STATUTORY INSTRUMENTS

# 1977 No. 2133

# MEDICINES

## The Medicines (Pharmacy and General Sale—Exemption) Order 1977

| | |
|---|---|
| *Made* - - - - | *20th December* 1977 |
| *Laid before Parliament* | *5th January* 1978 |
| *Coming into Operation* | *1st February* 1978 |

The Secretaries of State respectively concerned with health in England and in Wales, the Secretary of State concerned with health and with agriculture in Scotland, the Minister of Agriculture, Fisheries and Food, the Department of Health and Social Services for Northern Ireland and the Department of Agriculture for Northern Ireland, acting jointly, in exercise of powers conferred by section 57(1) of the Medicines Act 1968**(a)** and now vested in them **(b)** and the Secretaries of State concerned with health in England, in Wales and in Scotland and the Department of Health and Social Services for Northern Ireland, acting jointly, in exercise of powers conferred by section 55(2)(*b*) of that Act and now vested in them **(b)** and of all other powers enabling them in that behalf, after consulting such organisations as appear to them to be representative of interests likely to be substantially affected by the following order, and after taking into account the advice of the Medicines Commission, hereby make the following order:—

*Citation, commencement and interpretation*

**1.**—(1) This order may be cited as the Medicines (Pharmacy and General Sale—Exemption) Order 1977 and shall come into operation on 1st February 1978.

(2) In this order, unless the context otherwise requires—
"the Act" means the Medicines Act 1968;

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**(a)** 1968 c. 67.

**(b)** In the case of the Secretaries of State concerned with health in England and in Wales by virtue of Article 2(2) of, and Schedule 1 to, the Transfer of Functions (Wales) Order 1960 (S.I. 1969/388 (1969 I, p. 1070)), and in the case of the Northern Ireland Departments by virtue of section 40 of, and Schedule 5 to, the Northern Ireland Constitution Act 1973 (c. 36), and section 1(3) of, and paragraph 2(1)(*b*) of Schedule 1 to, the Northern Ireland Act 1974 (c. 28).

"the appointed day" means 1st February 1978 being the day appointed for the purposes of section 52 of the Act(a);

"controlled drug" has the meaning assigned to it by section 2 of the Misuse of Drugs Act 1971(b);

"cosmetic" means any substance or preparation intended to be applied to the various surfaces of the human body including epidermis, pilary system and hair, nails, lips and external genital organs, or the teeth and buccal mucosa wholly or mainly for the purpose of perfuming them, cleansing them, protecting them, caring for them or keeping them in condition, modifying their appearance (whether for aesthetic purposes or otherwise) or combating body odours or normal body perspiration;

"external use" means—

(*a*) in relation to medicinal products for use by being administered to human beings, application to the skin, teeth, mucosa of the mouth, throat, nose, ear, eye, vagina or anal canal,

(*b*) in relation to veterinary drugs, application to the skin, hair, fur, feathers, scales, hoof, horn, ear, eye, mouth or mucosa of the throat or prepuce,

in either case when a local action only is necessary and extensive systemic absorption is unlikely to occur and references to "medicinal products for external use" shall be read accordingly except that in relation to paragraph (*a*) above the references shall not include throat sprays, throat pastilles, throat lozenges, throat tablets, nasal drops, nasal sprays, nasal inhalations or teething preparations;

"food" includes beverages, confectionery and articles and substances used as ingredients in the preparation of food and includes any manufactured substance to which there has been added any vitamin and which is advertised (within the meaning of section 92 of the Act) as available and for sale to the general public as a dietary supplement;

"master" has the same meaning as in the Merchant Shipping Act 1894(c);

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

"pharmacy medicine" means any medicinal product which is not a prescription only medicine or a medicinal product on a general sale list;

"prescription only medicine" means a medicinal product of a description or falling within a class specified in Article 3 of the Medicines (Prescription Only) Order 1977(d);

"registered ophthalmic optician" means a person who is registered in either of the registers of ophthalmic opticians established and maintained under section 2(*a*) of the Opticians Act 1958(e);

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(a) The Medicines (Pharmacy and General Sale) (Appointed Day) Order 1977 (S.I. 1977/2126).(1977 III, P. 5818)).
(b) 1971 c. 38.
(c) 1894 c. 60.
(d) S.I. 1977/2127 (1977 III, p. 5820).
(e) 1958 c. 32.

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

"state registered chiropodist" means a person who is registered in the register established and maintained under section 2(1) of the Professions Supplementary to Medicine Act 1960(**a**) by the Chiropodists Board;

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

"unit preparation" means a preparation, including a mother tincture, prepared by a process of solution, extraction or trituration with a view to being further diluted tenfold, or serially in multiple powers of ten, in an inert diluent, and then used either in this diluted form or, where applicable, by impregnating tablets, granules, powders or other inert substances for the purpose of being administered to human beings;

and other expressions shall have the same meanings as in the Act.

(3) Except in so far as the context otherwise requires, any reference in this order to any enactment, order or regulations shall be construed as a reference to that enactment or order or to those regulations, as the case may be, as amended, extended or re-enacted by any other enactment, order or regulations.

(4) The rules for the construction of Acts of Parliament contained in the Interpretation Act 1889(**b**) shall apply for the purposes of the interpretation of this order as they apply for the purposes of the interpretation of an Act of Parliament.

*Transitional exemptions for products for human use*

**2.**—(1) The restrictions imposed by section 52 of the Act (sale or supply of medicinal products not on a general sale list) shall not apply for a period of two years from the appointed day to any person who sells, offers or exposes for sale or supplies a medicinal product which is not a medicinal product on a general sale list if and so long as the conditions specified in paragraph (2) of this Article are satisfied.

(2) The conditions referred to in the preceding paragraph are that—

(*a*) the medicinal product in question—

(i) is for use by being administered to human beings, and

(ii) is one in respect of which a product licence has been granted under Part II of the Act;

(*b*) the person selling or supplying the medicinal product in question could lawfully have sold or supplied that product immediately before the appointed day; and

(*c*) the conditions specified in section 53 of the Act are fulfilled.

*Temporary exemption for certain products for human use*

**3.**—(1) The restrictions imposed by section 52 of the Act shall not apply during the period set out in paragraph (2) below to the sale, offer or exposure for sale or supply of any medicinal product which is for use by being administered to human beings and either—

(**a**) 1960 c. 66. (**b**) 1889 c. 63.

(*a*) in respect of which a product licence has been granted after the appointed day containing a provision to the effect that the method of sale or supply of the medicinal product in question may be otherwise than by or under the supervision of a pharmacist, or

(*b*) in respect of which a product licence has been varied after the appointed day so as to contain a provision to the effect that the method of sale or supply may be otherwise than by or under the supervision of a pharmacist

if and so long as the conditions specified in section 53 of the Act are fulfilled.

(2) The period referred to in the preceding paragraph is the period starting with the date on which the product licence is granted or varied and ending one year from the date on which a notice that such a licence has been granted or varied, as the case may be, is published in the Gazette.

*Exemption in cases involving another's default*

**4.** The restrictions imposed by section 52 of the Act shall not apply to the sale, offer or exposure for sale or supply of a medicinal product by a person who, having exercised all due diligence, believes on reasonable grounds that the product is a medicinal product on a general sale list but which due to the act or default of another person is not such a medicinal product if and so long as the conditions specified in section 53 of the Act are fulfilled.

*Exemption for products used by midwives in the course of their professional practice*

**5.** There are hereby specified for the purposes of section 55(2)(*b*) of the Act (exemptions for certified midwives) the following descriptions and classes of medicinal products—

(1) any medicinal product which may lawfully be sold or supplied other than in accordance with a prescription given by a practitioner,

(2) any medicinal product which may lawfully be sold or supplied only in accordance with a prescription given by a practitioner but which, by virtue of an exemption conferred by section 58(4)(*a*) of the Act, may be sold or supplied by a certified midwife other than in accordance with such a prescription, and

(3) any medicinal product on a general sale list.

*Exemption for certain persons*

**6.**—(1) The restrictions imposed by section 52 of the Act shall not apply—

(*a*) to the sale, offer or exposure for sale or supply by any person listed in column 1 of Part I of Schedule 1 to this order, or

(*b*) to the supply by any person listed in column 1 of Part II of the said Schedule,

of the prescription only medicines and the pharmacy medicines specified in column 2 of Part I or Part II of the said Schedule as the case may be in relation to that person if and so long as the conditions specified in the corresponding paragraphs in column 3 of Part I or Part II of the said Schedule as the case may be are satisfied.

(2) The restrictions imposed by section 53 of the Act shall not apply to the sale, offer or exposure for sale or supply by any person listed in column 1 of Part I or Part II of Schedule 1 to this order of any medicinal product on a general sale list (if any) which is specified in column 2 of Part I or Part II of the said Schedule as the case may be in relation to that person if and so long as the conditions specified in the corresponding paragraphs in column 3 of Part I or Part II of the said Schedule as the case may be are satisfied.

(3) For the purposes of the sale, offer or exposure for sale or supply of any veterinary drug on a general sale list, section 53 of the Act shall have effect without the conditions in sub-section (3) of the said section being required to be fulfilled if and so long as the sale or supply is by the holder of a product licence granted under Part II of the Act in respect of such veterinary drugs and is to a person who has in his charge or maintains animals for the purposes of and in the course of carrying on a business, whether as his sole business activity or as a substantial part of his business activities.

*Exemption for medicinal products at high dilutions*

**7.**—(1) The restrictions imposed by section 52 of the Act shall not apply to the sale, offer or exposure for sale or supply of any medicinal product (not being a medicinal product for parenteral administration or a controlled drug) which consists solely of one or more unit preparations of any substance specified in the following sub-paragraphs diluted to the extent specified therein and for the use (if any) specified therein, namely—

(*a*) any substance where the unit preparation has been diluted to at least one part in a million (6x),

(*b*) any substance listed in Part I of Schedule 2 to this order where the unit preparation has been diluted to at least one part in a thousand (3x),

(*c*) any substance listed in column 1 of Table A of Schedule 3 to the Medicines (General Sale List) Order 1977(**a**) or in Part III of Schedule 2 to this order or, if the medicinal product in question is for external use only, any substance listed in column 1 of Table B of Schedule 3 to the Medicines (General Sale List) Order 1977 or in Part IV of Schedule 2 to this order, in each case where the unit preparation has been diluted to at least one part in ten (1x),

if and so long as the person selling or supplying the medicinal product as aforesaid has been requested by or on behalf of a particular person and in that person's presence to use his own judgment as to the treatment required.

(2) The said restrictions also do not apply to the sale, offer or exposure for sale or supply of any medicinal product (not being a medicinal product for parenteral administration or a controlled drug) which consists of one or more unit preparations of any substance specified in the following sub-paragraphs diluted to the extent specified therein and for the use (if any) specified therein, namely—

(*a*) any substance listed in Part II of Schedule 2 to this order where the unit preparation has been diluted to at least one part in a million (6x),

(**a**) S.I. 1977/2129. (1977 III, p. 5921).

(*b*) any substance listed in column 1 of Table A of Schedule 3 to the Medicines (General Sale List) Order 1977 or in Part III of Schedule 2 to this order or, if the medicinal product in question is for external use only, any substance listed in column 1 of Table B of Schedule 3 to the Medicines (General Sale List) Order 1977 or in Part IV of Schedule 2 to this order, in each case where the unit preparation has been diluted to at least one part in ten (1x),

if and so long as the conditions specified in section 53 of the Act are fulfilled.

*Exemption for foods and cosmetics*

**8.** For the purposes of the sale, offer or exposure for sale or supply of any medicinal product on a general sale list which is for sale either for oral administration as a food or for external use as a cosmetic, section 53 of the Act shall have effect without the condition in sub-section (2) of the said section being required to be fulfilled.

*David Ennals,*
Secretary of State for Social Services.

15th December 1977.

*John Morris,*
Secretary of State for Wales.

15th December 1977.

*Bruce Millan,*
Secretary of State for Scotland.

15th December 1977.

In witness whereof the official seal of the Minister of Agriculture, Fisheries and Food is hereunto affixed on 15th December 1977.

(L.S.)

*John Silkin,*
Minister of Agriculture, Fisheries and Food.

Sealed with the official seal of the Department of Health and Social Services for Northern Ireland this 19th day of December 1977.

(L.S.)

*N. Dugdale,*
Permanent Secretary.

Sealed with the official seal of the Department of Agriculture for Northern Ireland this 20th day of December 1977.

(L.S.)

*J. A. Young,*
Permanent Secretary.

## SCHEDULE 1

EXEMPTIONS FOR CERTAIN PERSONS FROM SECTIONS 52 AND 53 OF THE ACT

Article 6(1)(a) and (2) Part I

| Column 1 | Column 2 | Column 3 |
|---|---|---|
| Persons exempted | Medicinal products to which the exemption applies | Conditions |
| 1. State registered chiropodists. | 1. Medicinal products on a general sale list for external use either listed in Schedule 1 to the Medicines (General Sale List) Order 1977(a) or composed solely of substances listed in Schedule 3 to the said order and any of the following pharmacy medicines—<br><br>Paint containing not more than 9.0 per cent Borotannic complex<br><br>Ointment, tincture or dusting powder containing not more than 5.0 per cent Diamthazole hydrochloride<br><br>Ointment or lotion containing not more than 10.0 per cent Buclosamide or not more than 10.0 per cent Crotamiton<br><br>Cream, jelly or powder containing not more than 1.0 per cent Fenticlor<br><br>Pastes containing not more than 70.0 per cent Salicylic acid or not more than 70.0 per cent Pyrogallol<br><br>Powder or cream containing not more than 2.0 per cent 1-Phenoxypropan-2-ol<br><br>Dusting powder or jelly or tincture containing not more than 0.4 per cent Hydrargaphen<br><br>Potassium permanganate crystals or solution. | 1.(1) The sale or supply shall only be in the course of their professional practice.<br><br>(2) The medicinal product must have been made up for sale or supply in a container elsewhere than at the place at which it is sold or supplied. |

(a) S.I. 1977/2129. (1977 III, p. 5921).

| Column 1 | Column 2 | Column 3 |
|---|---|---|
| Persons exempted | Medicinal products to which the exemption applies | Conditions |
| 2. Registered ophthalmic opticians. | 2. Prescription only medicines that are eye drops or eye ointments containing Mafenide propionate or not more than 30.0 per cent of Sulphacetamide sodium or Sulphafurazole diethanolamine equivalent to not more than 4.0 per cent Sulphafurazole and prescription only medicines containing any of the following substances but no other substance listed in column 1 of Part I or Part II of Schedule 1 to the Medicines (Prescription Only) Order 1977(a)—<br><br>Atropine<br>Atropine methobromide<br>Atropine methonitrate<br>Atropine sulphate<br>Bethanecol chloride<br>Carbachol<br>Cyclopentolate hydrochloride<br>Ecothiopate iodide<br>Homatropine hydrobromide<br>Hyoscine hydrobromide<br>Naphazoline nitrate<br>Neostigmine methylsulphate<br>Physostigmine<br>Physostigmine salicylate<br>Physostigmine sulphate<br>Pilocarpine hydrochloride<br>Pilocarpine nitrate<br>Tropicamide. | 2. The sale or supply shall be only—<br><br>(*a*) in the course of their professional practice, and<br><br>(*b*) in an emergency. |
| 3. Holders of manufacturer's licences where the licence in question con- | 3. Medicinal products on a general sale list for external use either listed in | 3. The licence holder shall only sell or supply the medicinal product in |

(a) S.I. 1977/2127. (1977 III, p. 5820).

| Column 1 | Column 2 | Column 3 |
|---|---|---|
| Persons exempted | Medicinal products to which the exemption applies | Conditions |
| tains a provision that the licence holder shall only manufacture the medicinal product to which the licence relates for a particular person after being requested by or on behalf of that person and in that person's presence to use his own judgment as to the treatment required. | Schedule 1 to the Medicines (General Sale List) Order 1977 or composed solely of substances listed in Schedule 3 to the said order and pharmacy medicines which are for external use in the treatment of hair and scalp conditions and which contain any of the following substances—<br>not more than 5.0 per cent of Boric acid<br>Isopropyl myristate or Lauryl sulphate<br>not more than 0.004 per cent Oestrogens<br>not more than 1.0 per cent of Resorcinol<br>not more than 3.0 per cent of Salicylic acid<br>not more than 0.2 per cent of Sodium pyrithione or Zinc pyrithione. | question to a particular person after being requested by or on behalf of that person and in that person's presence to use his own judgment as to the treatment required. |
| 4. Persons selling or supplying medicinal products to universities, other institutions concerned with higher education or institutions concerned with research. | 4. All prescription only medicines, pharmacy medicines and medicinal products on a general sale list. | 4.(1) The sale or supply shall be for the purpose of that education or research.<br>(2) The sale or supply shall be subject to the presentation of an order signed by the principal of the institution concerned with that education or research or the appropriate head of department in charge of the specified course of research stating—<br>(*a*) the name of the institution for which the medicinal product is required,<br>(*b*) the purpose for which |

| Column 1<br>Persons exempted | Column 2<br>Medicinal products to which the exemption applies | Column 3<br>Conditions |
|---|---|---|
| | | the medicinal product is required, and<br>(*c*) the total quantity required. |
| 5. Persons selling or supplying medicinal products to any of the following—<br>(1) a public analyst appointed under section 89 of the Food and Drugs Act 1955(**a**), section 27 of the Food and Drugs (Scotland) Act 1956(**b**) or section 31 of the Food and Drugs Act (Northern Ireland) 1958(**c**),<br>(2) an agricultural analyst appointed under section 67 of the Agriculture Act 1970(**d**),<br>(3) a person duly authorised by an enforcement authority under sections 111 and 112 of the Act,<br>(4) a sampling officer within the meaning of the Food and Drugs Act 1955, the Food and Drugs (Scotland) Act 1956 or the Food and Drugs Act (Northern Ireland) 1958, or<br>(5) a sampling officer within the meaning of Schedule 3 to the Act. | 5. All prescription only medicines, all pharmacy medicines and all medicinal products on a general sale list. | 5. The sale or supply shall be subject to the presentation of an order signed by or on behalf of any person listed in sub-paragraphs (1), (2), (3), (4) or (5) of column 1 of this paragraph stating the status of the person signing it and the amount of the medicinal product required and shall be only in connection with the exercise by those persons of their statutory functions. |
| 6. Persons selling or supplying medicinal products to any person employed or engaged in connection with a scheme for testing the quality and amount of the drugs, preparations and appliances supplied under the National Health Service | 6. All prescription only medicines, all pharmacy medicines and all medicinal products on a general sale list. | 6.(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 stating the status of the person signing it and the amount of the medicinal product required. |

(**a**) 1955 c. 16 (4 & 5 Eliz. 2). (**b**) 1956 c. 30. (**c**) 1958 c. 27 (N.I.). (**d**) 1970 c. 40.

| Column 1 | Column 2 | Column 3 |
|---|---|---|
| Persons exempted | Medicinal products to which the exemption applies | Conditions |
| Act 1977(a), the National Health Service (Scotland) Act 1947(b) and the Health and Personal Social Services (Northern Ireland) Order 1972(c), or under any regulations made under the said Acts or Order. | | (2) The sale or supply shall be for the purposes of the said scheme. |
| 7. Persons providing a poultry vaccination service. | 7. The poultry vaccines listed in paragraph 11 of Schedule 1 to the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1977(d). | 7. The sale or supply shall be only to a person who has charge of animals for the purpose of and in the course of carrying on a business, whether as his sole business activity or as a substantial part of his business activities. |
| 8. Persons selling or supplying medicinal products to the persons referred to in paragraph 7 above. | 8. The prescription only medicines referred to in paragraph 7 above. | 8. The sale or supply shall be subject to the presentation of an order signed by the purchaser stating the amount of the prescription only medicine required. |
| 9. Persons selling or supplying medicinal proproducts to veterinary surgeons and veterinary practitioners. | 9. All prescription only medicines, all pharmacy medicines and all medicinal products on a general sale list. | 9. No conditions. |
| 10. Persons selling or supplying medicinal products to the British Standards Institution. | 10. All prescription only medicines, all pharmacy medicines and all medicinal products on a general sale list. | 10.(1) The sale or supply shall be subject to the presentation of an order signed on behalf of the British Standards Institution stating the status of the person signing it and the amount of the medicinal product required.<br>(2) The sale or supply shall be only for the purpose of testing or determining the standards for containers of medicinal products. |

(a) 1977 c. 49. (b) 1947 c. 27. (c) S.I. 1972/1265 (N.I. 14).
(d) S.I. 1977/2167 (1977 III, p. 6141).

PART II Article 6(1) (*b*) and (2)

| Column 1<br>Persons exempted | Column 2<br>Medicinal products to which the exemption applies | Column 3<br>Conditions |
|---|---|---|
| 1. Royal National Lifeboat Institution and certificated first aiders of the Institution. | 1. All prescription only medicines, all pharmacy medicines and all medicinal products on a general sale list. | 1. The supply shall be only so far as is necessary for the treatment of sick and injured persons. |
| 2. British Red Cross Society and certificated first aid and certificated nursing members of the Society. | 2. All pharmacy medicines and all medicinal products on a general sale list. | 2. The supply shall be only so far as is necessary for the treatment of sick and injured persons. |
| 3. St John Ambulance Association and Brigade and certificated first aid and certificated nursing members of the Association and Brigade. | 3. All pharmacy medicines and all medicinal products on a general sale list. | 3. The supply shall be only so far as is necessary for the treatment of sick and injured persons. |
| 4. St Andrew's Ambulance Association and certificated first aid and certificated nursing members of the Association. | 4. All pharmacy medicines and all medicinal products on a general sale list. | 4. The supply shall be only so far as is necessary for the treatment of sick and injured persons. |
| 5. Order of Malta Ambulance Corps and certificated first aid and certificated nursing members of the Corps. | 5. All pharmacy medicines and all medicinal products on a general sale list. | 5. The supply shall be only so far as is necessary for the treatment of sick and injured persons. |
| 6. Persons authorised by licences granted under regulation 5 of the Misuse of Drugs Regulations 1973 (**a**) or regulation 5 of the Misuse of Drugs (Northern Ireland) Regulations 1974(**b**) to supply a controlled drug. | 6. Such prescription only medicines and such pharmacy medicines as are specified in the licence. | 6. The supply shall be subject to such conditions and in such circumstances and to such an extent as may be specified in the licence. |
| 7. Persons requiring medicinal products for the purpose of enabling them, in the course of any business carried on by them, to comply with any requirements made by or in pursuance of any enactment with respect to the medical treatment of their employees. | 7. Such prescription only medicines and such pharmacy medicines as may be specified in the said enactments and medicinal products on a general sale list. | 7. The supply shall be—<br>(1) for the purpose of enabling them to comply with any requirements made by or in pursuance of any such enactment, and<br>(2) subject to such conditions and in such circumstances as may be specified in the relevant enactments. |

(**a**) S.I. 1973/797 (1973 I, p. 2549). (**b**) S.R. of N.I. 1974/272 (1974 II, p. 1287).

| Column 1 | Column 2 | Column 3 |
|---|---|---|
| Persons exempted | Medicinal products to which the exemption applies | Conditions |
| 8. The owner or the master of a ship which does not carry a doctor on board as part of her complement. | 8. All prescription only medicines, all pharmacy medicines and all medicinal products on a general sale list. | 8. The supply shall be only so far as is necessary for the treatment of persons on the ship. |

Article 7

## SCHEDULE 2

### PART I

*Dilutions of unit preparations diluted to at least one part in a thousand* (3x)

Agaricus muscarius
Ailanthus glandulosa
Apocynum cannabinum
Aurum iodatum
Belladonna
Bismuth Subgallate
Bryonia alba dioica
Calcium Fluoride
Cantharis
Cerium oxalicum
Chelidonium majus
Chenopodium oil
Cina
Colocynthis
Convallaria majalis
Gelsemium sempervirens
Hyoscyamus niger
Lycopodium
Manganese acetate
Ranunculus bulbosus
Terebinthinae oleum

### PART II

*Dilutions of unit preparations diluted to at least one part in a million* (6x)

Adonis vernalis
Agaricus bulbosus
Agaricus muscarius
Agnus castus
Ailanthus glandulosa
Alum
Amethyst
Ammonium Iodide
Amygdalae amarae
Apatite
Apocynum androsaemifolium
Apocynum cannabinum
Argentite
Argentum Chloride
Argentum Iodide
Artemisia cina
Aspidium filix-mas
Aspidium anthelmintica
Aurum Sulphide
Balsamum copaivae
Balsamum peruvianum
Barium Citrate
Barium Sulphate
Bismuth Metal
Bismuth Subgallate
Bismuth Subnitrate
Boletus laricis
Bovista
Cade Oil
Calcium Fluoride
Carduus marianus
Cedar Wood Oil
Cerium Oxalicum
Chalcopyrite
Chalcocite
Chelidonium majus
Chenopodium Oil
Colocynthis
Convallaria majalis
Copper Silicate, Nat.
Crotalus horridus
Cucurbita
Cucumis melo
Datura stramonium
Derris
Diamond
Ephedra vulgaris
Ferric Acetate
Ferrous Iodide
Ferrous Oxalate
Ferrous Sulphide
Formic Acid
Gall
Gelsemium sempervirens
Gneiss
Granatum (Pomegranate Bark)
Hamamelis virginiana
Hepar Sulfuris
Hyoscyamus niger
Iris florentine
Jaborandi
Juniperus sabina
Kaolinite
Lachmanthus tinctoria
Lapis Albus
Lycopodium
Magnesium
Magnesium Acetate
Magnesium Chloride
Magnetite
Manganese Acetate
Nicotiana tabacum
Nicotiana tabacum oil
Oleander
Opuntia vulgaris
Oxalic Acid
Petroleum
Phellandrum aquaticum

Pix Liquida
Platinum
Platinum Chloride
Potassium Hydroxide
Potassium Silicate
Pyrethrum
Pyrolusite
Ranunculus bulbosus
Ranunculus acris
Ranunculus flammula
Ranunculus repens
Ranunculus sceleratus
Rauwolfia serpentina
Rhodium Oxynitrate
Rhododendron chrysanthemum
Rhus toxicodendron
Salicylic Acid
Scrophularia aquatica
Sodium Aluminium Chloride
Sodium Auro-chloride
Sodium Hypochlorite
Sodium Nitrate
Squill
Stannum Metal
Sulphur Iodide
Tannic Acid
Terebinthinae Oleum
Topaz
Uric Acid
Zinc Hypophosphite
Zinc Isovalerate

## Part III

*Dilutions of unit preparations diluted to at least one part in ten* (1x)

Abies excelsa
Abies nigra
Abies nobilis
Acalypha indica
Agate
Alisma plantago Aq.
Alstonia scholaris
Aluminium
Amber (Succinum)
Ambra grisea
Ammonium Phosphate
Angostura vera
Anthoxanthum
Apis mellifera
Aqua Marina
Aqua Mellis
Aralia racemosa
Aranea diadema
Arum maculatum
Arum triphyllum
Asarum
Asperula odorata
Astacus fluviatilis
Auric Chloride
Badiaga
Beech (fagus sylvestris)
Bellis perennis
Berberis aquifolium
Borago officinalis
Butyric Acid
Calcium Metal
Calcium Chloride
Calcium Oxide
Calcium Sulphate
Castoreum
Ceanothus americanus
Cedron
Cerato (Ceratostigma Willmottiana)
Cherry Plum (Prunus cerasifera)
Chestnut, Red and Sweet
Cholesterinum
Chrysolite
Cistus canadensis
Clematis erecta
Conchae vera
Conchiolinum
Corallium Rubrum
Crab Apple
Crocus sativus
Erbium
Erigeron Canadense
Fuligo
Genista tinctoria
Geum urbanum
Glycogen
Gnaphalium leontopodium
Gold
Gorse (Ulex europocus)
Graphites
Gratiola officinalis
Gymnocladus (American Coffee Tree)
Haematoxylon campechianum
Hecla Lava (Ash from Mount Hecla)
Hedeoma pulegioides
Hedera helix
Heliotrope
Heracleum spondylium
Herniaria
Hornbeam (Carpinus betulus)
Iberis amara
Impatiens
Iris germanica
Iris pseudacorus
Jacaranda procera
Jatropha curcas
Juncus communis
Justicia adhatoda
Lamium album

Laurus nobilis oil
Laurocerasus
Ledum palustre
Lilium tigrinum
Lonicera caprifolium
Lysimachia vulgaris
Magnesium Phosphate
Magnesite
Magnolia
Marum verum
Melilotus officinalis
Menispermum canadense
Mephitis putorius
Mercurialis perennis
Mimulus (Mimullis guttatus)
Moschus
Myrica gale
Myrtus communis
Ocimum basilicum
Olive
Oxalis acetosella
Pangamic Acid
Paullinia cupana
Penthorum sedoides
Pollen (mixed)
Polygonatum multiflorum
Polygonum aviculare
Polypodium vulgare
Primula vulgaris
Prunella vulgaris
Ptelea trifoliata
Ratanhia
Robinia pseudoacacia
Rubia tinctorum
Rumex acetosella
Sal Marina
Sarcolactic Acid
Sarracenia purpurea
Scleranthus (Scleranthus annuus)
Silica
Silphium laciniatum
Sodium Benzoate
Spongia marina
Star of Bethlehem (Ornithogalum umbellatum)
Ulmus campestris
Vine
Walnut (juglerus regia)
Water Violet (Hottonia palustris)
Wild Oat
Wild Rose

## PART IV

*Dilutions of unit preparations diluted to at least one part in ten* (1x) *for external use*

Abies excelsa
Abies nigra
Abies nobilis
Acalypha indica
Adonis vernalis
Agaricus bulbosus
Agaricus muscarius
Agate
Agnus castus
Ailanthus glandulosa
Alisma plantago Aq.
Alstonia scholaris
Alum
Aluminium
Amber (Succinum)
Ambra grisea
Amethyst
Ammonium Iodide
Ammonium Phosphate
Amygdalae amarae
Angostura vera
Anthoxanthum
Apatite
Apis mellifera
Apocynum androsaemifolium
Apocynum cannabinum
Aqua Marina
Aqua Mellis
Arallia racemosa
Aranea diadema
Argentite
Argentum Chloride
Argentum Iodide
Artemisia cina
Arum maculatum
Arum triphyllum
Asarum
Asperula odorata
Aspidium filix-mas
Aspidium anthelmintica
Astacus fluviatilis
Auric Chloride
Aurum Sulphide
Balsamum copaivae
Balsamum peruvianum
Badiaga
Barium Citrate
Barium Sulphate
Beech (fagus sylvestris)
Bellis perennis
Berberis aquifolium
Bismuth Metal
Bismuth Subgallate
Bismuth Subnitrate
Boletus laricis
Borago officinalis
Bovista
Butyric Acid

Cade Oil
Calcium Fluoride
Calcium Metal
Calcium Chloride
Calcium Oxide
Calcium Sulphate
Carduus marianus
Castoreum
Ceanothus americanus
Cedar Wood Oil
Cedron
Cerato (Ceratostigma Willmottiana)
Cerium Oxalicum
Chalcopyrite
Chalcocite
Chelidonium majus
Chenopodium Oil
Cherry Plum (Prunus cerasifera)
Chestnut, Red and Sweet
Cholesterinum
Chrysolite
Cistus canadensis
Clematis erecta
Colocynthis
Conchae vera
Conchiolinum
Convallaria majalis
Copper Silicate, Nat
Corallium Rubrum
Crab Apple
Crocus sativus
Crotalus horridus
Cucurbita
Cucumis melo
Datura stramonium
Derris
Diamond
Ephedra vulgaris
Erbium
Erigeron canadense
Ferric Acetate
Ferrous Iodide
Ferrous Oxalate
Ferrous Sulphide
Formic Acid
Fuligo
Gall
Gelsemium sempervirens
Genista tinctoria
Geum urbanum
Glycogen
Gnaphalium leontopodium
Gneiss
Gold
Gorse (Ulex europocus)
Graphites
Gratiola officinalis
Gymnocladus (American Coffee Tree)
Haematoxylon campechianum
Hamamelis virginiana
Hecla Lava (Ash from Mount Hecla)
Hedeoma pulegioides
Hedera helix
Heliotrope
Hepar Sulfuris
Heracleum spondylium
Herniaria
Hornbeam (Carpinus betulus)
Hyoscyamus niger
Iberis amara
Impatiens
Iris florentine
Iris germanica
Iris pseudacorus
Jaborandi
Jacaranda procera
Jatropha curcas
Juncus communis
Juniperus sabina
Justicia adhatoda
Kaolinite
Lachmanthus tinctoria
Lamium album
Lapis Albus
Laurus nobilis oil
Laurocerasus
Ledum palustre
Lilium tigrinum
Lonicera caprifolium
Lycopodium
Lysimachia vulgaris
Magnesium
Magnesium Acetate
Magnesium Chloride
Magnesium Phosphate
Magnesite
Magnetite
Magnolia
Manganese Acetate
Marum verum
Melilotus officinalis
Menispermum canadense
Mephitis putorius
Mercurialis perennis
Mimulus (Mimullis Guttatus)
Moschus
Myrica gale
Myrtus communis
Nicotiana tabacum
Nicotiana tabacum oil
Ocimum basilicum
Oleander
Olive
Opuntia vulgaris
Oxalic Acid
Oxalis acetosella
Pangamic Acid
Paullinia cupana
Penthorum sedoides
Petroleum
Phellandrium aquaticum
Pix Liquida

Platinum
Platinum Chloride
Pollen (mixed)
Polygonatum multiflorum
Polygonum aviculare
Polypodium vulgare
Potassium Hydroxide
Potassium Silicate
Primula vulgaris
Prunella vulgaris
Ptelea trifoliata
Pyrethrum
Pyrolusite
Ranunculus bulbosus
Ranunculus acris
Ranunculus flammula
Ranunculus repens
Ranunculus sceleratus
Ratanhia
Rauwolfia serpentina
Rhodium Oxynitrate
Rhododendron chrysanthemum
Rhus toxicodendron
Robinia pseudoacacia
Rubia tinctorum
Rumex acetosella
Sal Marina
Salicylic Acid
Sarcolactic Acid
Sarracenia purpurea
Scleranthus (Scleranthus annuus)
Scrophularia aquatica
Silica
Silphium laciniatum
Sodium Aluminium Chloride
Sodium Auro-chloride
Sodium Benzoate
Sodium Hypochlorite
Sodium Nitrate
Spongia marina
Squill
Stannum Metal
Star of Bethlehem (Ornithogalum umbellatum)
Sulphur Iodide
Tannic Acid
Terebinthinae Oleum
Topaz
Ulmus campestris
Uric Acid
Vine
Walnut (juglerus regia)
Water Violet (Hottonia Palustris)
Wild Oat
Wild Rose
Zinc Hypophosphite
Zinc Isovalerate

## EXPLANATORY NOTE

*(This Note is not part of the Order.)*

This Order provides exemptions from the restrictions imposed by sections 52 and 53 of the Medicines Act 1968. Section 52 provides that medicinal products not on a general sale list shall only be sold or supplied on premises which are a registered pharmacy and by or under the supervision of a pharmacist. Section 53 provides that medicinal products on a general sale list may only be sold elsewhere than at a registered pharmacy if the conditions specified in that section are fulfilled. The exemptions from section 52 include exemptions for certain categories of persons (listed in column 1 of Schedule 1 to this order), exemptions for certain homoeopathic products and transitional and temporary exemptions. The exemptions from section 53 include exemptions for foods and cosmetics and exemptions for holders of product licences granted in respect of veterinary drugs on a general sale list. The order also specifies the medicinal products which by virtue of section 55(2)(*b*) of the Medicines Act 1968 certified midwives may sell or supply other than in a registered pharmacy.

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