STATUTORY INSTRUMENTS

1984 No. 1861

MEDICINES

The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1984

Made - - - -	*26th November* 1984
Laid before Parliament	*5th December* 1984
Coming into Operation	*1st January* 1985

The Secretary of State concerned with health in England, the Secretaries of State respectively concerned with health and with agriculture in Scotland and in Wales, 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 the powers conferred by sections 57(1), (2), (2A) and 129(4) of the Medicines Act 1968 (**a**) and now vested in them (**b**), and of all the 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 in accordance with section 129(6) of the said Act and with the consent of the Treasury, hereby make the following order:—

Title and commencement

1. This order may be cited as the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1984 and shall come into operation on 1st January 1985.

Interpretation

2.—(1) In this order, unless the context otherwise requires—

"the Act" means the Medicines Act 1968;

"agricultural requisites" means things used in the cultivation of the soil or in the keeping of animals for the production of food or as game and equipment used in or for the collection of produce from animals kept for the production of food and things used for the maintenance of such equipment, and includes any protective clothing but does not include any other kind of human apparel;

"the Department" means the Department of Health and Social Services for Northern Ireland;

"the Department's register" means the register of merchants in veterinary drugs kept by the Department under Article 3(7);

(**a**) 1968 c.67; section 57(2A) was inserted by section 14 of the Animal Health and Welfare Act 1984 (c.40).

(**b**) In the case of the Secretaries of State concerned with health in England and Wales by virtue of S.I. 1969/388, in the case of the Secretary of State concerned with agriculture in Wales by virtue of S.I. 1978/272 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).

"dosage unit" means—

(a) where a veterinary drug is in the form of a tablet or capsule or is an article in some other pharmaceutical form that tablet, capsule or other similar article, and

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

"external use" means application to the skin, hair, fur, feathers, scales, hoof, horn, ear, eye, mouth or mucosa of the throat or prepuce, when local action only is necessary and extensive systemic absorption is unlikely to occur;

"maximum strength" means either the maximum quantity of the substance by weight or volume contained in a dosage unit of a veterinary drug or the maximum percentage of the substance contained in a veterinary drug calculated in terms of weight in weight, weight in volume, volume in weight or volume in volume, as appropriate;

"the Minister" means the Minister of Agriculture, Fisheries and Food;

"qualifying business" means a business in respect of which more than one half of the total sales for its last accounting period was derived from the retail sale of agricultural requisites;

"self-service methods" means any method of sale which allows a purchaser to help himself on or before payment;

"sell by retail" includes offer or expose for sale by retail and supply in circumstances corresponding to retail sale, and cognate expressions shall be construed accordingly;

"the Society" means the Pharmaceutical Society of Great Britain;

"the Society's register" means the register of merchants in veterinary drugs kept by the Society under Article 3(7);

"a specially authorised person" means, in relation to a veterinary drug, either—

(a) a person specially authorised, by virtue of a direction of the licensing authority under Article 3(1) of the Medicines (Exemptions from Licences) (Special and Transitional Cases) Order 1971 **(a)** , to assemble that drug otherwise than in accordance with a manufacturer's licence; or

(b) a person specially authorised by the product licence in respect of that drug to sell the drug under the alternative product name specified in the licence;

"veterinary drug on a general sale list" means a veterinary drug of a description, or falling within a class, specified in an order under section 51 of the Act which is for the time being in force.

(2) A reference in this order to a numbered Article or Schedule is to the Article of, or Schedule to, this order which bears that number.

Exemption for merchants in veterinary drugs

3.—(1) The restrictions imposed by section 52 of the Act (restrictions on sale or supply of medicinal products not on a general sale list) shall not apply to the sale by retail of any veterinary drug not on a general sale list by the

(a) S.I. 1971/1450.

holder of the product licence in respect thereof, by a specially authorised person or by a person who is for the time being carrying on a qualifying business, if—

(a) that veterinary drug either—

(i) is a veterinary drug consisting of or contained in a medicinal product in respect of which there has been granted a product licence, being a licence of right and is not on a general sale list by reason of its consisting of or containing one or more of the substances classified in the first column and specified in the second column of Part A of Schedule 1, or

(ii) is specified in the second column of Part B of Schedule 1, and

(b) the conditions contained in this Article are complied with.

(2) No veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above shall be sold by retail except—

(a) in the container in which it was made up for sale by the manufacturer or, as the case may be, the assembler of the drug;

(b) in a container which has not been opened since the drug was made up for sale in it;

(c) on premises which are occupied by, and under the control of, the seller at the time of sale and which are capable of being closed so as to exclude the public; and

(d) to a person whom the seller knows, or has reasonable cause to believe, to be a person who has in his charge or maintains animals for the purposes of, and in the course of carrying on, a business, either as his sole business activity or as a part of his business activities;

except that, when a person has lawfully purchased a veterinary drug on the premises of the seller, condition *(c)* above shall not apply to the subsequent delivery of that drug to that person.

(3) No veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above shall be sold by retail by self-service methods.

(4) Where in relation to a veterinary drug containing one or more of the substances classified in the first column and listed in the second column of Part A of Schedule 1—

(a) a maximum strength or concentration is specified in the third column of the said Part A, that drug shall not be sold by retail except in containers or packages labelled so as to show a strength or concentration not exceeding that so specified;

(b) a pharmaceutical form is specified in the fourth column of the said Part A, that drug shall not be sold by retail except in the form so specified;

(c) a form of administration is specified in the said fourth column, that drug shall not be sold by retail except for use in the form so specified;

(d) any other restriction is specified in the fifth column of the said Part A, that drug shall not be sold by retail except in compliance with the restriction so specified.

(5)*(a)* In respect of any sale by retail of any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above the seller shall make a record of the sale containing the particulars specified in sub-paragraph *(b)* below and shall keep such record for a period of two years from the date of the sale.

(b) The particulars referred to in sub-paragraph *(a)* above are—

(i) the date on which the veterinary drug was sold;

(ii) the name, quantity and, except where it is apparent from the name, the pharmaceutical form and strength of the veterinary drug sold; and

(iii) the name and address of the person to whom the veterinary drug was sold.

(6) No person shall, in the course of a qualifying business carried on by him, sell by retail any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above unless his name is entered in the Society's register or the Department's register in respect of the premises on which the drug is sold.

(7) The Society and the Department shall keep, for the purposes of paragraph (6) above, a register of persons as being persons entitled, in the course of qualifying businesses carried on by them, to sell by retail on premises in respect of which their names are entered in the register, any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above free from the restrictions imposed by section 52 of the Act, if and so long as the conditions contained in this Article are complied with.

(8) Where a person who, whilst carrying on a qualifying business elsewhere than in Northern Ireland, makes an application in writing to the Society for his name to be entered in the Society's register in respect of any premises on which any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above is to be sold by him in the course of that qualifying business, the Society, shall, subject to paragraphs (12) and (13) below, enter his name in the Society's register in respect of those premises.

(9) Where a person who, whilst carrying on a qualifying business in Northern Ireland, makes an application in writing to the Department for his name to be entered in the Department's register in respect of any premises on which any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above is to be sold by him in the course of that qualifying business, the Department shall, subject to paragraphs (12) and (13) below, enter his name in the Department's register in respect of those premises.

(10) Subject to paragraphs (14) and (15) below a person whose name is entered in the Society's register or the Department's register in respect of any premises shall, in order to retain his name on the register in respect of those premises in any year subsequent to the year in which his name is first entered in it, in the month of January in any such year make an application in writing to the Society or the Department (as the case may be) for his name to be retained in the Society's register or the Department's register (as the case may be) in respect of those premises.

(11) There shall be paid to the Society or the Department in respect of the entry or the retention in the Society's register or the Department's register (as the case may be) of the name of any person in respect of any premises a fee of £55.00.

(12) The Society or the Department shall refuse to enter in the Society's register or the Department's register (as the case may be) the name of any person in respect of any premises unless that person—

(a) has paid to the Society or the Department (as the case may be) the fee specified in paragraph (11) above for the entry of his name in the register; and

(b) has given to the Society or the Department (as the case may be) an undertaking in writing that he will comply with the provisions of the Code of Practice for Merchants Selling or Supplying Veterinary Drugs dated 30th October 1984 and published by the Ministry of Agriculture, Fisheries and Food (being a code relating to the sale or supply of such veterinary drugs as are described in paragraphs (1)*(a)* (i) and (ii) above).

(13) The Society, with the approval of the Minister, or the Department, with the approval of the Department of Agriculture for Northern Ireland, may refuse to enter in the Society's register or the Department's register (as the case may be) the name of any person in respect of any premises if, in the opinion of the Society or the Department (as the case may be), the premises are unsuitable for the storage or safekeeping of any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above.

(14) The Society or the Department shall refuse to retain in the Society's register or the Department's register (as the case may be) the name of any person in respect of any premises unless that person has paid to the Society or the Department (as the case may be) the fee specified in paragraph (11) above for the retention of his name in the register.

(15) The Society, with the approval of the Minister, or the Department, with the approval of the Department of Agriculture for Northern Ireland, may refuse to retain in, or may remove from, the Society's register or the Department's register (as the case may be) the name of any person in respect of any premises if, in the opinion of the Society or the Department (as the case may be),—

(a) that person has failed to observe any of the provisions of the code of practice referred to in paragraph (12)*(b)* above; or

(b) the conditions under which any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above is sold by retail on the premises or under which it is stored on the premises prior to retail sale thereon, are unsuitable for that purpose.

(16) In paragraph (2)*(c)* above "premises" includes a stall of a permanent nature situated at a market or an agricultural showground.

Exemptions in respect of veterinary drugs to be incorporated in animal feeding stuffs

4.—(1) The restrictions imposed by section 52 of the Act shall not apply to the sale by retail of any veterinary drug not on a general sale list by the holder of the product licence in respect thereof, by a specially authorised person or by a person for the time being carrying on a business wholly or mainly comprising either the manufacture of animal feeding stuffs for sale or the sale or supply in bulk of veterinary drugs, if—

(a) that veterinary drug either—

(i) is a veterinary drug consisting of or contained in a medicinal product in respect of which there has been granted a product licence, being a licence of right and is not on a general sale list by reason only of its consisting of or containing one or more substances classified in the first column and specified in the second column of Part A of Schedule 2 or specified in Part A of Schedule 3, or

(ii) is specified in the second column of Part B of Schedule 2 or the second column of Part B of Schedule 3, and

(b) the conditions set out in paragraphs (2) to (6) below are complied with.

(2) No veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above shall be sold by retail except—

(a) for incorporation in animal feeding stuffs; and

(b) to a person whom the seller knows, or has reasonable cause to believe, to be a person carrying on a business wholly or mainly comprising the manufacture of animal feeding stuffs for sale.

(3) No veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above shall be sold by retail by self-service methods.

(4)*(a)* In respect of any sale by retail of any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above the seller shall make a record of the sale containing the particulars specified in sub-paragraph *(b)* below and shall keep such record for a period of two years from the date of the sale.

(b) The particulars referred to in sub-paragraph *(a)* above are—

(i) the date on which the veterinary drug was sold;

(ii) the name, quantity and, except where it is apparent from the name, the pharmaceutical form and strength of the veterinary drug sold; and

(iii) the name and address of the person to whom the veterinary drug was sold.

(5) No person shall, in the course of a business carried on by him, sell by retail any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) above unless—

(a) before making any such sale he, or a previous owner of the business, has notified the Society, or in the case of a business carried on in Northern Ireland, the Department, of the relevant particulars;

(b) every twelve months after the first notification, whether made by him or by a previous owner, he notifies the Society or the Department, as appropriate, of the relevant particulars; and

(c) he notifies the Society or the Department, as appropriate, of any change in the relevant particulars which has occurred since the last notification thereof as soon after such change occurs as is reasonably practicable.

(6) In paragraph (5) above "the relevant particulars", in relation to a business, means the name of the business and the address, or, where appropriate, the location of every premises on which, during the course of the carrying on of that business, veterinary drugs such as are described in paragraph (1)*(a)* (i) or (ii) above are being, or are during the next twelve months to be, sold.

Exemptions for merchants in horse wormers

5.—(1) The restrictions imposed by section 52 of the Act shall not apply during the period referred to in paragraph (15) below to the sale by retail of any veterinary drug not on general sale list by the holder of the product licence in respect thereof, by a specially authorised person or by a person who is for the time being carrying on a qualifying business or a saddlery business if—

(a) that veterinary drug is specified in the second column of Schedule 4, and

(b) the conditions contained in this Article are complied with.

(2) No veterinary drug such as is described in paragraph (1)*(a)* above shall be sold by retail except—

(a) in the container in which it was made up for sale by the manufacturer or, as the case may be, the assembler of the drug;

(b) in a container which has not been opened since the drug was made up for sale in it;

(c) on premises which are occupied by, and under the control of, the seller at the time of sale and which are capable of being closed so as to exclude the public; and

(d) to a person whom the seller knows, or has reasonable cause to believe, to be a person who has in his charge horses or ponies;

except that, where a person has lawfully purchased a veterinary drug on the premises of the seller, condition *(c)* above shall not apply to the subsequent delivery of that drug to that person.

(3) No veterinary drug such as is described in paragraph (1)*(a)* above shall be sold by retail by self-service methods.

(4)*(a)* In respect of any sale by retail of any veterinary drug such as is described in paragraph (1)*(a)* above the seller shall make a record of the sale containing the particulars specified in sub-paragraph *(b)* below and shall keep such record for a period of two years from the date of the sale.

(b) The particulars referred to in sub-paragraph *(a)* above are—

(i) the date on which the veterinary drug was sold; and

(ii) the name, quantity and, except when it is apparent from the name, the pharmaceutical form and strength of the veterinary drug sold.

(5) No person shall, in the course of a qualifying business or a saddlery business carried on by him, sell by retail any veterinary drug such as is described in paragraph (1)*(a)* above unless his name is entered in the register kept by the Society or the Department under paragraph (6) below in respect of the premises on which the drug is sold.

(6) The Society and the Department shall keep, for the purposes of paragraph (5) above, a register of persons as being persons entitled, in the course of qualifying businesses or saddlery businesses carried on by them, to sell by retail on premises in respect of which their names are entered in the register, any veterinary drug such as is described in paragraph (1)*(a)* above free from the restrictions imposed by section 52 of the Act, if and so long as the conditions contained in this Article are complied with.

(7) Where a person who, whilst carrying on a qualifying business or a saddlery business elsewhere than in Northern Ireland, makes an application in writing to the Society for his name to be entered in the register kept by the Society under paragraph (6) above in respect of any premises on which any veterinary drug such as is described in paragraph (1)*(a)* above is to be sold by him in the course of that qualifying business or saddlery business, the Society shall, subject to paragraphs (11) and (12) below, enter his name in that register in respect of those premises.

(8) Where a person who, whilst carrying on a qualifying business or a saddlery business in Northern Ireland, makes an application in writing to the Department for his name to be entered in the register kept by the Department under paragraph (6) above in respect of any premises on which any veterinary drug such as is described in paragraph (1)*(a)* above is to be sold by him in the course of that qualifying business or saddlery business, the Department shall, subject to paragraphs (11) and (12) below, enter his name in that register in respect of those premises.

(9) Subject to paragraphs (13) and (14) below a person whose name is entered in the register kept by the Society or the Department under paragraph (6) above in respect of any premises shall, in order to retain his name on the register in respect of those premises in any year subsequent to the year in which his name was first entered in it, in the month of January in any such year make an application in writing to the Society or the Department (as the case may be) for his name to be retained on that register in respect of those premises.

(10) There shall be paid to the Society or the Department in respect of the entry in, or the retention in, the register kept by the Society or the Department (as the case may be) under paragraph (6) above of the name of any person in respect of any premises, a fee of £25.00, except that no such fee shall be payable in respect of a person whose name is for the time being entered in the Society's register or the Department's register (as the case may be) in respect of those premises as being a person entitled to sell thereon, during the course of a qualifying business carried on by him, any veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) of Article 3.

(11) The Society or the Department shall refuse to enter in the register kept by the Society or the Department (as the case may be) under paragraph (6) above the name of any person in respect of any premises unless that person—

(a) has paid to the Society or the Department (as the case may be) the fee specified in paragraph (10) above for the entry of his name in the register; and

(b) has given to the Society or the Department (as the case may be) an undertaking in writing that he will comply with the provisions of the Code of Practice for Saddlers Selling or Supplying Horse Wormers dated 30th October 1984 and published by the Ministry of Agriculture, Fisheries and Food (being a code of practice relating to the sale or supply of such veterinary drugs as are described in paragraph (1)*(a)* above).

(12) The Society, with the approval of the Minister, or the Department, with the approval of the Department of Agriculture for Northern Ireland, may refuse to enter in the register kept by the Society or the Department (as the case may be) the name of the person in respect of any premises if, in the opinion of the Society or the Department (as the case may be), the premises are unsuitable for the storage or safekeeping of any veterinary drug such as is described in paragraph (1)*(a)* above.

(13) The Society or the Department shall refuse to retain in the register kept by the Society or the Department (as the case may be) under paragraph (6) above the name of any person in respect of any premises unless that person has paid to the Society or the Department (as the case may be) the fee specified in paragraph (11) above for the retention of his name in the register.

(14) The Society, with the approval of the Minister, or the Department, with the approval of the Department of Agriculture for Northern Ireland,

may refuse to retain in, or may remove from, the register kept by the Society or the Department (as the case may be) under paragraph (6) above the name of any person in respect of any premises if, in the opinion of the Society or the Department (as the case may be),—

(a) that person has failed to observe any of the provisions of the code of practice referred to in paragraph (11)*(b)* above; or

(b) the conditions under which any veterinary drug such as is described in paragraph (1)*(a)* above is sold by retail on the premises or under which it is stored on the premises prior to retail sale thereon, are unsuitable for that purpose.

(15) The period referred to in paragraph (1) above is that of 3 years from the date of the coming into operation of this order.

(16) In paragraph (2)*(c)* above "premises" includes a stall of a permanent nature situated at a market or agricultural showground.

(17) For the purposes of this Article—

(a) "saddlery business" means a business in respect of which more than one half of the total sales for its last accounting period was derived from the retail sale of saddlery requisites; and

(b) "saddlery requisites" means equipment used in the keeping of horses or ponies and things used for the maintenance of such equipment and includes any human apparel used in the keeping of horses or ponies.

Exemption for supply, subsequent to sale, by pharmacists

6. The restrictions imposed by section 52 of the Act on the supply of medicinal products shall not apply to the supply in circumstances corresponding to retail sale of a veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) of Article 4 by a pharmacist, or his agent, to the person to whom the pharmacist has, in accordance with the provisions of the said section 52, sold the drug by retail.

Exemption in cases involving another's default

7.—(1) The restrictions imposed by section 52 of the Act shall not apply to the sale by retail, in compliance with the conditions contained in Article 3, of a veterinary drug by a person for the time being carrying on a qualifying business, which drug that person, having exercised all due diligence, on reasonable grounds believes to be a veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) of Article 3, but which, due to the act or default of another person, is not such a veterinary drug.

(2) The restrictions imposed by section 52 of the Act shall not apply to the sale by retail, in compliance with the conditions set out in paragraphs (2) to (6) of Article 4, of a veterinary drug by a person for the time being carrying on a business as is described in Article 4(1), which drug that person, having exercised all due diligence, on reasonable grounds believes to be a veterinary drug such as is described in paragraph (1)*(a)* (i) or (ii) of Article 4 but which, due to the act or default of another person, is not such a veterinary drug.

(3) The restrictions imposed by section 52 of the Act shall not apply to the sale by retail, in compliance with the conditions set out in Article 5, of a veterinary drug by a person for the time being carrying on a qualifying business or a saddlery business (as defined in Article 5(17)), which drug that person, having exercised all due diligence, on reasonable grounds believes to

be a veterinary drug such as is described in paragraph (1)*(a)* of Article 5, but which, due to the act or default of another person is not such a veterinary drug.

Revocation

8. The orders listed in Schedule 5 are hereby revoked.

Norman Fowler,
Secretary of State for
Social Services.

26th November 1984.

George Younger,
Secretary of State for Scotland.

14th November 1984.

Nicholas Edwards,
Secretary of State for Wales.

15th November 1984.

In Witness whereof the Official Seal of the Minister of Agriculture, Fisheries and Food is hereunto affixed on 12th November 1984.

L.S.

Michael Jopling,
Minister of Agriculture,
Fisheries and Food.

Sealed with the Official Seal of the Department of Health and Social Services for Northern Ireland this 15th day of November 1984.

L.S.

Maurice N. Hayes,
Permanent Secretary.

Sealed with the Official Seal of the Department of Agriculture for Northern Ireland this 15th day of November 1984.

L.S.

W. H. Jack,
Permanent Secretary.

We consent,

A. G. Hamilton,
T. Garel Jones,
Two of the Lords Commissioners
of Her Majesty's Treasury.

14th November 1984.

SCHEDULE 1

PART A

LICENCE OF RIGHT VETERINARY DRUGS

Article 3

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
1. GROWTH PROMOTERS	Bacitracin Zinc	6,300,000 i.u./kg	Incorporation in feed	
	Bambermycin	30 g/kg	Incorporation in feed	
	Nitrovin		Incorporation in feed	
	Tylosin Phosphate		Incorporation in feed	For pigs
	Virginiamycin	20 g/kg	Incorporation in feed	
2. COCCIDIOSTATS	Amprolium Hydrochloride			
	Clopidol	33%		
	Decoquinate	80 g/kg	Incorporation in feed	
	Diaveridine			
	Dinitolmide	33%		
	Ethopabate			
	Methyl Benzoquate	1.75%	Incorporation in feed	When combined with not more than 20.6% of Clopidol
	Pyrimethamine		Incorporation in feed	For broiler chickens at levels not exceeding 5 ppm
	Robenidine			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
	Sulphaquinoxaline	12%	Incorporation in feed	When combined with not more than 20% of Amprolium hydrochloride and 1% of Ethopabate with or without 1% of Pyrimethamine
3. ANTI-BLACKHEAD PREPARATIONS	Acinitrazole			
	Aminonitrothiazole			
	Nifursol			
4. SHEEP DIPS AND ECTO-PARASITICIDES	Allethrin			
	Amitraz			
	Benzuldazic Acid, Sodium Salt		External use only	
	Benzyl Benzoate			
	Bromocyclen		External use only	
	Bromophos			
	Bucarpolate			
	Butacarb			
	Carbaryl			
	Carbophenothion			
	Chlorfenvinphos			
	Chlorpyrifos			
	Coal Tar Phenols		External use only	
	Coumaphos			
	Crotoxyphos			
	Cresol	4%		

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
4. SHEEP DIPS AND ECTO-PARASITICIDES (continued)	Cresylic Acid		External use only	
	Derris Resins			
	Diazinon			
	Dichlofenthion			
	Dicophane			
	Dioxathion			
	Dursban			
	Fenchlorphos			
	Fenitrothion			
	Gamma BHC			
	Iodofenphos			
	Lethane			
	Malathion			
	Phosalone			
	Pyractone			
	Pyrimithate			
	Rotenone			
5. ANTHELMINTICS	Bephenium and its salts			
	Bunamidine and its salts			
	Cyacetazide			
	Dichlorvos			
	Diethyl-carbamazine and its salts			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
	Haloxon			
	Levamisole and its salts			
	Mebendazole			
	Metriphonate			
	Morantel and its salts			
	Naphthalophos			
	Niclosamide			
	Parbendazole			
	Phenothiazine			
	Piperazine Carbon Disulphide Complex			
	Pyrantel and its salts			
	Sodium Glycarsamate			
	Tetramisole and its salts			
	Thiabendazole			
	Thiophanate			
6. MILK FEVER PREPARATIONS	Calcium borogluconate Injection whether or not containing all or any of the following substances:			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
6. MILK FEVER PREPARATIONS (continued)	Dextrose, Magnesium and Phosphorus			
7. WARBLE FLY DRESSINGS	Crufomate			
	Famphur			
	Fenchlorphos			
	Fenthion			
	Metriphonate			
	Prolate			
8. LIVER FLUKE REMEDIES	Brotianide			
	Diamphenethide			
	Hexachloroethane			
	Hexachlorophane			
	Nitroxynil and its salts			
	Oxyclozanide			
	Rafoxanide			
	Tribromsalan			
9. SHEEP AND CATTLE CLOSTRIDIAL VACCINES AND ANTISERA	Black Disease Antisera and Vaccines			
	Blackleg (Black-quarter) Vaccines and Antisera			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
	Braxy Vaccines and Antisera			
	Enterotoxaemia Vaccines and Antisera			
	Lamb Dysentry Antisera and Vaccines			
	Pulpy Kidney Vaccines and Antisera			
	Struck Vaccines and Antisera			
	Tetanus Toxoids			
	Combinations of two or more of Braxy, Blackleg (Black-quarter), Lamb Dysentry, Pulpy Kidney, Enterotoxaemia, Struck, Tetanus, Black Disease and Pasteurella Vaccines			
10. POULTRY VACCINES	Avian Encephalomyelitis Vaccine (Living and Inactivated)			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
10. POULTRY VACCINES (continued)	Duck Hepatitis Vaccines (Living) Fowl Pox Vaccines Fowl Typhoid Vaccines (Salmonella gallinarum) Infectious Bronchitis Vaccines (Living and Inactivated) Infectious Bursal Disease Vaccines Infectious Laryngotracheitis Vaccines (Living) Marek's Disease Vaccines Newcastle Disease Vaccines (Living and Inactivated) Combinations of Newcastle Disease Vaccines and Avian Encephalomyelitis Vaccines			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
	Combinations of Newcastle Disease Vaccines with Infectious Bronchitis Vaccines			
	Pasteurella Vaccines			
11. ERYSIPELAS VACCINES	Avian Erysipelas Vaccines			
	Swine Erysipelas Vaccines			
12. SALMONELLA AND E. COLI VACCINES	E. Coli Vaccines (Killed)			
	Salmonella Vaccines (Killed)			
	E. Coli and Salmonella Sero Vaccines			
13. OTHER SHEEP AND CATTLE VACCINES	Foot Rot Vaccines			
	Louping Ill Vaccines (Killed)			
	Ovine Enzootic Abortion Vaccines			
	Pasteurella Vaccines (Killed)			
	Pneumonia Combined Vaccines (Pasteurella)			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
14. MISCELLANEOUS VACCINES	Botulism Vaccines (Mink) Combinations of E. Coli, Salmonella and Pasteurella Vaccines (Killed) Pigeon Pox Vaccines (Living)			
15. SULPHANILAMIDE SURFACE WOUND DRESSINGS	This group comprises powdered surface wound dressings containing not more than 5% of sulphanilamide for application to farm animals			
16. LOCAL ANAESTHETICS	This group comprises injections containing not more than 5% of procaine hydrochloride, lignocaine, or lignocaine hydrochloride with or without not more than 0.002% of adrenaline,			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
16. LOCAL ANAESTHETICS (continued)	adrenaline acid tartrate or noradrenaline			
17. OTHERS	Ammonia Solution Conc.	4%		
	Broxyquinoline			
	Butafosfan			
	Butyl Amino Benzoate		Non-parenteral use only	
	Butynorate			
	Clioquinol	5%		
	Cobalt Carbonate			For use only in anthelmintics
	Cobalt Oxide			For use only in combination with anthelmintics and in ruminal pellets
	Copper its inorganic salts and organic preparations		Non-parenteral use only	
	Creosote		Internal use	
	Dextrose Injection			
	Dill, Oil of			
	Halquinol	5%		
	Iron organic complexes with or without Vitamin B_{12}			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
17. OTHERS (continued)	Magnesium salts		(1) Subcutaneous injection	
			(2) Slow intravenous injection	(2) When combined with any of calcium borogluconate, phosphorus or dextrose
	Menandione Sodium Bisulphite			
	Menandione Dimethyl Pyrimidinol Bisulphite			
	Phenol			
	Poloxalene			
	Turpentine	40%	Internal use	
	Vitamin A			
	Vitamin B group			Excluding:— (1) preparations recommended for administration by intravenous route and (2) those recommended for use for conditions requiring a veterinary diagnosis
	Vitamin D_2			
	Vitamin D_3			
	Vitamin E Injection			

SCHEDULE 1 Article 3

PART B

VETERINARY DRUGS

Product Licence No.	Name of Product*
1. Growth Promoters	
PL 4131/4000	Advantage with Romensin[R]
PL 0095/4026	Avotan 50 Avoparcin / Avotan 50
PL 0095/4036	*Avotan 100*
PL 0095/4028	Avotan 50c Avoparcin
PL 0010/4043	Bayo-n-ox 10% Premix
PL 3832/4031	Eskalin 100
PL 0002/4045 / *PL 3832/4017*	Eskalin 500
PL 0002/4055 / PL 3832/4021	Eskalin S-400
PL 0029/4102	Fedan 10% Premix
PL 0086/4124	Flavomycin 40
PL 0086/4137	*Flavomycin 50*
PL 3405/4016	Nitrovin
PL 3405/4018	*Nitrovin – 20*
PL 2592/4075	Nitrozone 25
PL 4869/4000	Panazone 250–Nitrovin
PL 0095/4007	Payzone 50 MA Nitrovin Milk Replacer Additive
PL 4188/4008	Pentazone 250
PL 0006/4052	Romensin (Monensin Sodium) Premix
PL 2969/4006	Rumevite with Romensin
PL 0012/4170	SPIRA 200
PL 0006/4055	Tylamix Premix 100 g/kg
PL 0006/4062	Tylamix Premix 250 g/kg
PL 3405/4007	Tylosin 100 Premix
PL 3405/4002	ZB 100
PL 3405/4015	*ZB – 100*
PL 3405/4005	ZB 150
PL 3734/4000	Zinc Bacitracin Dumex Feed Grade
PL 3734/4001	Zinc Bacitracin "Dumex" 150 Premix
PL 0109/4001	Zinc Bacitracin Premix
2. Implants	
PL 0829/4119	Ralgro
3. Coccidiostats	
PL 0025/4035	Arpocox
PL 0031/4011	Avatec Premix
PL 3405/4017	Clopidol
PL 0109/4000	Dinormix SR 25
PL 4188/4004	Dinitolmide (DOT) 3.5 – Dinitro-ortho-toluamide / Unicox Pure
PL 4869/4005	*DOT*
PL 0109/4002	DOT (dinitolmide)
PL 1598/4032	DOT Premix 12.5%
PL 1598/4033	DOT Premix 25%

* Alternative product names used by specially authorised persons are not shown.

Items shown in italics did not appear in the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1979 as amended.

Product Licence No. Name of Product

3. Coccidiostats (continued)

PL 0006/4047 { Elancoban / Elancoban Premix
PL 3405/4006 { Elancoban Premix / Monensin 200
PL 0621/4015 Lerbek
PL 0006/4061 Monteban 100 Premix
PL 0025/4019 Nicrazin (Premix)
PL 0086/4135 Sacox 60 Premix
PL 0086/4117 Stenorol
PL 0025/4003 Supacox

4. Anti-Blackhead Preparations

PL 3405/4009 Dazole Premix
PL 4869/4003 Dimetridazole BP (Vet)
PL 3636/4001 Dimetridazole – PML Turkeys
PL 0012/4176 'Emtryl' premix
PL 0012/4174 'Emtryl' Pure
PL 0012/4175 'Emtryl' Soluble

5. Sheep Dips and Ectoparasiticides

PL 0010/4041 Asuntol Scab Dip
PL 1300/4010 Barricade
PL 0676/4089 Battles Improved Organo-Phosphorus Single-Dipping Fluid Dip
PL 0676/4088 Battles Improved Special Single-Dipping Fluid Dip
PL 0676/4086 Battles Liquid Summer Fly Dip (Scab Approved)
PL 0676/4087 Battles Organo-Phosphorus Single-Dipping Fluid Dip
PL 2613/4003 Cheviot Sheep Head Ointment
PL 1300/4004 Ciodrin Insecticide
PL 0805/4015 Cooper MD Powder Dip (BHC)
PL 0003/4124 Cooper – Summer Dip 400
PL 0003/4116 Cooper Winter Dip 200
PL 1300/4011 C Tag 97 Fly Tag/Flectron Fly Tag
PL 4149/4001 Deodorised Malathion Premium Grade
PL 1476/4018 Deosan Dysect
PL 1476/4026 Deosan Flectron
PL 1476/4021 Deosan Summer Dip
PL 1476/4020 Deosan Winter Dip
PL 0829/4127 Dermol
PL 1978/4001 Ectoral Tablets No. 1, 2 and 3
PL 4055/4031 Farmers Fly and Tick Dip
PL 0038/4061 Flockcare Fly and Scab Dip
PL 0038/4063 Flockcare Winter Dip
PL 4436/4005 Fly Dip
PL 2759/4006 Killgerm and Marstan Sheep Dip
PL 2759/4009 Killgerm Scab and Fly Sheep Dip
PL 2759/4007 Killgerm Single Dipping Type Liquid – Sheep Dip
PL 4055/4012 Lice and Mange Remedy
PL 1826/4004 Lice Tick and Mange Dressing (LTM)
PL 0430/4001 Lorexane Medicated Shampoo
PL 2759/4007 Marstan Single Dipping Type Liquid – Sheep Dip
PL 3317/4070 Milocide 50%
PL 0015/4003 Nexion (Bromophos) 2% Dusting Powder
PL 1826/4001 Northern Fly Dip
PL 4055/4003 Northern Fly Dip
PL 1728/4055 Nuvanol Vet
PL 1826/4023 Osmond's Wintol Sheep Dip (1–200)
PL 0025/4044 Ovidip Scab Approved Sheep Dip
PL 0028/4068 Porect
PL 0086/4138 Prodip

Product Licence No.	Name of Product

5. Sheep Dips and Ectoparasiticides (continued)

PL 2100/4034	Rodgers No. 10 Neu-Fly and Tick Dip
PL 0003/4113	Stomoxin
PL 0003/4148	Stomoxin Fly Tags
PL 4055/4001	Supona Fly and Tick Dip
PL 1300/4005	Supona Sheep Dip
PL 4436/4001	Taktic
PL 1345/4040	Taskill
PL 0086/4140	*Tirade Fly Tags*
PL 1728/4070	*Topclip Parasol*
PL 1826/4025	Viper Dip
PL 4055/4000	Viper Winter Dip
PL 4055/4002	Vipex 200 Liquid Dip
PL 4436/4003	Winter Dip
PL 1447/4052	Young's 200 Liquid Tick Dip
PL 1447/4087	Young's 400 Fly Dip
PL 1447/4096	Young's Cypor
PL 1447/4086	Young's Dursban 400 Winter Dip
PL 1447/4080	Young's Ectomort Sheep Dip Young's Scab Approved Summer Dip
PL 1447/4015	Young's Powder Fly Dip
PL 1447/4058	Young's Scab Approved 200 Liquid Tick Dip
PL 1447/4050	Young's Scab Approved 200 Winter Dip
PL 1447/4063	Young's Scab Approved 400 Fly Dip
PL 1447/4068	Young's Scab Approved Bromophos Winter Dip
PL 1447/4085	Young's Scab Approved Diazinon Winter Dip
PL 1447/4071	Young's Scab Approved Dursban 400 Winter Dip
PL 1447/4070	Young's Scab Approved Dursban Winter Dip
PL 1447/4103	*Young's Scab Approved Ectomort Summer Dip*
PL 1447/4073	Young's Scab Approved Iodofenphos Winter Dip
PL 1447/4055	Young's Scab Approved Killtick Liquid Tick Dip
PL 1447/4041	Young's Scab Approved Liquid Fly Dip
PL 1447/4056	Young's Scab Approved Powder Fly Dip
PL 1447/4056	Young's Scab Approved Summer Mycotic Dip
PL 1447/4060	Young's Sheep Blowfly Spray
PL 1447/4083	Young's SP Fly Spray
PL 1447/4015	Young's Summer Mycotic Dip
PL 3893/4069	*Zeprox*

6. Anthelminitics

PL 0829/4135	Action Paranil Pellets
PL 0029/4103	'Amatron' Cattle Drench
PL 0029/4105	'Amatron' Sheep Drench
PL 1447/4092	Anthelpor
PL 4318/4003	Ashmintic Drench
PL 4318/4013	Ashmintic Injection
PL 1732/4059	*Astrobot 5*
PL 1732/4060	*Astrobot 10*
PL 0010/4054	Bayverm L.V. Paste
PL 0010/4058	*Bayverm Pellets 1.9%*
PL 0010/4049	Bayverm Premix 0.6%
PL 0010/4050	Bayverm Premix 2.4%
PL 0010/4047	Bayverm Suspension 2.5%
PL 0010/4048	Bayverm Suspension 10%
PL 3974/4021	*Cevasol Injection*
PL 3974/4020	*Cevasol Worm Drench*
PL 0095/4040	*Cyverm 11.5% Gel*
PL 0095/4038	Cyverm Levamisole 3.2% Drench
PL 0095/4037	Cyverm Levamisole 7.5% Injection
PL 1861/4055	Day's Worm Drench

Product Licence No. Name of Product

6. Anthelminitics (continued)

PL 0100/4047 Dicarocide Forte Injection
PL 0010/4046 Droncit
PL 1596/4071 Duphamisole 7.5% Oral
PL 0025/4023 Equiben
PL 0002/4074, PL 3832/4012 Equitac
PL 0829/4044 Equivurm Plus
PL 0829/4058 Equivurm Plus Paste
PL 0025/4027 Equizole Pony Paste
PL 0025/4042 Eqvalan Paste for Horses
PL 0829/4120 Flubenol
PL 0829/4131 Flubenol Pellets
PL 0010/4055 Flukombin
PL 3763/4000 Gapex
PL 0002/4004 Helmatac In-Feed Wormer
PL 0002/4014, *PL 3832/4051* Helmatac Wormer Pellets
PL 2592/4076 Helminate Sow Wormer Pellets
PL 0025/4041 Ivomec Drench
PL 0025/4043 Ivomec Injection for Pigs
PL 2000/4054 Levacide - C Worm Drench
PL 2000/4049 Levacide Injection
PL 2000/4050 Levacide Worm Drench
PL 1447/4091 Levanthel
PL 3832/4066 Loditac 3% Wormer Pellets
PL 0829/4126 Mebatreat
PL 0829/4113 Mebenvet (1.2%)
PL 0829/4123 Mebenvet (5%)
PL 0010/4026 Neguvon
PL 0012/4003 Nemafax Drench
PL 0012/4150 Nemafax P Wormer Pellets
PL 0012/4149 Nemafax 5
PL 0012/4149 Nemafax 14
PL 0012/4151 Nemafax Cattle, Sheep and Goat Wormer Pellets
PL 0012/4151 Nemafax Sow
PL 0012/4153 Nemafax Wettable Powder
PL 0029/4101 Nemicide Cattle Drench
PL 1345/4069 Nilvax Under 30 kg
PL 0029/4100 Nilverm C. Small Dose Cattle Drench, Nilverm C. Cattle Drench
PL 0029/4101 Nilverm Cattle Special
PL 0029/4114 Nilverm Plus Drench
PL 0029/4118 Nilverm Super
PL 0029/4098 Nilzan C
PL 0029/4115 'Nilzan' Drench Plus
PL 0029/4117 'Nilzan' Drench Super
PL 0829/4114 Ovitelmin
PL 0829/4162 Ovitelmin Bolus
PL 0829/4163 Ovitelmin SC
PL 0086/4121 Panacur 1.5% Pellets
PL 0086/4105 Panacur 2.5% Sheep Wormer
PL 0086/4105 Panacur 2.5% Suspension
PL 0086/4110 Panacur 4% Paste
PL 0086/4106 Panacur 10% Suspension
PL 0086/4107 Panacur 22% Granules
PL 0086/4119 Panacur Paste
PL 0086/4130 Panacur SC, Panacur SC Sheep Wormer
PL 0086/4136 Panacur SC Cattle Wormer

Product Licence No. Name of Product

6. Anthelminitics (continued)

PL 0057/4075 Paratect Sustained Release Bolus
PL 0025/4031 Porcam
PL 0025/4038 Ranizole Paste
PL 0829/4150 Ripercol
PL 0829/4133 Ripercol 3.2% Oral Solution
PL 0829/4140 Ripercol 5% Injectable Solution
PL 0829/4151 Ripercol 7.5% Injectable
PL 0829/4132 Ripercol 15% Injectable Solution
PL 0086/4115 Rumevite Wormablok with Panacur for Cattle
PL 0086/4114 Rumevite Wormablok with Panacur for Sheep
PL 1447/4094 Rycovet Horse and Pony Wormer
PL 0029/4099 Spectril
PL 0057/4060 Strongid-P (Granules)
PL 0057/4062 Strongid-P Paste
PL 0057/4063 Suiminth (Morantel Tartrate)
PL 0286/4032 Synanthic
PL 0286/4034 Synanthic DC
PL 0286/4039 Synanthic Horse Paste
PL 0286/4035 Synanthic Horse Pellets
PL 0003/4127 Systamex Paste 18.5% Cattle and Horse Wormer
PL 0003/4127 Systamex Paste 18.5% Horse and Pony Wormer
PL 0003/4121 Systamex 906 Concentrated Cattle Wormer
PL 0003/4112 Systamex Worm Drench for Cattle and Sheep
PL 1300/4002 Task
PL 0829/4112 Telmin KH
PL 0829/4114 Telmin Liquid
PL 4462/4002 Tetramisole Hydrochloride BP (Vet)
PL 0025/4024 Thibenzole 50% Paste
PL 0025/4024 Thibenzole Paste
PL 1728/4060 Topclip Wormer
PL 1728/4061 Topclip Wormer Pellets
PL 0829/4136 Triban Drench
PL 0002/4061 Valbazen 2.5% Total Spectrum Wormer
PL 3832/4022 Valbazen 2.5% Total Spectrum Wormer
PL 0002/4062 Valbazen 10% Total Spectrum Wormer
PL 3832/4023 Valbazen 10% Total Spectrum Wormer
PL 3832/4015 Valbazen 40% Paste
PL 3832/4025 Valbazen C 10% Total Spectrum Wormer
PL 3832/4026 Valbazen SC 2.5% Total Spectrum Wormer
PL 3832/4016 Valbazen Cattle Wormer Pellets
PL 0012/4172 Vermadex
PL 0086/4139 Wormex
PL 1447/4091 Young's Anthelpor 20
PL 1447/4075 Young's Anthelworm
PL 1447/4075 *Rycovet Widespec*
PL 1447/4090 Young's Anthelworm Feed Pellets
PL 1447/4076 Young's Anthelworm L
PL 1447/4100 Young's Endozal
PL 1447/4079 Young's Nemtrem

7. Milk Fever Preparations

PL 4134/4003 Astracalc (Calcium Borogluconate 20%) No. 1
PL 4134/4004 Astracalc (Calcium Borogluconate 40%) No. 2
PL 4134/4005 Astracalc (Calcium Borogluconate PM) No. 3
PL 4134/4006 Astracalc (Calcium Borogluconate PM 29) No. 4
PL 4134/4007 Astracalc (Calcium Borogluconate PM 40) No. 5
PL 4134/4008 Astracalc (Calcium Borogluconate PMD) No. 6
PL 4134/4009 Astracalc (Calcium Borogluconate M) No. 7
PL 0100/4028 Boracalinate 20 Injection

Product Licence No. Name of Product

7. Milk Fever Preparations (continued)

PL 0100/4027 Boracalinate 40 Injection
PL 0100/4025 Boracalinate MP Injection
PL 0100/4048 Boracalinate MPD Injection
PL 0123/4034 Calcibor C.B.G. 20% w/v
PL 0123/4035 Calcibor C.B.G. 40% w/v
PL 0123/4036 Calcibor C.M.P. 20
PL 0123/4037 Calcibor C.M.P. 30
PL 0123/4038 Calcibor C.M.P. 40
PL 0123/4039 Calcibor C.M.P. and D
PL 2324/4077 Calcium Borogluconate 30% and Magnesium Hypophosphite 2.2% Solution CMP 30
PL 0829/4118 Calcium Borogluconate 40% with Magnesium and Phosphorus
PL 2848/4018 Calcium Borogluconate Injection B Vet C 20%
PL 2848/4019 Calcium Borogluconate Injection B Vet C 30%
PL 2848/4020 Calcium Borogluconate Injection B Vet C 40%
PL 2324/4076 Calcium Borogluconate Solution CBG 20
PL 4134/4003 Flexopax (Calcium Borogluconate 20%) No. 1
PL 4134/4004 Flexopax (Calcium Borogluconate 40%) No. 2
PL 4134/4005 Flexopax (Calcium Borogluconate PM) No. 3
PL 4134/4008 Flexopax (Calcium Borogluconate PMD) No. 6
PL 2324/4079 Glucose Saline Injection
PL 2324/4078 Injection of Calcium Borogluconate 40% and Magnesium Hypophosphite 2.2% Solution CMP 40
PL 4134/4012 Novocalc
PL 1345/4007 TVL Calcium Borogluconate "Borocal"

8. Warble Fly Dressings

PL 0003/4115 Cooper Warble Fly Liquid
PL 0829/4127 Dermol
PL 0010/4045 Neguvon Spot-on
PL 0038/4062 Orbisect Warble Fly Liquid and Louse Liquid for Cattle
PL 0621/4013 Trolene FM
PL 0095/4024 Warbex 16.7% Famphur Pour-on
PL 4436/4000 Warbexol – Ready to Use Systemic Warble Fly Dressing
PL 1447/4059 Young's Concentrated Poron
PL 1447/4074 Young's New Poron
PL 1447/4077 Young's Poron 20

9. Liver Fluke Remedies

PL 0010/4031 Dirian
PL 0025/4036 Flukanide
PL 1826/4000 Hexol
PL 0002/4061, *PL 3832/4022* Valbazen 2.5% Total Spectrum Wormer
PL 0002/4062, *PL 3832/4023* Valbazen 10% Total Spectrum Wormer
PL 3832/4025 Valbazen C 10% Total Spectrum Wormer
PL 3832/4026 Valbazen SC 2.5% Total Spectrum Wormer
PL 1447/4065 Young's Flukol
PL 1447/4101 Young's New Flukol

10. Sheep and Cattle Clostridial Vaccines and Antisera

PL 0086/4132 Heptavac P
PL 1345/4063 Nilvax
PL 1345/4069 Nilvax under 30 kg
PL 0086/4129 Ovivac P
PL 1345/4062 Tasvax 8
PL 1728/4066 Topclip Ewe Vaccine 8 in 1

Product Licence No.	Name of Product
11. Poultry Vaccines	
PL 1531/4001	Addervax ND Vaccine (Living) HB 1
PL 3359/4024	Avian Encephalomyelitis Vaccine Delvax AE
PL 1598/4001	Avian Encephalomyelitis Vaccine (Living) Calnek Strain
PL 1708/4133	Avian Encephalomyelitis Vaccine (Living) Nobilis
PL 3317/4086	Avivac-Avian Encephalomyelitis Vaccine (Live)
PL 0002/4053 *PL 3832/4033*	Bronchimune IB Vaccine *Infectious Bronchitis Vaccine (Living) Strain HL Massachusetts type (Bronchimune)*
PL 0002/4034	Combimune
PL 1598/4029	Combined ND (HB1) and IB (Massachusetts MM) Vaccine (Living)
PL 1598/4028	Combined ND La Sota and IB Vaccine (Living)
PL 3359/4004	Delvax IB H52
PL 3359/4003	Delvax IB H120
PL 3359/4001	Delvax Marek THV Freeze-dried
PL 3359/4035	Delvax ND Hitchner
PL 3359/4006	Delvax ND La Sota
PL 3359/4005	Delvax ND HB1
PL 2592/4055	Eavax
PL 3317/4083	Fowl Laryngotracheitis Vaccine (Modified Live Virus)
PL 1598/4055	Fowl Pox Vaccine Poxine
PL 1598/4053	Fowl Pox Vaccine Poxinet
PL 1708/4139	Gumboro Disease Vaccine (Living) Nobilis
PL 4978/4004	Iblin Emulsion
PL 2592/4037	Ibvax
PL 0002/4002 PL 3832/4056	IB Vaccine (Living) Massachusetts H52 Strain
PL 0002/4003 PL 3832/4036	IB Vaccine (Living) Massachusetts H120 Strain
PL 1708/4135	Inactivated ND Vaccine (oil emulsion) Newcavac Nobilis
PL 1598/4056	Infectious Laryngotracheitis Vaccine (LT-VAC)
PL 2592/4074	Ivamarek Marek's Disease Vaccine
PL 2592/4044	Lentogen HB1
PL 2592/4043	Lentogen La Sota
PL 0086/4004	Marek's Disease Vaccine, Behringwerke
PL 0002/4001 *PL 3832/4039*	Marek's Disease Vaccine (Living) THV (Marimune) *Marek's Disease Vaccine (Living) THV (Strain FC 126) Freeze-dried (Marimune)*
PL 3317/4085	Marek's Disease Vaccine (Live) THV
PL 1598/4026	Marek's Disease Vaccine MD-VAC (Living) THV (Winter Strain) Frozen (Wet)
PL 1598/4027	Marek's Disease Vaccine (Lyophilised) MD-VAC
PL 1708/4141	Marexine MD
PL 1708/4169	Marexine THV/CA
PL 4978/4005	Maternalin Emulsion
PL 4978/4003	Myxilin Emulsion
PL 2592/4033	Newcastle Disease Vaccine (Inactivated) Oil Emulsion (Layer Plus)
PL 0039/4040 PL 4978/4002	Newcadin Day Old
PL 3317/4087	Newcastle Disease Vaccine K2C (Inactivated)
PL 1708/4142	Newcastle Disease Vaccine (Living) Nobilis Clone 30
PL 3317/4086	Newcastle Disease Vaccine (Live) La Sota Strain
PL 1708/4150	Newcavac + EDS'76 Vaccine
PL 1598/4000	NDV (Living) La Sota
PL 0020/4000	ND Vaccine (Living) La Sota
PL 0039/4000 PL 4978/4000	ND Vaccine (Living) HB 1 Strain (Newcadin L)
PL 0039/4029 PL 4978/4001	ND Oil Adjuvant Vaccine (Inactivated) (Newcadin Emulsion)

Product Licence No. Name of Product

11. Poultry Vaccines (continued)

PL 3318/4000 ND Vaccine (Inactivated) Oil Emulsion
PL 1708/4143 Nobi-Vac Egg Drop Syndrome 76 Vaccine BC14 (Inactivated)
PL 1708/4155 Nobi-Vac Gumboro Inactivated
PL 1596/4034 Poulvac AE
PL 1596/4040 Poulvac EDS
PL 1596/4029 Poulvac IB Vaccine H52 (Living)
PL 1596/4030 Poulvac IB Vaccine H120 (Living)
PL 1596/4045 Poulvac Marek HVT Vaccine
PL 1596/4025 Poulvac Marek THV
PL 1596/4026 Poulvac ND Vaccine (Living) HB 1
PL 1596/4027 Poulvac ND Vaccine (Living) La Sota
PL 0002/4005, *PL 3832/4024* Tremimune

12. Erysipelas Vaccines

PL 1531/4012 Ferrovac Ery Vaccine
PL 3317/4110 Swine Erysipelas Vaccine (Inactivated)
PL 1345/4004 Swine Erysipelas Vaccine, Inactivated (Oil Adjuvant) Erysivax

13. Salmonella and E. coli Vaccines

PL 0086/4134 Coliovac
PL 3832/4009 Ecopig
PL 0003/4110 Gletvax K88-Porcine E. coli Vaccine (Polyvalent)
PL 0003/4110 Gletvax-Porcine E. coli Vaccine (Polyvalent)
PL 0003/4110 Gletvax-Porcine E. coli Vaccine (Polyvalent) + K88
PL 0086/4113 Porcovac AT
PL 3832/4004 Scourguard I

14. Other Sheep and Cattle Vaccines

PL 0003/4135 Ovine Enzootic Abortion (Improved) Vaccine
PL 0086/4133 Ovipast

15. Miscellaneous Vaccines

PL 1708/4152 Nobi-Vac L.T. K88
PL 3317/4088 Pigeon Pox Vaccine (Live Virus-Chicken Embryo Origin)

16. Sulphanilamide Surface Wound Dressings

PL 2428/4002 Sulphonamide Wound and Navel Dressing Powder

17. Local Anaesthetics

PL 3317/4049 Lignavet Plus Injection
PL 2324/4074 Lignocaine Anaesthetic Injection
PL 2000/4029 Lignocaine and Adrenalin Injection
PL 2428/4021 Lignocaine Injection
PL 0123/4052 Lignol
PL 1599/4005 Ruby Freezaject
PL 0123/4068 Willcain

18. Others

PL 4318/4002 Ashfer 100
PL 4134/4010 Astracalc (Glucose (Dextrose) 40%) No. 8
PL 4134/4011 Astracalc (Magnesium Sulphate 25%) No. 9
PL 2428/4026 Bactasorb Tablets
PL 0002/4043, *PL 3832/4034* Bloat Guard
PL 0002/4054 Bloat Guard Drench
PL 0002/4051, *PL 3832/4065* Bloat Guard Liquid
PL 3514/4002 Boar Mate

Product Licence No. Name of Product

18. Others (continued)

PL 4261/4000 Bovinyl
PL 1754/4003 Calf Intagen Premix
PL 2613/4000 Cheviot Veterinary Oil
PL 2545/4009 Codifer 10
PL 0010/4009 Coforta 10
PL 0676/4091 Colostrene Watery Mouth Drench for Young Lambs
PL 3317/4010 Copavet
PL 2987/4003 Copper (Cupric) Carbonate
PL 0038/4088 Copporal 2 g
PL 0038/4089 Copporal 4 g
PL 0038/4078 Copprite 2 g
PL 0038/4084 Copprite 4 g
PL 1345/4012 Cujec
PL 2987/4002 Cupric Oxide
PL 2987/4001 Cuprous Chloride
PL 3656/4012 Dio-Iron
PL 1596/4031 Ducrofer
PL 1532/4026 Ferriphor
PL 3317/4041 Ferrofax 10
PL 0113/4005 Fisons Multivitamin Injection
PL 0113/4006 Fisons Vitamin A, D & E Injection
PL 3026/4009 "Flex Flac" Pack for infusion 25% Dextrose Injection BP
PL 0113/4007 Gleptosil
PL 2324/4079 Glucose Saline Injection
PL 0100/4031 Glucose Saline Injection
PL 1708/4121 Haemalift
PL 1754/4009 HI-FAT Baby Calf Food 'Intagen'
PL 0100/4029 Injection Magnesium Sulphate 25%
PL 0100/4022 Injection of Dextrose 40%
PL 1754/4000 Intagen Premix
PL 2000/4017 Intravit 12
PL 0829/4117 Iron Dextran 10% (Pharmacosmos)
PL 0025/4040 Ivomec Injection
PL 0043/4000 Leodex
PL 0043/4042 Leodex 20%
PL 0043/4036 Leodex Plus
PL 2000/4043 Magnesium Sulphate Injection 25% w/v
PL 4127/4000 Micro Anti-Bloat Premix
PL 2592/4059 Microdex
PL 0101/4001 MS 222 Sandoz
PL 3317/4069 Multivet Soluble Powder
PL 0676/4090 Orfoids – Capsules for Orf
PL 0032/4060 Pegasus DE Mineral Mixture
PL 0032/4041 Pegasus Minerals JGW 343
PL 0032/4087 Pegasus OCU Mineral Mixture
PL 1345/4042 Permaco C
PL 1345/4041 Permaco S
PL 1345/4051 Permasel-C
PL 1345/4052 Permasel-S
PL 0295/4000 Poudre Armoricaine
PL 4134/4000 Proviton
PL 4262/4000 Quay-Iron
PL 0829/4133 Ripercol 3.2% Oral
PL 0829/4140 Ripercol 5% Injectable Solution
PL 0829/4132 Ripercol 15% Injectable
PL 2100/4032 Rogers 1–80 Purl Dip
PL 1011/4001 Roscofer 10% Vet
PL 1011/4000 Roscoral Vet
PL 3317/4077 Sildex

Product Licence No. Name of Product

18. Others (continued)

PL 1754/4002 Sow Intagen 0/I
PL 3317/4022 Super Suntax
PL 1599/4004 Swipoul
PL 0032/4039 Telmin Pellet JGW 343
PL 0829/4117 Tendex
PL 5923/4002 Tracerglass C
PL 2686/4000 Vache Ointment
PL 2428/4000 Vetrion 200
PL 3317/4047 Vetrivite Plus
PL 0829/4121 Vital Multivitamin Solution
PL 0038/4060 Vitament Vitamin A, D_3 & E Injection
PL 1532/4020 Vitamin AD_3 E Oral
PL 3317/4069 Vitapol
PL 2969/4001 Vituramag
PL 2969/4005 Vituramol 60 with Romensin
PL 0038/4057 Whitmoyer V – Mix
PL 1447/4036 Young's Swaycop

SCHEDULE 2 Article 4

PART A

LICENCE OF RIGHT VETERINARY DRUGS

Group/Class	Substance
1. GROWTH PROMOTERS	Bacitracin Zinc Bambermycin Nitrovin Tylosin Phosphate Virginiamycin
2. COCCIDIOSTATS	Amprolium Hydrochloride Clopidol Decoquinate Diaveridine Dinitolmide Ethopabate Pyrimethamine Robenidine
3. ANTI-BLACKHEAD PREPARATIONS	Acinitrazole Aminonitrothiazole Nifursol
4. ANTHELMINTICS	Haloxon Mebendazole Parbendazole Phenothiazine Piperazine Carbon Disulphide Complex Tetramisole Thiabendazole
5. OTHERS	Menandione Dimethyl Pyrimidinol Bisulphite Menandione Sodium Bisulphite

Article 4

SCHEDULE 2

PART B

VETERINARY DRUGS

Product Licence No.	Name of Product*
1. Growth Promoters	
PL 0095/4026	Avotan 50 Avoparcin
	Avotan 50
PL 0095/4028	Avotan 50c Avoparcin
PL 0095/4036	*Avotan 100*
PL 0010/4043	Bayo-n-ox 10% Premix
PL 3832/4031	Eskalin 100
PL 0002/4045	Eskalin 500
PL 3832/4017	
PL 0002/4055	Eskalin S-400
PL 3832/4021	
PL 0029/4102	Fedan 10% Premix
PL 0086/4124	Flavomycin 40
PL 0086/4137	Flavomycin 50
PL 3405/4016	Nitrovin
PL 3405/4018	*Nitrovin – 20*
PL 2592/4075	Nitrozone 25
PL 4869/4000	Panazone 250 Nitrovin
PL 0095/4007	Payzone 50 MA Nitrovin Milk Replacer Additive
PL 4188/4008	Pentazone 250
PL 2969/4006	Rumevite with Romensin
PL 0006/4052	Romensin (Monensin Sodium) Premix
PL 0012/4170	SPIRA 200
PL 0006/4055	Tylamix Premix 100 g/kg
PL 0006/4062	Tylamix Premix 250 g/kg
PL 3405/4007	Tylosin 100 Premix
PL 3734/4000	Zinc Bacitracin Dumex Feed Grade
PL 3734/4001	Zinc Bacitracin "Dumex" 150 Premix
PL 0109/4001	Zinc Bacitracin Premix
PL 3405/4015	*ZB-100*
PL 3405/4002	ZB100
PL 3405/4005	ZB150
2. Coccidiostats	
PL 0025/4035	Arpocox
PL 0031/4011	Avatec Premix
PL 3405/4017	Clopidol
PL 0012/4056	"Deccox" Pure
PL 0109/4000	Dinormix SR 25
PL 4188/4004	Dinitolmide (DOT) 3.5-Dinitro-ortho-toluamide
	Unicox Pure
PL 4869/4005	*DOT*
PL 0109/4002	DOT (dinitolmide)
PL 1598/4032	DOT Premix 12.5%
PL 1598/4033	DOT Premix 25%
PL 0006/4047	Elancoban
	Elancoban Premix
PL 3405/4006	Elancoban Premix
	Monensin 200

* Alternative product names used by specially authorised persons are not shown.

Items shown in italics did not appear in the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1979 as amended.

Product Licence No.	Name of Product
2. Coccidiostats (continued)	
PL 0621/4015	Lerbek
PL 0006/4061	Monteban 100 Premix
PL 0025/4019	Nicrazin (Premix)
PL 0025/4010	Pancoxin
PL 0086/4135	Sacox 60 Premix
PL 1598/4036	Salcostat
PL 0086/4117	Stenorol
PL 0025/4003	Supacox
3. Anti-Blackhead Preparations	
PL 4869/4003	*Dimetridazole BP (Vet)*
PL 3636/4001	*Dimetridazole – PML Turkeys*
PL 3405/4009	Dazole Premix
PL 0012/4176	'Emtryl' Premix
PL 0012/4174	'Emtryl' Pure
PL 0012/4175	'Emtryl' Soluble
4. Anthelmintics	
PL 0010/4049	Bayverm Premix 0.6%
PL 0010/4050	Bayverm Premix 2.4%
PL 0829/4131	Flubenol Pellets
PL 0002/4004	Helmatac In-Feed Wormer
PL 0829/4113	Mebenvet (1.2%)
PL 0829/4123	Mebenvet (5%)
PL 0012/4149	Nemafax 5
PL 0012/4149	Nemafax 14
PL 0012/4153	Nemafax Wettable Powder
5. Others	
PL 0002/4043 *PL 3832/4034*	Bloat Guard
PL 1754/4003	Calf Intagen Premix
PL 1754/4009	HI-FAT Baby Calf Food 'Intagen'
PL 1754/4000	Intagen Premix
PL 4127/4000	Micro Anti-Bloat Premix
PL 0032/4041	Pegasus Minerals JGW 343
PL 1754/4002	Sow Intagen O/I
PL 0032/4039	Telmin Pellet JGW 343
PL 2969/4005	Vituramol 60 with Romensin
PL 0038/4057	Whitmoyer V – Mix

Article 4

SCHEDULE 3

PART A

LICENCE OF RIGHT OF VETERINARY DRUGS

Aklomide
Ampicillin Trihydrate
Arsanilic Acid
Benzylpenicillin
Chlortetracycline
Erythromycin
Framycetin Sulphate
Furazolidone
4 hydroxy-3 nitrophenyl arsonic acid
Lincomycin Hydrochloride
Methyl Benzoquate
Nitrofurazone
Oxytetracycline
Procaine Penicillin
Sulphadimidine
Sulphanitran
Sulphaquinoxaline
Tylosin Phosphate

SCHEDULE 3 Article 4

PART B

VETERINARY DRUGS

Product Licence No.	Name of Product*
PL 0006/4053	Apralan Soluble Powder
PL 0006/4057	Apralan 20 Premix
PL 0006/4058	Apralan 100 Premix
PL 4188/4002	Chlortetracycline Feedgrade Auromix 100
PL 0095/4029	Cycostat 66R Robenidine Feed Additive
PL 3405/4010	*Dazole Prescription Premix*
PL 3636/4002	*Dimetridazole – POM Swine and Turkeys*
PL 0034/4031	Dynamutilin 2% Premix
PL 0006/4063	Elancoban for Turkeys and Replacement Chickens
PL 0012/4159	'Emtryl' Prescription Premix
PL 0012/4160	'Emtryl' Prescription Pure
PL 0012/4161	'Emtryl' Prescription Soluble
PL 0012/4158	Emtrymore
PL 1596/4018	Engemycin 5% Soluble Powder
PL 3832/4018	*Eskalin 20 POM for laying and breeding hens*
PL 0002/4071 *PL 3832/4019*	'Eskalin' 500 POM for laying and breeding hens
PL 0057/4068	Fortigro S Premix
PL 1654/4012	Fortracin BMD[R]
PL 3317/4023	Framomycin Feed Additive
PL 3317/4031	Framomycin Soluble Powder 25%
PL 3405/4018	*Furazolidone – 200*
PL 3405/4012	Furazolidone BP
PL 0131/4002	Furazolidone BPC 68
PL 4188/4003	Furazolidone BPC 68 or USNF 13 Unidone
PL 3058/4000	Furazolidone NF BVC
PL 2592/4036	Furazolidone Premix
PL 0006/4050	Granulated Tylosin Concentrate
PL 0032/4084	Lincocin Premix
PL 2592/4065	Micro-Bio Sulphadimidine Premix
PL 3832/4060	*Neftin 200*
PL 0364/4003	Neftin Premix
PL 0364/4004	Neftin Supplement
PL 1598/4037	Nifulidone Premix 11.6%
PL 1598/4037	Nifulidone Premix 22.4%
PL 1598/4037	Nifulidone Premix 44.8%
PL 4188/4001	Oxytetracycline HCI Feedgrade
PL 0034/4001	Quixalud Feed Additive
PL 0034/4026	Quixalud Premix 12%
PL 0025/4028	Ridzol 12% Premix
PL 1728/4041	Sermix
PL 0086/4120	Stenorol for Turkeys
PL 4219/4000	Sulphadimidine
PL 3405/4003	*Sulphadimidine*
PL 3405/4020	*Sulphadimidine – 100*
PL 0057/4061	Terramycine Concentrate 20%
PL 0057/4031	Terramycine 5% Feed Supplement

* Alternative product names used by specially authorised persons are not shown.

Items shown in italics did not appear in the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1979 as amended.

Product Licence No.	Name of Product
PL 0057/4065	Terramycine 20% Feed Supplement
PL 0003/4105	Tribrissen Powder
PL 0006/4045	Tylan Premix 20 g/kg
PL 0006/4001	Tylasul Premix Veterinary Tylasul Premix
PL 0006/4064	Tylasul Premix 100
PL 4188/4000	Unidim
PL 0131/4008	Unidim
PL 4188/4007	Unidim 100
PL 4188/4012	Unizole S Soluble – For Pigs
PL 4188/4011	Unizole S – For Pigs and Poultry
PL 3317/4076	Vi-Mycin Soluble Powder
PL 0038/4037	Whitsyn 10
PL 0038/4047	Whitsyn – S

SCHEDULE 4 Article 5

HORSE WORMERS

Product Licence No.	Name of Product*
PL 1732/4059	*Astrobot 5*
PL 1732/4060	*Astrobot 10*
PL 0010/4054	Bayverm LV Paste
PL 1745/4005	*Equigard 5*
PL 1745/4006	*Equigard 10*
PL 0829/4043	*Equilox*
PL 0002/4074	Equitac
PL 3832/4012	Equitac
PL 0829/4044	Equivurm Plus
PL 0829/4058	Equivurm Plus Paste
PL 0829/4043	Equivurm Syringe
PL 0025/4004	*Equizole Feed Pellets*
PL 0025/4027	Equizole Pony Paste
PL 0025/4005	*Equizole Powder*
PL 0025/4042	Eqvalan Paste for Horses
PL 0844/4055	*Multiwurma (Horses)*
PL 0086/4109	Panacur 22% Granules
PL 0086/4119	Panacur Paste
PL 1599/4001	*Ruby Horse Wormer*
PL 1447/4094	Rycovet Horse and Pony Wormer Paste
PL 0057/4060	Strongid P (Granules)
PL 0057/4062	Strongid P Paste
PL 0286/4039	Synanthic Horse Paste
PL 0286/4035	Synanthic Horse Pellets
PL 0003/4127	Systamex Paste 18.5% Horse and Pony Wormer
PL 0829/4058	*Telmin*
PL 0829/4044	*Telmin Granules*

* Alternative product names used by specially authorised persons are not shown.

Items shown in italics did not appear in the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1979 as amended.

Article 8

SCHEDULE 5

REVOCATION

Orders revoked	References
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1979	S.I. 1979/45
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1979	S.I. 1979/1008
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1980	S.I. 1980/283
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) (No. 2) Order 1980	S.I. 1980/1650
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1981	S.I. 1981/793
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) (No. 2) Order 1981	S.I. 1981/1872
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1982	S.I. 1982/1019
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) (No. 2) Order 1982	S.I. 1982/1805
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1983	S.I. 1983/274
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) (No. 2) Order 1983	S.I. 1983/1156
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1984	S.I. 1984/349

EXPLANATORY NOTE

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

This order re-enacts, with amendments, the provisions of the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1979 and its amending instruments, which are revoked.

The order continues to provide for certain exemptions from the restrictions imposed by section 52 of the Medicines Act 1968. Section 52 restricts the retail sale or supply of medicinal products not on a general sale list (a general sale list being a list of freely sellable medicinal products specified in an order made under section 51 of the Act) to sale or supply from a registered pharmacy by or under the supervision of a pharmacist.

Article 3 of the order exempts from the restrictions imposed by section 52 the sale or supply of any veterinary drug described in Article 3(1)*(a)* of the order by (1) the product licence holder, (2) a specially authorised person (as defined in the order) or (3) a person carrying on a qualifying business (that is to say, a business in respect of which more than one half of the total sales for its last accounting period was derived from the retail sale of agricultural requisites) provided, in each case, that the relevant conditions contained in Article 3 are complied with.

These conditions include a requirement that a person who, in the course of carrying on a qualifying business, wishes to sell any veterinary drug described in Article 3(1)*(a)* must have his name and details of the relevant premises entered in a register of merchants in veterinary drugs ("the Register"). The Register is kept by the Pharmaceutical Society of Great Britain ("the Society") and the Department of Health and Social Services for Northern Ireland ("the Department"). A fee of £55 is to be paid to the Society or the Department for the initial entry in the Register of the name of a person in respect of any premises and a further fee of £55 will be payable annually for the retention of a person's name in the Register in respect of any premises.

A person's name will not be entered in the Register unless he has paid the prescribed fee and has given an undertaking that he will comply with the provisions of the Code of Practice for Merchants Selling or Supplying Veterinary Drugs dated 30th October 1984. This Code is published by the Ministry of Agriculture, Fisheries and Food.

In addition, the Society (with the prior approval of the Minister of Agriculture, Fisheries and Food) or the Department (with the prior approval of the Department of Agriculture for Northern Ireland) may refuse to retain in, or may remove from, its Register, the name of any person in respect of any premises if that person has failed to observe any of the provisions of the Code of Practice referred to above or, if the conditions under which any veterinary drug described in Article 3(1)*(a)* is sold by retail on the premises, or under which it is stored on the premises prior to retail sale, are unsuitable for that purpose.

Article 4 of the order exempts from the restrictions imposed by section 52 the sale or supply of any veterinary drug described in Article 4(1)*(a)* of the order by (1) the product licence holder, (2) a specially authorised person or (3) a person carrying on a business comprising either the manufacture of animal feeding-stuffs for sale or the sale or supply in bulk of veterinary drugs provided, in each case, that the conditions contained in Article 4 are complied with.

Article 5 of the order exempts from the restrictions imposed by section 52 the sale by retail of any veterinary drug (being a horse wormer) described in Article 5(1)*(a)* of the order by (1) the product licence holder, (2) a specially authorised person and (3) a person carrying on a qualifying business or a saddlery business (that is to say, a business in respect of which more than one half of its total sales for its last accounting period was derived from the retail sale of saddlery requisites) provided, in each case, that the conditions contained in Article 5 are complied with.

These conditions are similar to those contained in Article 3 except that horse wormers may only be sold by retail to keepers of horses and ponies. The code of practice to be complied with is the Code of Practice for Saddlers Selling or Supplying Horse Wormers dated 30th October 1984 (published by the Ministry of Agriculture, Fisheries and Food) and the registration and retention fee is £25. (This fee is not payable by a person carrying on a qualifying business on premises in respect of which his name is entered in the Register for the purposes of the exemption contained in Article 3.)

The exemption contained in Article 5 will apply for a period of three years from 1st January 1985.

Article 6 of the order provides for exemptions from the restrictions imposed by section 52 in the case of the supply, subsequent to retail sale, of veterinary drugs described in Article 4(1)*(a)* by pharmacists and Article 7 provides for further exemptions from the restrictions imposed by section 52 in cases involving another person's default.

The changes of substance made by this order are:—

(1) The inclusion of a registration requirement as one of the conditions of the exemption contained in Article 3(Article 3(6)) and of ancilliary provisions relating to that requirement (Article 3(7) to (15)) including conditions in respect of the inclusion and the retention of persons in the registers kept by the Society and the Department for the purposes of that exemption and in respect of the payment of registration and retention fees;

(2) the inclusion, in Article 5, of an exemption for sellers of horse wormers subject to certain conditions being complied with. Such conditions include a registration requirement (Article 5(5)) and conditions in respect of the inclusion and the retention of persons in the registers kept by the Society and the Department for the purposes of the exemption and in respect of registration and retention fees;

(3) the inclusion of a requirement that persons registered in accordance with Article 3(1)*(a)* and Article 5(1)*(a)* have to comply with the provisions of the relevant Codes of Practice (Article 3(12)*(b)* and Article 5(11)*(b)*).

(These Codes of Practice are priced publications and are available from MAFF Publications Unit, Willowburn Estate, Alnwick, Northumberland, NE66 2PF).

SI 1984/1861
ISBN 0-11-047861-4