

PART 3

(Article 3(4))

NAMED PRODUCTS THAT ARE NOT PRESCRIPTION ONLY MEDICINES

TABLE A	
Name and product licence number of medicinal products that are not prescription only medicines	
Adcortyl in Orabase for Mouth Ulcers	0034/0321
Anusol Plus HC Ointment	0018/0223
Anusol Plus HC Suppositories	0018/0224
Beechams Hydrocortisone Cream	0079/0203
Boots Hydrocortisone Ointment	0014/0364
Calacort Cream	12650/0001
Canesten Hydrocortisone Cream	0010/0216
Corlan Pellets	0039/0397
Cortaid Cream 1%	0032/0126
Corteze Cream	0001/0107
Cortiderm	2855/0010
Cortril Topical Ointment 1% (non-greasy)	0057/0251
Dermacort Hydrocortisone Cream	8265/0002
Dioderm Hydrocortisone Cream	0173/0153
Efcortelan Eczema Cream	10949/0234
Efcortelan Eczema Ointment	10949/0235
Eurax HC Cream	0001/5010R
Hc45 Hydrocortisone Cream	0327/0039
Herpetad Cold Sore Cream	4986/0007
Jungle Formula Bite & Sting Relief Cream	2855/0010
Lanacort Cream	3157/0008
Lanacort Ointment	3157/0011
Perinal Spray	0173/0049
Pharmacort Cream 0.5%	0011/0077
Proctocream HC	0036/0065
Soothelip Cold Sore Cream	0142/0426
Timocort Hydrocortisone Cream	0044/0090
Timocort Hydrocortisone Cream 1%	0063/0076
Wasp-Eze Hydrocortisone Cream	8452/0012
Zaclovir Cold Sore Cream	4986/0007
Zenoxone Cream	0181/0033
Zovirax Cold Sore Cream	0003/0304

TABLE B	
Relevant product licence holder and name and product licence number of medicinal products that are not prescription only medicines	
Leo Laboratories Limited:	
Hydrocortisone Acetate Cream BP 0.5%	0043/0150
Hydrocortisone Acetate Cream BP 1.0%	0043/0151
Richard Daniel and Son Limited:	
Hydrocortisone Cream BP 1.0%	0842/0011

PART 4[12]

(Article 3(4))

OTHER MEDICINAL PRODUCTS THAT ARE NOT PRESCRIPTION ONLY MEDICINES

1 A medicinal product shall not be a prescription only medicine by reason that it contains the substance aciclovir, where –

(a) the maximum strength of the aciclovir in the medicinal product does not exceed 5%;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 2 g of the medicinal product; and

(c) the medicinal product is indicated only for external application for the treatment of herpes simplex virus infections of the lips and face (Herpes labialis).

2 A medicinal product shall not be a prescription only medicine by reason that it contains the substance acrivastine, where –

(a) the medicinal product is sold or supplied in a container, or package, containing not more than 240 mg of acrivastine; and

(b) the container or package is labelled to show a maximum daily dose of 24 mg of acrivastine.

3 A medicinal product shall not be a prescription only medicine by reason that it contains the substance aloxiprin, where –

(a) the medicinal product is in the form of non-effervescent tablets or capsules;

(b) the maximum strength of the medicinal product in each tablet or capsule does not exceed 620 mg;

(c) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied in one container or package does not exceed 32; and

(d) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied to a person at any one time does not exceed 100.

4 A medicinal product shall not be a prescription only medicine by reason that it contains the substance aloxiprin, where it is not in the form of a non-effervescent tablet or capsule.

5 A medicinal product shall not be a prescription only medicine by reason that it contains the substance aspirin, where –

(a) the medicinal product is in the form of non-effervescent tablets or capsules;

(b) the maximum strength of the medicinal product in each tablet or capsule does not exceed 75 mg;

(c) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied in one container or package does not exceed 100; and

(d) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied to a person at any one time does not exceed 100.

6 A medicinal product shall not be a prescription only medicine by reason that it contains the substance aspirin, where –

(a) the medicinal product is in the form of non-effervescent tablets or capsules;

(b) the maximum strength of the medicinal product in each tablet or capsule does not exceed 325 mg;

(c) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied in one container or package does not exceed 32; and

(d) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied to a person at any one time does not exceed 100.

7 A medicinal product shall not be a prescription only medicine by reason that it contains the substance azelastine hydrochloride, where –

(a) the medicinal product is in non-aerosol, aqueous form for nasal administration;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 36 doses each of which contains not more than 140 mcg of azelastine hydrochloride;

(c) the container or package is labelled to show a maximum dose of 140 mcg per nostril and a maximum daily dose of 280 mcg per nostril of azelastine hydrochloride; and

(d) the medicinal product is indicated only for the treatment of seasonal allergic rhinitis or perennial allergic rhinitis, in persons aged not less than 5 years.

8 A medicinal product shall not be a prescription only medicine by reason that it contains the substance azelastine hydrochloride, where –

(a) the medicinal product is in the form of eye drops; and

(b) it is indicated only for the treatment of allergic conjunctivitis, in persons aged not less than 12 years.

9 A medicinal product shall not be a prescription only medicine by reason that it contains the substance beclomethasone dipropionate, where –

(a) the medicinal product is in non-aerosol form for nasal administration;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 5,600mcg of beclomethasone diopropionate;

(c) the container or package is labelled to show a maximum dose of 100 mcg per nostril and a maximum daily dose of 200 mcg per nostril of beclomethasone dipropionate; and

(d) the medicinal product is indicated only for the prevention of treatment of allergic rhinitis, in persons aged not less than 12 years.

10 A medicinal product shall not be a prescription only medicine by reason that it contains the substance budesonide, where –

(a) the medicinal product is in non-aerosol, aqueous form for nasal administration;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 10 mg of the medicinal product;

(c) the container or package is labelled to show a maximum dose, and a maximum daily dose, of 200 mcg per nostril of budesonide; and

(d) the medicinal product is indicated only for the prevention of treatment of seasonal allergic rhinitis, in persons aged not less than 12 years.

11 A medicinal product shall not be a prescription only medicine by reason that it contains the substance carbenoxolone sodium, where –

(a) the medicinal product is in the form of granules;

(b) the maximum strength of the carbenoxolone sodium in the medicinal product does not exceed 1%, calculated in terms of weight in weight;

(c) the medicinal product is sold or supplied in a container, or package, containing not more than 560 mg of carbenoxolone sodium;

(d) the container or package is labelled to show a maximum dose of 20 mg and a maximum daily dose of 80 mg of carbenoxolone sodium; and

(e) the medicinal product is indicated only for treatment by mouthwash, in persons aged not less than 12 years.

12 A medicinal product shall not be a prescription only medicine by reason that it contains the substance cetirizine, where –

(a) the medicinal product is sold or supplied in a container, or package, containing not more than 100 mg of cetirizine; and

(b) the container or package is labelled to show a maximum daily dose of 10mg of cetirizine.

13 A medicinal product shall not be a prescription only medicine by reason that it contains the substance cimetidine, where –

(a) the container or package in which the medicinal product is sold or supplied is labelled to show a maximum dose of 200 mg and a maximum daily dose of 800 mg of cimetidine for a maximum period of 14 days; and

(b) the medicinal product is indicated for the short-term symptomatic relief of heartburn, dyspepsia and hyperacidity and for the prophylaxsis of meal-induced heartburn.

14 A medicinal product shall not be a prescription only medicine by reason that it contains the substance cimetidene, where –

(a) the medicinal product is for the prophylactic management of nocturnal heartburn; and

(b) the container or package in which the medicinal product is sold or supplied is labelled to show a maximum dose of 100 mg of cimetidine to be taken once daily at night for a maximum period of 14 days.

15 A medicinal product shall not be a prescription only medicine by reason that it contains the substance clobetasone butyrate, where –

(a) the medicinal product is in the form of a cream;

(b) the maximum strength of the clobetasone butyrate in the medicinal product does not exceed 0.05%;

(c) the medicinal product is sold or supplied in a container, or package, containing not more than 15 g of the medicinal product; and

(d) the medicinal product is indicated only for external application for the short-term treatment of eczema and dermatitis, in persons aged not less than 12 years.

16 A medicinal product shall not be a prescription only medicine by reason that it contains the substance diclofenac diethylammonium, where –

(a) the maximum strength of the diclofenac diethylammonium in the medicinal product does not exceed 1.16%, calculated in terms of weight in weight;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 30 g of the medicinal product;

(c) the container or package is labelled to show a maximum period of use of 7 days; and

(d) the medicinal product is indicated for external application for the local symptomatic relief of pain and inflammation in trauma of the tendons, ligaments, muscles and joints and in localized forms of soft tissue rheumatism, in persons aged not less than 12 years.

17 A medicinal product shall not be a prescription only medicine by reason that it contains the substance domperidone, where –

(a) the medicinal product is indicated for the relief of post-prandial symptoms of excessive fullness, nausea, epigastric bloating and belching, occasionally accompanied by epigastric discomfort and heartburn;

(b) the medicinal product is sold or supplied in a container or package containing not more than 200 mg of domperidone; and

(c) the container or package is labelled to show a maximum dose of 10 mg of domperidone and a maximum daily dose of 40 mg of domperidone.

18 A medicinal product shall not be a prescription only medicine by reason that it contains the substance domperidone maleate, where –

(a) the medicinal product is sold in a container, or package, containing not more than 200 mg of domperidone maleate;

(b) the container or package is labelled to show a maximum dose of 10 mg and a maximum daily dose of 40 mg; and

(c) the medicinal product is indicated for use for the relief of postprandial symptoms of excessive fullness, nausea, epigastric bloating and belching, accompanied by epigastric discomfort and heartburn.

19 A medicinal product shall not be a prescription only medicine by reason that it contains the substance famotidine, where –

(a) the container or package in which the medicinal product is sold or supplied is labelled to show a maximum dose of 10 mg and a maximum daily dose of 20 mg of famotidine for a maximum period of 14 days; and

(b) the medicinal product is indicated for –

(i) the short-term symptomatic relief of heartburn, dyspepsia, indigestion, acid indigestion or hyperacidity, or

(ii) the prevention of the symptoms of heartburn, dyspepsia, indigestion, acid indigestion or hyperacidity where they are associated with the consumption of food or drink, including the prevention of sleep disturbance because of those symptoms.

20 A medicinal product shall not be a prescription only medicine by reason that it contains the substance felbinac, where –

(a) the maximum strength of the felbinac in the medicinal product does not exceed 3.17%, calculated in terms of weight in weight;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 50 g of the medicinal product;

(c) the container or package is labelled to show a maximum period of use of 7 days; and

(d) the medicinal product is indicated for external application for the relief of symptoms associated with soft tissue injury such as strains, sprains and contusions, in persons aged not less than 12 years.

21 A medicinal product shall not be a prescription only medicine by reason that it contains the substance fluconazole, where –

(a) the medicinal product is sold or supplied in a container, or package, containing not more than 150 mg of the medicinal product;

(b) the container or package is labelled to show a maximum dose of 150 mg of fluconazole; and

(c) the medicinal product is indicated for oral administration for the treatment of vaginal candidiasis or associated candidal balanitis, in persons aged not less than 16 years but less than 60 years.

22 A medicinal product shall not be a prescription only medicine by reason that it contains the substance flunisolide, where –

(a) the medicinal product is in the form of a non-pressurized nasal spray;

(b) the maximum strength of the flunisolide in the medicinal product does not exceed 0.025%, calculated in terms of weight in volume;

(c) the medicinal product is sold or supplied in a container, or package, containing not more than 240 metered doses of the medicinal product;

(d) the container or package is labelled to show a maximum dose of 50 mcg per nostril and a maximum daily dose of 100 mcg per nostril of flunisolide in the case of persons aged not less than 16 years, and a maximum dose of 25 mcg per nostril and a maximum daily dose of 75 mcg per nostril in the case of children aged not less than 12 years but less than 16 years; and

(e) the medicinal product is indicated for the prevention and treatment of seasonal allergenic rhinitus, including hay fever, in persons aged not less than 12 years.

23 A medicinal product shall not be a prescription only medicine by reason that it contains the substance hydrocortisone, where –

(a) the maximum strength of the hydrocortisone in the medicinal product does not exceed 0.5%, calculated in terms of weight in weight;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 15 g of the medicinal product; and

(c) the medicinal product is indicated for external use in combination with nystatin of a maximum strength of 3.0%, for intertrigo, in persons aged not less than 10 years.

24 A medicinal product shall not be a prescription only medicine by reason that it contains the substance hydrocortisone, where –

(a) the medicinal product is in the form of a cream, ointment or spray;

(b) the maximum strength of the hydrocortisone in the medicinal product does not exceed 1.0%, calculated in terms of weight in weight;

(c) the medicinal product is sold or supplied in a container, or package, containing –

(i) where the medicinal product is in the form of a cream or ointment, not more than 15 g of the medicinal product, or

(ii) where the medicinal product is in the form of a spray, not more than 30 ml of the medicinal product;

(d) the medicinal product is indicated for external use, either alone or in conjunction with crotamiton in irritant dermatitis, contact allergic dermatitis, insect bite reactions or mild to moderate eczema, and either in combination with clotrimazole or miconazole nitrate for athlete's foot and candidal intertrigo or in combination with lignocaine for anal and perianal itch associated with haemorrhoids; and

(e) the medicinal product is indicated for use in persons aged not less than 10 years.

25 A medicinal product shall not be a prescription only medicine by reason that it contains the substance hydrocortisone acetate, where –

(a) the medicinal product is in the form of a cream or ointment, or suppositories;

(b) the maximum strength of the hydrocortisone acetate in the medicinal product is equivalent to 1.0% of hydrocortisone, calculated in terms of weight in weight;

(c) the medicinal product is sold or supplied in a container, or package, containing –

(i) where the medicinal product is in the form of a cream or ointment, not more than 15 g of the medicinal product, or

(ii) where the medicinal product is in the form of suppositories, not more than 12 suppositories;

(d) the medicinal product is indicated for external use -

(i) for irritant dermatitis, contact allergic dermatitis, insect bite reactions or mild to moderate eczema,

(ii) in combination with one or more of the following, namely benzyl benzoate, bismuth oxide, bismuth subgallate, peru balsam, pramoxine hydrochloride and zinc oxide, for haemorrhoids, or

(iii) in combination with miconazole nitrate, for tinea pedis or candidal intertrigo; and

(e) the medicinal product is indicated for use in persons aged not less than 10 years.

26 A medicinal product shall not be a prescription only medicine by reason that it contains the substance hydrocortisone sodium succinate, where –

(a) the medicinal product is in the form of pellets;

(b) the maximum strength of the hydrocortisone sodium succinate in the medicinal product is equivalent to 2.5 mg of hydrocortisone, calculated in terms of weight in weight;

(c) the medicinal product is sold or supplied in a container, or package, containing the equivalent of 50 mg of hydrocortisone; and

(d) the medicinal product is indicated for external use for aphthous ulceration of the mouth, in persons aged not less than 12 years.

27 A medicinal product shall not be a prescription only medicine by reason that it contains the substance hydroxyzine hydrochloride, where –

(a) the medicinal product is sold or supplied in a container, or package, containing not more than 750 mg of the medicinal product;

(b) the container or package is labelled to show a maximum dose of 25 mg, and to show a maximum daily dose of 75 mg in the case of persons aged not less than 12 years and a maximum daily dose of 50 mg in the case of children aged not less than 6 years but less than 12 years; and

(c) the medicinal product is indicated for the management of pruritus associated with acute or chronic urticaria or atopic dermatitis or contact dermatitis, in persons aged not less than 6 years.

28 A medicinal product shall not be a prescription only medicine by reason that it contains the substance hyoscine butylbromide, where –

(a) the route of administration of the medicinal product is internal and is otherwise than by means of an inhaler;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 240 mg of the medicinal product; and

(c) the container or package is labelled to show a maximum dose of 20 mg and a maximum daily dose of 80 mg of the medicinal product.

29 A medicinal product shall not be a prescription only medicine by reason that it contains the substance Ibuprofen, where –

(a) the medicinal product is indicated for the relief of rheumatic and muscular pain, pain of non-serious arthritic conditions, backache, neuralgia, migraine, headache, dental pain, dysmenorrhoea, feverishness, symptoms of colds and influenza; and either

(b) the route of the administration of the medicinal product is internal; and –

(i) in the case of a prolonged release preparation the container or package is labelled to show a maximum dose of 600 mg and a maximum daily dose of 1200 mg, or

(ii) in any other case the container or package is labelled to show a maximum dose of 400 mg and a maximum daily dose of 1200 mg; or

(c) the route of administration of the medicinal product is external; and

(i) the maximum strength of the Ibuprofen in the medicinal product does not exceed 5%, or

(ii)

(A) the maximum strength of the Ibuprofen in the medicinal product does not exceed 10%, and

(B) the medicinal product is sold or supplied in a container or package containing not more than 50 g of medicinal product which is labelled to show a maximum dose of 125 mg and a maximum daily dose of 500 mg.

30 A medicinal product shall not be a prescription only medicine by reason that it contains the substance ketoconazole, where –

(a) the medicinal product is in the form of a shampoo;

(b) the maximum strength of the ketoconazole in the medicinal product does not exceed 2%, calculated in terms of weight in weight;

(c) the medicinal product is sold or supplied in a container, or package, containing not more than 120 ml of the medicinal product and containing in the medicinal product not more than 2,400 mg of ketoconazole;

(d) the container or package is labelled to show a maximum frequency of application of once every 3 days; and

(e) the medicinal product is indicated for the prevention and treatment of dandruff and seborrhoeic dermatitis of the scalp.

31 A medicinal product shall not be a prescription only medicine by reason that it contains the substance ketoprofen, where –

(a) the maximum strength of the ketoprofen in the medicinal product does not exceed 2.5%, calculated in terms of weight in weight;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 30 g of the medicinal product; and

(c) the medicinal product is indicated only for treatment by external topical application, for rheumatic and muscular pain, in persons aged not less than 12 years, for a maximum period of 7 days.

32 A medicinal product shall not be a prescription only medicine by reason that it contains the substance levocabastine hydrochloride, where –

(a) the medicinal product is in the form of a nasal spray;

(b) the maximum strength of the medicinal product does not exceed the equivalent of 0.05% levocabastine;

(c) the medicinal product is sold or supplied in a container, or package, containing not more than 10 ml of the medicinal product; and

(d) the medicinal product is indicated for the symptomatic treatment of seasonal allergic rhinitis.

33 A medicinal product shall not be a prescription only medicine by reason that it contains the substance levocabastine hydrochloride, where –

(a) the medicinal product is in the form of aqueous eye drops;

(b) the maximum strength of the medicinal product does not exceed the equivalent of 0.05% levocabastine;

(c) the medicinal product is sold or supplied in a container, or package, containing not more than 4ml of the medicinal product; and

(d) the medicinal product is indicated for the symptomatic treatment of seasonal allergic conjunctivitis.

34 A medicinal product shall not be a prescription only medicine by reason that it contains the substance loratadine, where –

(a) the medicinal product is sold or supplied in a container, or package, containing not more than 100 mg of loratadine; and

(b) the container or package is labelled to show a maximum daily dose of 10 mg of loratadine.

35 A medicinal product shall not be a prescription only medicine by reason that it contains the substance mebendazole, where –

(a) the medicinal product is sold or supplied in a container, or package, containing not more than 800 mg of mebendazole;

(b) the container or package is labelled to show a maximum dose of 100 mg of mebendazole; and

(c) the medicinal product is indicated for oral use in the treatment of enterobiasis, in persons aged not less than 2 years.

36 A medicinal product shall not be a prescription only medicine by reason that it contains the substance nedocromil sodium, where –

(a) the maximum strength of the nedocromil sodium in the medicinal product does not exceed 2.0%, calculated in terms of weight in volume;

(b) the medicinal product is sold in a container, or package, containing not more than 3 ml of the medicinal product; and

(c) the medicinal product is indicated for the prevention, relief and treatment of seasonal and perennial allergic conjunctivitis.

37 A medicinal product shall not be a prescription only medicine by reason that it contains the substance nizatidine, where –

(a) the container or package in which the medicinal product is sold or supplied is labelled to show a maximum dose of 75 mg of nizatidine and a maximum of 4 such doses in any period of 14 days; and

(b) the medicinal product is indicated only for the prevention of the symptoms of food-related heartburn, in persons aged not less than 16 years.

38 A medicinal product shall not be a prescription only medicine by reason that it contains the substance nystatin, where –

(a) the maximum strength of the nystatin in the medicinal product does not exceed 3.0%, calculated in terms of weight in weight;

(b) the medicinal product is sold in a container, or package, containing not more than 15 g of the medicinal product; and

(c) the medicinal product is indicated for external use in combination with hydrocortisone of a maximum strength of 0.5% for intertrigo, in persons aged not less than 10 years.

39 A medicinal product shall not be a prescription only medicine by reason that it contains the substance oxethazaine, where –

(a) the medicinal product is sold or supplied in a container, or package, containing not more than 400 ml of oxethazaine; and

(b) the container or package is labelled to show a maximum dose of 10 ml and a maximum daily dose of 30 ml of oxethazaine.

40 A medicinal product shall not be a prescription only medicine by reason that it contains the substance paracetamol, where –

(a) the medicinal product is in the form of non-effervescent tablets or capsules;

(b) the maximum strength of the medicinal product in each tablet or capsule does not exceed 500 mg;

(c) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied in one container or package does not exceed 32;

(d) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied to a person at any one time does not exceed 100; and

(e) the medicinal product is indicated for use by administration wholly or mainly to persons aged not less than 12 years.

41 A medicinal product shall not be a prescription only medicine by reason that it contains the substance paracetamol, where –

(a) the medicinal product is in the form of non-effervescent tablets or capsules;

(b) the maximum strength of the medicinal product in each tablet or capsule does not exceed 250 mg;

(c) the quantity (of tablets or capsules, or of any combination of tablets and capsules) that is sold or supplied in one container or package does not exceed 32; and

(d) the quantity (of tablets and capsules, or of any combination of tablets and capsules) that is sold or supplied to a person at any one time does not exceed 100.

42 A medicinal product shall not be a prescription only medicine by reason that it contains the substance piroxicam, where –

(a) the maximum strength of the piroxicam in the medicinal product does not exceed 0.5%;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 30 mg of the medicinal product;

(c) the container or package is labelled to show a maximum period of use of 7 days; and

(d) the medicinal product is indicated for external application for the relief of rheumatic pain, pain of non-serious arthritic conditions and muscular aches, pains and swellings such as strains, sprains and sports injuries, in persons aged not less than 12 years.

43 A medicinal product shall not be a prescription only medicine by reason that it contains the substance prochlorperazine maleate, where –

(a) the medicinal product is in the form of tablets;

(b) the maximum amount of the prochlorperazine maleate in the medicinal product, in each tablet, does not exceed 3 mg;

(c) the medicinal product is sold or supplied in a container, or package, containing not more than 8 tablets; and

(d) the medicinal product is indicated only for nausea, and vomiting, in previously diagnosed migraine, in persons aged not less than 18 years.

44 A medicinal product shall not be a prescription only medicine by reason that it contains the substance pyrantel embonate, where –

(a) the medicinal product is sold or supplied in a container, or package, containing not more than 750 mg of the medicinal product;

(b) the container or package is labelled to show a maximum daily dose (to be taken as a single dose) of pyrantel embonate of 750 mg in the case of persons aged not less than 12 years, of 500 mg in the case of children aged not less than 6 years but less than 12 years, and of 250 mg in the case of children aged not less than 2 years but less than 6 years; and

(c) the medicinal product is indicated for the treatment of enterobiasis, in persons aged not less than 2 years.

45 A medicinal product shall not be a prescription only medicine by reason that it contains the substance ranitidine hydrochloride, where –

(a) the container or package in which the medicinal product is sold or supplied is labelled to show a maximum dose equivalent to 75 ml and a maximum daily dose equivalent to 300 ml of ranitidine for a maximum period of use of 14 days; and

(b) the medicinal product is indicated for the short-term symptomatic relief of heartburn, dyspepsia and hyperacidity, or the prevention of those symptoms when associated with the consumption of food and drink.

46 A medicinal product shall not be a prescription only medicine by reason that it contains the substance sodium cromoglycate, where –

(a) the medicinal product is in the form of aqueous eye drops;

(b) the maximum strength of the sodium cromoglycate in the medicinal product does not exceed 2%, calculated in terms of weight in volume;

(c) the medicinal product is sold or supplied in a container containing not more than 10 ml of the medicinal product; and

(d) the medicinal product is indicated for treatment of acute seasonal allergic conjunctivitis.

47 A medicinal product shall not be a prescription only medicine by reason that it contains the substance sodium cromoglycgate, where –

(a) the medicinal product is in the form of an eye ointment;

(b) the maximum strength of the sodium cromoglycate in the medicinal product is 4%, calculated in terms of weight in weight;

(c) the medicinal product is sold or supplied in a container, or package, containing not more than 5 g of the medicinal product; and

(d) the medicinal product is indicated for the treatment of acute seasonal allergic conjunctivitis or perennial allergic conjunctivitis.

48 A medicinal product shall not be a prescription only medicine by reason of the fact that it contains terbinafine, where –

(a) the maximum strength of the terbinafine in the medicinal product does not exceed 1%;

(b) the medicinal product is sold or supplied in a container, or package, containing not more than 30 g of the medicinal product; and

(c) the medicinal product is indicated for external use as a gel for the treatment of tina corporis, tinea pedis and tinea cruris.

49 A medicinal product shall not be a prescription only medicine by reason of the fact that it contains terbinafine hydrochloride where –

(a) the maximum strength of the terbinafine hydrochloride in the medicinal product does not exceed 1%;

(b) the medicinal product is indicated for external use for the treatment of tinea pedis and tinea cruris; and

(c) the medicinal product is sold or supplied in a container or package containing not more than 15 g of medicinal product.

50 A medicinal product shall not be a prescription only medicine by reason of the fact that it contains terbinafine hydrochloride, where –

(a) the maximum strength of the terbinafine hydrochloride in the medicinal product does not exceed 1%;

(b) the medicinal product is sold or supplied in a container containing not more than 30 ml of the medicinal product; and

(c) the medicinal product is indicated for external use as a spray solution for the treatment of tina corporis, tinea pedis and tinea cruris.

51 A medicinal product shall not be a prescription only medicine by reason of the fact that it contains triamcinolone acetonide where –

(a) the medicinal product is in the form of a non-pressurised nasal spray;

(b) the medicinal product is indicated for the treatment of symptoms of seasonal allergic rhinitis in persons aged 18 years and over for a maximum period of 3 months;

(c) the container or package is labelled to show a maximum dose of 110 mcg per nostril and a maximum daily dose of 110 mcg per nostril; and

(d) the medicinal product is sold or supplied in a container or package containing not more than 3.375 mg of triamcinolone acetonide.

SCHEDULE 2

EXEMPTION FOR CERTAIN PERSONS FROM ARTICLE 57(2) OF THE LAW

PART 1[13]

(Articles 1(2)(a) and 8(1))

Column 1	*Column 2*	*Column 3*
Persons exempted	**Prescription only medicines to which the exemption applies**	**Conditions**
1. Persons selling or supplying prescription only medicines to universities, other institutions concerned with higher education or institutions concerned with research.	1. All prescription only medicines.	1.(1) The sale or supply shall be subject to the presentation of an order, signed by the principal of the institution or the appropriate head of department in charge of a specified course of research.
		(2) The order shall specify–
		(a) the name of the institution for which the prescription only medicine is required;
		(b) the purpose for which the prescription only medicine is required; and
		(c) the total quantity required.
		(3) The sale or supply shall be only for the purposes of the education or research with which the institution is concerned.
2. Persons selling or supplying prescription only medicines to any of the following persons – (a) the Official Analyst appointed under the Official Analyst (Jersey) Law 2022, or any person appointed under that Law to deputise for the Official Analyst;	2. All prescription only medicines.	2.(1) The sale or supply shall be subject to the presentation of an order signed by or on behalf of any person listed in any of paragraph 2(a), (b) and (c) of column 1 of this Part of this Schedule. (2) The order shall specify the status of the person signing it, and the amount

Column 1	*Column 2*	*Column 3*
Persons exempted	**Prescription only medicines to which the exemption applies**	**Conditions**
		of the prescription only medicine required.
(b) an authorized officer within the meaning of the Food Safety (Jersey) Law 1966; and		(3) The sale or supply shall be only in connection with the exercise by the person of his or her statutory functions.
(c) a person duly authorized by the Minister under Article 96 or 97 of the Law.		
3. Persons selling or supplying prescription only medicines to any person employed or engaged in connection with a scheme for testing the quality and checking the amount of drugs and appliances supplied under the Health Insurance (Jersey) Law 1967, or under any subordinate legislation made under that law.	3. All prescription only medicines.	3.(1) The sale or supply shall be subject to the presentation of an order signed by or on behalf of the person so employed or engaged. (2) The order shall specify the status of the person signing it, and the amount of the prescription only medicine required.
		(3) The sale or supply shall be only for the purposes of a scheme to which paragraph 3 of column 1 of this Part of this Schedule refers.
4. Certified midwives.	4. Prescription only medicines containing any of the following substances – Chloral hydrate Dichloral-phenazone Ergometrine maleate Pentazocine hydrochloride Phytomenadrone Triclofos sodium	4. The sale or supply shall be only in the course of the midwife's professional practice and, in the case of Ergometrine maleate, only when contained in a medicinal product that is not for parenteral administration.
5. Persons lawfully conducting retail pharmacy businesses.	5. Prescription only medicines (not being for parenteral administration) that are of any of the following descriptions –	5. The sale or supply shall be subject to the presentation of an order signed by a registered optometrist.
	(a) eye drops, or eye ointments, that are prescription only medicines	

Column 1	*Column 2*	*Column 3*
Persons exempted	**Prescription only medicines to which the exemption applies**	**Conditions**
	by reason only that they contain – (i) 30.0% Sulphacetamide sodium; or (ii) 0.5% Chloramphenicol;	
	(b) eye ointments that are prescription only medicines by reason only that they contain – (i) 30.0% Sulpha-cetamide sodium; or (ii) 0.5%Chloramphenicol; or	
	(c) medicinal products that are prescription only medicines by reason only that they contain any of the following substances – Atropine sulphate Bethanecol chloride Carbachol Cyclopentolate hydrochloride Homatropine hydrobromide Hyoscine hydrobromide Naphazoline hydrochloride Naphazoline nitrate Neostigmine methyl-sulphate Physostigmine salicylate Physostigmine sulphate Pilocarpine hydrochloride Pilocarpine nitrate Tropicamide.	
6. Registered optometrists.	6. Prescription only medicines listed in paragraph 5 of column 2 of this Part of this Schedule.	6.(1) The sale or supply shall be only in the course of the optician's professional practice. (2) The sale shall be only in an emergency.
7.(1) Holders of product licences.	7. Prescription only medicines to which the licence relates.	7. The sale or supply shall be only –
(2) Holders of		(a) to a pharmacist, so as to

Column 1	*Column 2*	*Column 3*
Persons exempted	**Prescription only medicines to which the exemption applies**	**Conditions**
manufacturer's licences.		enable that pharmacist to prepare an entry relating to the prescription only medicine in question in a tablet or capsule identification guide or similar publication; and
		(b) of no greater quantity than is reasonably necessary for that purpose.
8. Pharmacists selling or supplying to persons to whom cyanide salts may be sold lawfully under the Poisons (Jersey) Law 1952.	8. Amyl nitrite.	8. The sale or supply shall only be so far as is necessary to enable an antidote to be available to persons at risk of cyanide poisoning.

PART 2

(Articles 1(2) and 8(1))

Column 1 **Persons exempted**	Column 2 **Prescription only medicines to which the exemption applies**	Column 3 **Conditions**
1.(1) The Royal National Lifeboat Institution. (2) Certificated first aiders of the Institution.	1. All prescription only medicines.	1. The supply shall be only so far as is necessary for the treatment of sick or injured persons.
2. The owner or the master of a ship that does not carry a doctor on board as part of the ship's complement.	2. All prescription only medicines.	2. The supply shall be only so far as is necessary for the treatment of persons on the ship.
3. The operator or commander of an aircraft.	3. Prescription only medicines that – (a) are not for parenteral administration; and (b) have been sold or supplied to the operator or commander of the aircraft in response to an order in writing signed by a doctor.	3.(1) The supply shall be only so far as is necessary for the immediate treatment of sick or injured persons on the aircraft. (2) The supply shall be in accordance with the written instructions of a doctor as to the circumstances in which prescription only medicines of the description in question are to be used on the aircraft.
4. Persons authorized by licences granted under Article 4 of the Misuse of Drugs (General Provisions) (Jersey) Order 1989 to supply a controlled drug.	4. Prescription only medicines (being controlled drugs) whose supply is authorized by the licence.	4. The supply shall be subject to the conditions, in the circumstances and to the extent specified in the licence.

Column 1	Column 2	Column 3
5. Persons requiring prescription only medicines to enable them, in the course of any business carried on by them, to comply with any requirements under any enactment in respect of the medical treatment of their employees.	5. Prescription only medicines specified in the enactment.	5.(1) The supply shall be only to enable the person to comply with any such requirements. (2) The supply shall be subject to such conditions and in such circumstances as may be specified in the enactment.
6. Persons operating an occupational health scheme.	6. Prescription only medicines sold or supplied to such a person in response to an order in writing signed by a doctor or a registered nurse.	6.(1) The supply shall be only in the course of the scheme. (2) The person supplying the prescription only medicine shall be – (a) a doctor; or
		(b) a registered nurse acting in accordance with the written instructions of a doctor as to the circumstances in which prescription only medicines of the description in question are to be used in the course of the scheme.

PART 3[14]

(Articles 1(2) and 8(2))

Column 1	Column 2	Column 3
Persons exempted	Prescription only medicines to which the exemption applies	Conditions
1. Chiropodists, registered under the Health Care (Registration) (Jersey) Law 1995, who hold certificates of competence in the use of analgesics issued by or with the approval of the Chiropodists Board of the United Kingdom.	1. Prescription only medicines for parenteral administration that contain, as the sole active ingredient, not more than one of the following substances –	1. The administration shall be only in the course of the chiropodist's professional practice.
	Bupivacaine hydrochloride	
	Bupivacaine hydrochloride with adrenaline, where the maximum strength of the adrenaline does not exceed 1 mg in 200 ml of bupivacaine hydrochloride	
	Lignocaine hydrochloride	
	Lignocaine hydrochloride with adrenaline, where the maximum strength of the adrenaline does not exceed 1 mg in 200 ml of lignocaine hydrochloride	
	Mepivacaine hydrochloride	
	Prilocaine hydrochloride	
2. Certified midwives.	2. Prescription only medicines for parenteral administration that contain any of the following substances (but no other substance specified in column 1 of Part I of the First Schedule to this Order) –	2. The administration shall be only in the course of the midwife's professional practice and, in the case of Lignocaine, Lignocaine hydrochloride and Promazine hydrochloride, shall be only while attending on a woman in childbirth.
	Ergometrine maleate	
	Levallorphan tartrate	
	Lignocaine	
	Lignocaine hydrochloride	
	Naloxone hydrochloride	
	Oxytocins,	

Column 1	Column 2	Column 3
	Natural and Synthetic	
	Pentazocine lactate	
	Pethidine	
	Pethidine hydrochloride	
	Phytomenadione	
	Promazine hydrochloride.	
3. The owner or the master of a ship that does not carry a doctor on board as part of the ship's complement.	3. All prescription only medicines that are for parenteral administration.	3. The administration shall be only so far as is necessary for the treatment of persons on the ship.
4. The operator or commander of an aircraft.	4. Prescription only medicines for parenteral administration that have been sold or supplied to the operator or commander of the aircraft in response to an order in writing signed by a doctor.	4.(1) The administration shall be only so far as is necessary for the immediate treatment of sick or injured persons on the aircraft.
		(2) The administration shall be in accordance with the written instructions of a doctor as to the circumstances in which prescription only medicines of the description in question are to be used on the aircraft.
5. Persons operating an occupational health scheme.	5. Prescription only medicines for parenteral administralion that have been sold or supplied to such a person in response to an order in writing signed by a doctor or a registered nurse	5.(1) The administration shall be only in the course of the scheme. (2) The person administering the prescription only medicine shall be – (a) a doctor;
		(b) a person acting in accordance with the directions of a doctor; or
		(c) a registered nurse acting in accordance with the written instructions of a doctor as to the circumstances in which prescription only medicines of the description in question are to be used in the course of the scheme.
6. Persons who hold certificates of proficiency	6. The following prescription only	6. The administration shall be only for the immediate,

Column 1	Column 2	Column 3
in ambulance paramedical skills issued by or with the approval of the Secretary of State of the United Kingdom, or persons who are state registered paramedics.	medicines for parenteral administration –	necessary treatment of sick or injured persons and, in the case of a prescription only medicine containing Heparin sodium, shall be only for the purpose of cannula flushing.
	(a) Diazemuls (product licence number 10183/00001);	
	(b) Gelofusine (product licence number 00183/5025R); and	
	(c) medicines containing the substances Ergometrine Maleate 500mcg per ml with Oxytocin 5 iu per ml, but no other active ingredient:	
	(d) prescription only medicines that contain one or more of the following substances, (but no other active ingredient)-	
	Adrenaline acid tartrate	
	Benzylpenicillin	
	Frusemide	
	Glucose	
	Heparin sodium	
	Lignocaine hydrochloride	
	Metoclopramide	
	Morphine Sulphate	
	Nalbuphine hydrochloride	
	Naloxone hydrochloride	
	Polygeline	
	Sodium bicarbonate	
	Sodium chloride.	
	Streptokinase	

SCHEDULE 3

(Articles 9(3)(c) and (4)(a))

SUBSTANCES THAT MUST NOT BE CONTAINED IN A PRESCRIPTION ONLY MEDICINE EXEMPTED BY ARTICLE 9

Ammonium Bromide	
Amylobarbitone	
Amylobarbitone Sodium	
Barbitone	
Barbitone Sodium	
Butobarbitone	
Butobarbitone Sodium	
Calcium Bromide	
Calcium Bromidolactobionate	
Cyclobarbitone	
Cyclobarbitone Calcium	
Embutramide	
Fencamfamin Hydrochloride	
Fluanisone	
Heptabarbitone	
Hexobarbitone	
Hexobarbitone Sodium	
Hydrobromic Acid	
Meclofenoxate Hydrochloride	
Methohexitone Sodium	
Methylphenobarbitone	
Pemoline	
Pentobarbitone	
Pentobarbitone Sodium	
Phenobarbitone	
Phenobarbitone Sodium	
Phenylmethylbarbituric Acid	
Piracetam	
Potassium Bromide	
Prolintane Hydrochloride	
Quinalbarbitone	
Quinalbarbitone Sodium	
Quinidine Phenylethylbarbiturate	
Secbutobarbitone	
Secbutobarbitone Sodium	
Sodium Bromide	
Strychnine Hydrochloride	
Tacrine Hydrochloride	
Thiopentone Sodium	
Note (for information):	The restriction in Article 9(3)(c) is subject to Article 9(4), in respect of Phenobarbitone and Phenobarbitone Sodium

Ammonium Bromide	
	for use in the treatment of epilepsy.

SCHEDULE 4[15]

(Articles 6(2)(ca) and 6A(2))

CLINICAL MANAGEMENT PLAN

1 Information to be included in clinical management plan

A clinical management plan must include –

(a) the name of the patient to whom the plan relates;

(b) the illness or conditions in relation to which the supplementary prescriber may give a prescription or administer (or direct the administration of) a medicinal product;

(c) the date on which the plan is to take effect and the date or dates when it is subject to review by the doctor or dentist who is a party to the plan;

(d) the class or description of medicinal product that may be prescribed by a supplementary prescriber or administered by, or under the direction of, a supplementary prescriber;

(e) any restrictions or limitations as to the strength or dose, or period of use, of any medicinal product which may be prescribed by or administered by, or under the direction of, the supplementary prescriber;

(f) any relevant warnings about the known sensitivities of the patient to, or known difficulties of the patient with, particular medicinal products;

(g) arrangements for the notification of suspected or known adverse reactions to –

(i) the medicinal product referred to in paragraph (d), and

(ii) any other medicinal product taken at the same time or over the same period;

(h) the circumstances in which the supplementary prescriber should refer to, or seek the advice of, the doctor or dentist who is a party to the plan.

ENDNOTES

Table of Legislation History

Legislation	Year and No	Commencement
Medicines (Prescription Only) (Jersey) Order 1997	R&O.9140	1 January 1998
Medicines (Prescription Only) (Amendment) (Jersey) Order 1998	R&O.9326	1 January 1999
Medicines (Prescription Only) (Amendment No. 2) (Jersey) Order 2000	R&O.1/2000	1 February 2000
Medicines (Prescription Only) (Amendment No. 3) (Jersey) Order 2001	R&O.2/2001	1 February 2001
Medicines (Prescription Only) (Amendment No. 4) (Jersey) Order 2002	R&O.94/2002	1 October 2002
Medicines (Prescription Only) (Amendment No. 5) (Jersey) Order 2003	R&O.75/2003	13 August 2003
Medicines (Prescription Only) (Amendment No. 6) (Jersey) Order 2004	R&O.65/2004	12 July 2004
Medicines (Prescription Only) (Amendment No. 7) (Jersey) Order 2005	R&O.174/2005	11 November 2005
States of Jersey (Amendments and Construction Provisions No. 5) (Jersey) Regulations 2005	R&O.45/2005	9 December 2005
Pharmacists and Pharmacy Technicians (Registration) (Jersey) Law 2010	L.6/2010	16 May 2010
Medicines (Prescription Only) (Amendment No. 8) (Jersey) Order 2013	R&O.82/2013	1 July 2013
Opticians (Registration) (Amendment No.2) (Jersey) Law 2017	L.13/2017	19 May 2017
Data Protection (Jersey) Law 2018	L.3/2018	25 May 2018
Medicines (Prescription Only) (Amendment No. 9) (Jersey) Order 2019	R&O.108/2019	15 October 2019
Official Analyst (Jersey) Law 2022	L.30/2022	12 August 2022

Table of Renumbered Provisions

Original	Current

Original	Current
3(3)	revoked by R&O.9326
10A	11
11	12
12	13
13	14
14	15
15	16
FIRST SCHEDULE	SCHEDULE 1
PART I	PART 1
PART II	PART 2
PART III	PART 3
PART IV	PART 4
7A	8
8	9
9	10
10	11
11	12
12	13
13	14
13A	15
14	16
14A	17
15	18
16	19
17	20
18	21
19	22
20	23
21	24
22	25
23	26
24	27
25	28
25A	29
26	30
27	31
28	32
29	33
30	34
31	35
32	36
33	37
34	38
35	39
36	40
37	41
38	42

Original	Current
38A	43
39	44
40	45
41	46
42	47
42A	48
43	49
43A	50
44	51
SECOND SCHEDULE	SCHEDULE 2
PART I	PART 1
PART II	PART 2
PART III	PART 3
6(bb)	6(c)
6(c)	6(d)
THIRD SCHEDULE	SCHEDULE 3

Table of Endnote References

[1]	*This Order has been amended by the States of Jersey (Amendments and Construction Provisions No. 5) (Jersey) Regulations 2005. The amendments replace all references to a Committee of the States of Jersey with a reference to a Minister of the States of Jersey, and remove and add defined terms appropriately, consequentially upon the move from a committee system of government to a ministerial system of government*
[2] *Article 1(1)*	*amended by R&O.2/2001, R&O.82/2013, L.13/2017, L.3/2018, R&O.108/2019*
[3] *Article 2*	*amended by R&O.82/2013*
[4] *Article 5*	*substituted by R&O.82/2013, R&O.108/2019*
[5] *Article 6*	*heading amended by R&O.82/2013*
[6] *Article 6(2)*	*amended by R&O.82/2013*
[7] *Article 6A*	*inserted by R&O.82/2013*
[8] *Article 9(3)*	*amended by R&O.9326, R&O.1/2000*
[9] *Article 10*	*substituted by R&O.82/2013*
[10] *Article 11*	*inserted by R&O.65/2004*
[11] *Schedule 1*	*Part 1 amended by R&O.9326, R&O.1/2000, R&O.2/2001, R&O.94/2002, R&O.174/2005*
[12] *Schedule 1*	*Part 4 substituted by R&O.1/2000, amended by R&O.2/2001, R&O.94/2002,*
[13] *Schedule 2*	*Part 1 amended by R&O.1/2000, L.6/2010, L.13/2017, L.30/2022*
[14] *Schedule 2*	*Part 3 amended by R&O.9326, R&O.1/2000, R&O.2/2001*
[15] *Schedule 4*	*inserted by R&O.82/2013*