STATUTORY INSTRUMENTS

1979 No. 45

MEDICINES

The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1979

Made - - - -	16*th January* 1979
Laid before Parliament	19*th January* 1979
Coming into Operation	11*th February* 1979

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) and 129(4) of the Medicines Act 1968**(a)** 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 and commencement

1. This order may be cited as the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1979, and shall come into operation on 11th February 1979.

Interpretation

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

"the Act" means the Medicines Act 1968;

"appointed day" means 1st February 1978, being the day appointed for the purposes of Section 52 of the Act**(c)**;

"dosage unit" means—

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

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(a) 1968 c. 67.

(b) In the case of the Secretaries of State concerned with health in England and Wales by virtue of Article 2(2) of, and Schedule 1 to, the Transfer of Functions (Wales) Order 1969 (S.I. 1969/388), in the case of the Secretary of State concerned with agriculture in Wales by virtue of Article 2(3) of, and Schedule 1 to, the Transfer of Functions (Wales) (No. 1) Order 1978 (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 paragraph 2(1)(*b*) of Schedule 1 to the Northern Ireland Act 1974 (c. 28).

(c) By the Medicines (Pharmacy and General Sale) (Appointed Day) Order 1977 (S.I. 1977/2126).

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

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

"maximum daily dose" means the maximum quantity of the substance contained in the amount of the veterinary drug for internal use which it is recommended should be administered in any period of twenty-four hours;

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

"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;

"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) Any references in this order—

(*a*) to any enactment are, unless the context otherwise requires, references to that enactment as amended or extended by or under any other enactment, and

(*b*) to a numbered article or schedule are references to the article or schedule so numbered in this order.

Exemptions for dealers in veterinary drugs

3.—(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 who is for the time being carrying on a business wholly or mainly comprising the sale by retail of veterinary drugs and other agricultural requisites, provided that—

(*a*) that veterinary drug either—

(i) is not on a general sale list by reason of its being 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

(a) S.I. 1971/1450.

(*b*) the conditions set out in paragraphs (2) to (8) 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*) 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, whether as his sole business activity or as a substantial part of his business activities:

Provided that, where a person has lawfully purchased a veterinary drug on the premises of the seller, condition (*c*) 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*) a maximum daily dose is specified in the fifth 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 daily dose not exceeding the maximum daily dose so specified;

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

(5) 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 stating the particulars specified in paragraph (3) of Regulation 7 of the Medicines (Sale or Supply) (Miscellaneous Provisions) Regulations 1977**(a).**

(6) 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 Pharmaceutical Society of Great Britain or, in the case of a business carried on in Northern Ireland, the Department of Agriculture for Northern Ireland, of the relevant particulars,

(a) S.I. 1977/2132, to which there is an amendment not relevant to this order.

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

(*c*) he notifies the said Society or the said 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.

(7) In paragraph (2)(*c*) above "premises" means—

(*a*) a building or part of a building of a permanent nature, or

(*b*) a stall or other similar structure of a permanent nature situated at a market or agricultural showground.

(8) In paragraph (6) above "the relevant particulars" means the name of the business and the address, or where appropriate, the location, of every premises on or from which the business is being or is during the next twelve months to be carried on, but so far only as such business comprises or includes the retail sale of veterinary drugs such as are described in paragraph (1)(*a*)(i) or (ii) above.

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, provided that—

(*a*) that veterinary drug either—

(i) is not on a general sale list by reason only of its being or containing one or more of the 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) 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 stating the particulars specified in paragraph (3) of Regulation 7 of the Medicines (Sale or Supply) (Miscellaneous Provisions) Regulations 1977.

(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 Pharmaceutical Society of Great Britain, or, in the case of a business carried on in Northern Ireland, the Department of Agriculture for Northern Ireland, of the relevant particulars,

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

(*c*) he notifies the said Society or the said 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" means the name of the business and the address of every premises on or from which the business is being or is during the next twelve months to be carried on, but so far only as such business consists in the retail sale of veterinary drugs such as are described in paragraph (1)(*a*)(i) or (ii) above.

Exemption for supply, subsequent to sale, by pharmacists

5. 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

6.—(1) 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 (8) of Article 3, of a veterinary drug by a person for the time being carrying on a business such as is described in Article 3(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 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 such 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.

Temporary exemptions

7.—(1) The restrictions imposed by section 52 of the Act shall not apply during the period set out in paragraph (4) below to the sale by retail of any veterinary drug not on a general sale list in respect of which either—

(*a*) a product licence has been granted after the appointed day containing a provision (not being such a provision as is described in paragraph (2) or paragraph (3) below) to the effect that the method of sale or supply of the veterinary drug in question may be otherwise than by or under the supervision of a pharmacist, or

(*b*) a product licence has been varied after the appointed day so as to contain a provision (not being such a provision as is described in paragraph (2) or paragraph (3) below) 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 subsection (3) of section 53 of the Act are complied with.

(2) Where a product licence in respect of a veterinary drug has been granted after the appointed day and contains a provision, or has been varied after the appointed day so as to contain a provision, to the effect that the method of sale may be by or under the supervision of a pharmacist, or by the holder of the product licence or by a specially authorised person or by a person carrying on a business such as is described in Article 3(1) (whether or not it also provides for sale by a person carrying on a business such as is described in Article 4(1)), the restrictions imposed by section 52 of the Act shall not apply during the period set out in paragraph (4) below to the sale by retail of that veterinary drug by the holder of the product licence in respect thereof, or by a specially authorised person or by a person who is for the time being carrying on a business such as is described in Article 3(1), if and so long as the conditions set out in paragraphs (2) to (8) of Article 3 are complied with.

(3) Where a product licence in respect of a veterinary drug has been granted after the appointed day and contains a provision, or has been varied after the appointed day so as to contain a provision, to the effect that the method of sale may be by or under the supervision of a pharmacist, or by the holder of the product licence or by a specially authorised person or by a person carrying on a business such as is described in Article 4(1) (whether or not it also provides for sale by a person carrying on a business such as is described in Article 3(1)), the restrictions imposed by section 52 of the Act shall not apply during the period set out in paragraph (4) below to the sale by retail of that veterinary drug by the holder of the product licence in respect thereof, or by a specially authorised person or by a person who is for the time being carrying on a business such as is described in Article 4(1), if and so long as the conditions set out in paragraphs (2) to (6) of Article 4 are complied with.

(4) The period referred to in each of the preceding paragraphs is the period starting with the date of the granting or, as the case may be, variation of the product licence in question and ending one year from the date of publication in the Gazette of the notice stating that the licence has been granted, or, as the case may be, varied.

Transitional exemptions

8. (1) The restrictions imposed by section 52 of the Act shall not apply until 1st February 1980 to any person who sells by retail a veterinary drug not on a general sale list, if and so long as the conditions specified in paragraph (2) of this Article are complied with.

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

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

(*b*) the person selling the veterinary drug in question could lawfully have sold that drug by retail immediately before the appointed day; and

(*c*) the conditions specified in subsection (3) of section 53 of the Act are complied with.

Revocation

9. The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1977**(a)** and the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1978**(b)** are hereby revoked.

David Ennals,
Secretary of State for Social Services.

8th January 1979.

John Morris,
Secretary of State for Wales.

11th January 1979.

Bruce Millan,
Secretary of State for Scotland.

9th January 1979.

In Witness whereof the Official Seal of the Minister of Agriculture, Fisheries and Food is hereunto affixed on 28th December 1978.
(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 16th day of January 1979.
(L.S.)

N. Dugdale,
Permanent Secretary.

Sealed with the Official Seal of the Department of Agriculture for Northern Ireland this 16th day of January 1979.
(L.S.)

J. A. Young,
Permanent Secretary.

(a) S.I. 1977/2167. **(b)** S.I. 1978/1001.

SCHEDULE 1: PART A *Article 3*

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
1. GROWTH PROMOTERS	Avoparcin		Incorporation in feed		
	Bacitracin Zinc	6,300,000 i.u/kg	Incorporation in feed		
	Bambermycin	30g/kg	Incorporation in feed		
	Dimetridazole	225g/kg	Incorporation in feed		For pigs, turkeys and chickens
	Monensin Sodium		Incorporation in feed		For cattle for fattening at levels between 10 and 40 ppm
	Nitrovin		Incorporation in feed		
	Virginiamycin	20g/kg	Incorporation in feed		
2. IMPLANTS	Hexoestrol	15 mg for Poultry or Sheep 45 mg for Cattle			
	Zeranol				
3. COCCIDIOSTATS	Amprolium hydrochloride				
	Clopidol	33%			
	Decoquinate	80g/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

	Monensin Sodium		Incorporation in feed	For chickens for fattening at levels between 100 and 125 ppm
	Nicarbazin Pyrimethamine		Incorporation in feed	For broiler chickens at levels not exceeding 5 ppm
	Robenidine 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
4. ANTI-BLACKHEAD PREPARATIONS	Acinitrazole Aminonitrothiazole Dimetridazole Nifursol	225g/kg	Incorporation in feed	
5. SHEEP DIPS AND ECTO-PARASITICIDES	Allethrin Amitraz Benzyl Benzoate Bromophos Bucarpolate Butacarb Carbaryl Carbophenothion Chlorfenvinphos Chlorpyrifos			
	Coal Tar Phenols		External use only	
	Coumaphos Crotoxyphos Cresol	4%		
	Cresylic Acid		External use only	

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
5. SHEEP DIPS AND ECTO-PARASITICIDES (continued)	Derris Resins Diazinon Dichlofenthion Dicophane Dioxathion Dursban Fenchlorphos Fenitrothion Gamma BHC Iodofenphos Lethane Malathion Phosalone Pyractone Pyrimithate Rotenone				
6. ANTHEL-MINTICS	Bephenium and its salts 4-brom-2, 6 dihydroxy-benzanilide Bunamidine and its salts Cambendazole Cyacetazide Dichlorvos Diethylcarbamazine and its salts Extract Filicis BP Haloxon Levamisole and its salts Mebendazole Methyridine				

	Metriphonate
	Morantel and its salts
	Naphthalophos
	Niclosamide
	Parbendazole
	Phenothiazine
	Piperazine Carbon Disulphide Complex
	Pyrantel and its salts
	Sodium glycarsamate
	Tetramisole and its salts
	Thenium and its salts
	Thiabendazole
	Thiophanate
7. MILK FEVER PREPARATIONS	Calcium borogluconate Injection whether or not containing all or any of the following substances: Dextrose, Magnesium and Phosphorus
8. WARBLE FLY DRESSINGS	Crufomate
	Famphur
	Fenchlorphos
	Fenthion
	Metriphonate
	Prolate
9. LIVER FLUKE REMEDIES	Brotianide
	Carbon Tetrachloride
	Diamphenethide
	Hexachloroethane
	Hexachlorophane
	Nitroxynil and its salts
	Oxyclozanide
	Rafoxanide
	Tribromsalan

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
10. SHEEP AND CATTLE CLOSTRIDIAL VACCINES AND ANTISERA	Black Disease Antisera Black Disease Vaccines Blackleg (Blackquarter) Vaccine and Antiserum Braxy Vaccines and Antisera Enterotoxaemia Vaccines and Antisera Lamb Dysentery Antisera Lamb Dysentery Vaccines Pulpy Kidney Vaccines and Antisera Struck Vaccine and Antiserum Tetanus Antitoxins Tetanus Toxoids Combinations of two or more of Braxy, Blackleg (Blackquarter), Lamb Dysentery, Pulpy Kidney, Enterotoxaemia, Struck, Tetanus, Black Disease and Pasteurella Vaccines				
11. POULTRY VACCINES	Avian Encephalomyelitis Vaccine (Living and Inactivated) Duck Hepatitis Vaccine (Living)				

	Fowl Pox Vaccine
	Fowl Typhoid Vaccines (Salmonella gallinarum)
	Infectious Bronchitis Vaccines (Living)
	Infectious Bronchitis Vaccines (Inactivated)
	Infectious Bursal Disease Vaccines
	Infectious Laryngotracheitis Vaccines (Living)
	Marek's Disease Vaccine
	Newcastle Disease Vaccine (Living)
	Newcastle Disease Vaccine (Inactivated)
	Combinations of Newcastle Disease Vaccines and Avian Encephalomyelitis Vaccines
	Combinations of Newcastle Disease Vaccines with Infectious Bronchitis Vaccines
12. ERYSIPELAS VACCINES	Avian Erysipelas Vaccines
	Swine Erysipelas Vaccines
13. SALMONELLA AND E. COLI VACCINES	E. coli Vaccines (Killed)
	Salmonella Vaccines (Killed)
	E. coli and Salmonella Sero Vaccines
14. OTHER SHEEP AND CATTLE VACCINES	Foot Rot Vaccine
	Louping 111 Vaccines (Killed)

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
14. OTHER SHEEP AND CATTLE VACCINES (continued)	Orf Vaccine (Live) Ovine Enzootic Abortion Vaccines Pasteurella Vaccines (Killed) Pneumonia Combined Vaccine (Pasteurella)				
15. MISCELLANEOUS VACCINES	Botulism Vaccine (Mink) Epidemic Tremors Vaccine Combinations of E. coli, Salmonella and Pasteurella Vaccines (Killed) Pigeon Pox Vaccine (Living)				
16. SULPHANILAMIDE SURFACE WOUND DRESSINGS	This group comprises powdered surface wound dressings containing not more than 5% of sulphanilamide for application to farm animals				

17. LOCAL ANAESTHETICS	This group comprises injections containing not more than 2% of procaine hydrochloride, lignocaine, or lignocaine hydrochloride with or without not more than 0·002% of adrenaline, adrenaline acid tartrate or noradrenaline				
18. OTHERS	Ammonia Solution Conc.	4%			
	Azaperone				For pigs
	Broxyquinoline				
	Butyl amino benzoate		Non-parenteral use only		
	Butynorate		External use only		
	Chlorprothixene				For pigs
	Clioquinol	5%			
	Cobalt Carbonate				For use only in combination with anthelmintics
	Cobalt Oxide				For use only in combination with anthelmintics and in ruminal pellets
	Copper its salts and organic preparations except Copper Sulphate		Non-parenteral use only		
	Copper Sulphate	21%	External use only		
	Creosote		Internal use		
	Dextrose Injection				
	Dill, oil of				
	Dimethyl Sulphoxide				
	Dimetridazole	225g/kg	Incorporation in feed		For swine dysentery
	Etisazole		External use only		

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
18. OTHERS (Continued)	Halquinol	5%			
	Iron Dextran Complex				
	Iron Dextrin				
	Iron Galactan				
	Iron Organic Complexes				
	Magnesium salts				
	Menaphthone Sodium Bisulphite				
	Menaphthone Dimethyl Pyrimidinol Bisulphite				
	Methimazole				
	1-(n-butylamino)-1-methylethyl phosphorus acid				
	Phenol				
	Poloxalene				
	Turpentine	40%	Internal use		
	Vitamin A				
	Vitamin D_2				
	Vitamin D_3				
	Vitamin E Injection				

SCHEDULE 1: PART B *Article 3*

VETERINARY DRUGS

Product Licence No.	Name of Product*
1. Growth Promoters	
PL 0095/4026	Avotan 50 Avoparcin
PL 0006/4052	Romensin (Monensin Sodium) Premix
PL 0095/4007	Payzone 50 MA Nitrovin Milk Replacer Additive
PL 3734/4000	Zinc Bacitracin Dumex Feed Grade
2. Implants	
PL 0829/4119	Ralgro
3. Coccidiostats	
PL 0109/4000	Dinormix SR 25
PL 1598/4032	DOT Premix 12·5%
PL 1598/4033	DOT Premix 25%
PL 0006/4047	Elancoban
PL 0621/4015	Lerbek
PL 0025/4019	Nicrazin (Premix)
PL 0025/4003	Supacox
4. Anti-Blackhead Preparations	
5. Sheep Dips and Ectoparasiticides	
PL 0010/4041	Asuntol Scab Dip
PL 0676/4087	Battle's Organo-Phosphorus Single-Dipping Fluid Dip
PL 0676/4086	Battle's Supona Based Summer Fly Dip (Scab Approved)
PL 0014/4064	Boots Fly Dip
PL 2613/4003	Cheviot Sheep Head Ointment
PL 1300/4004	Ciodrin Insecticide
PL 0805/4015	Cooper MD Powder Dip (BHC)
PL 1978/4000	Ectoral Emulsifiable Concentrate
PL 1978/4001	Ectoral Tablets No. 1, 2 and 3
PL 2513/4036	Insect Powder
PL 2513/4019	Lice and Mange Remedy
PL 1826/4004	Lice Tick and Mange Dressing (LTM)
PL 0430/4001	Lorexane Medicated Shampoo
PL 3317/4070	Milocide 50%
PL 0015/4003	Nexion (Bromophos) 2% Dusting Powder
PL 1826/4001	Northern Fly Dip
PL 2513/4003	Northern Fly Dip
PL 2513/4001	Supona Fly and Tick Dip
PL 1300/4005	Supona Sheep Dip
PL 0014/4065	Taktic
PL 1345/4040	Taskil
PL 1728/4050	Top Clip Dri-Dress
PL 2513/4000	Viper Winter Dip
PL 2513/4002	Vipex 200 Liquid Dip
PL 0014/4066	Winter Dip
PL 1447/4041	Young's Scab Approved Liquid Fly Dip
PL 1447/4052	Young's 200 Liquid Tick Dip
PL 1447/4055	Young's Scab Approved Killtick Liquid Tick Dip

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

Product Licence No.	Name of Product
PL 1447/4058	Young's Scab Approved 200 Liquid Tick Dip
PL 1447/4060	Young's Sheep Blowfly Spray
PL 1447/4063	Young's Scab Approved 400 Fly Dip
PL 1447/4050	Young's Scab Approved 200 Winter Dip
PL 1447/4071	Young's Scab Approved Dursban 400 Winter Dip
PL 1447/4070	Young's Scab Approved Dursban Winter Dip
PL 1447/4068	Young's Scab Approved Bromophos Winter Dip
PL 1447/4073	Young's Scab Approved Iodofenphos Winter Dip
PL 1447/4015	Young's Powder Fly Dip Young's Summer Mycotic Dip
PL 1447/4056	Young's Scab Approved Powder Fly Dip Young's Scab Approved Summer Mycotic Dip
6. Anthelmintics	
PL 1447/4072	Anthelworm L
PL 0025/4032	Bonlam
PL 0025/4029	Cambendazole Paste (Bovicam)
PL 2513/4009	Dicarbazine 40
PL 0025/4022	Equizole A Granules
PL 0025/4027	Equizole Pony Paste
PL 0829/4044	Equivurm Plus
PL 0829/4058	Equivurm Plus Paste
PL 3763/4000	Gapex
PL 0002/4004	Helmatac In-Feed Wormer
PL 0829/4113	Mebenvet (1·2%)
PL 0010/4026	Neguvon
PL 0012/4003	Nemafax
PL 0012/4150	Nemafax P Wormer Pellets
PL 0012/4149	Nemafax Premix
PL 0012/4151	Nemafax Sow Wormer Pellets Nemafax Cattle, Sheep and Goat Wormer Pellets
PL 0012/4153	Nemafax Wormer Drench for Cattle, Sheep and Goats Nemafax Wettable Powder for feed medication of cattle and sheep
PL 0829/4114	Ovitelmin Telmin Liquid
PL 0086/4110	Panacur 4% Powder
PL 0086/4106	Panacur 10% Suspension
PL 0086/4105	Panacur 2·5%
PL 0086/4107	Panacur 22% Granules
PL 0025/4031	Porcam
PL 0057/4060	Strongid-P (Granules)
PL 0057/4062	Strongid-P Paste
PL 0057/4063	Suiminth (Morantel Tartrate)
PL 0286/4032	Synanthic Sheep and Cattle Wormer
PL 0003/4112	Systamex Worm Drench for Cattle and Sheep
PL 1300/4002	Task
PL 0829/4112	Telmin KH
PL 0025/4020	Thibenzole Crumbles
PL 0025/4024	Thibenzole 50% Paste
PL 0025/4021	Thiprazole Granules
PL 0002/4062	Valbazen 10% Drench
PL 0002/4061	Valbazen 2·5% Suspension
PL 0308/4017	Whithelmin Premix
PL 1447/4064	Young's Anthelworm
PL 1447/4075	Young's Anthelworm (New Formula)
PL 1447/4076	Young's Anthelworm L (New Formula)

Product Licence No.	Name of Product
PL 1447/4066	Young's Nemtrem Cattle
PL 1447/4067	Young's Nemtrem Sheep
7. Milk Fever Preparations	
PL 2513/4026	Calcium Borogluconate 20% with Magnesium Phosphorus and Dextrose
PL 2513/4027	Calcium Borogluconate 30% with Magnesium and Phosphorus
PL 2513/4028	Calcium Borogluconate 30% with Magnesium
PL 2513/4029	Calcium Borogluconate 40%
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 1345/4007	TVL Calcium Borogluconate "Borocal"
8. Warble Fly Dressings	
PL 0621/4017	Ruelene Ready to Use (New Formulation)
PL 0621/4014	Trolene 18
PL 0621/4013	Trolene FM
PL 0095/4024	Warbex 16·7% Famphur Pour-On
PL 1447/4059	Young's Concentrated Poron
PL 1447/4074	Young's New Poron
9. Liver Fluke Remedies	
PL 1937/4012	Carbon Tetrachloride Capsules BPC
PL 0010/4031	Dirian
PL 1826/4000	Hexol
PL 2513/4019	Hexol
PL 1826/4002	Osmond's Fluke Drench
PL 0002/4062	Valbazen 10% Drench
PL 0002/4061	Valbazen 2·5% Suspension
PL 1447/4066	Young's Nemtrem Cattle
PL 1447/4067	Young's Nemtrem Sheep
PL 1447/4065	Young's Flukol
10. Sheep and Cattle Clostridial Vaccines and Antisera	
11. Poultry Vaccines	
PL 1531/4001	Addervax ND Vaccine (Living) HB1
PL 1598/4001	Avian Encephalomyelitis Vaccine (Living) Calnek Strain
PL 1708/4133	Avian Encephalomyelitis Vaccine (Living) Nobilis
PL 1531/4005	Avivac—Avian Encephalomyelitis Vaccine (Live)
PL 1531/4009	Avivac—Infectious Bronchitis Vaccine H52 (Live) Massachusetts Strain (IB two)
PL 1531/4010	Avivac—Infectious Bronchitis Vaccine H120 (Live) Massachusetts Strain (IB one)
PL 1531/4008	Avivac—Newcastle Disease Vaccine (Live) Hitchner B1 Strain
PL 0002/4053	Bronchimune IB Vaccine
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)

Product Licence No.	Name of Product
PL 3359/4004	Delvax IB H52
PL 3359/4003	Delvax IB H120
PL 3359/4001	Delvax Marek THV Freeze-dried
PL 3359/4006	Delvax ND La Sota
PL 3359/4005	Delvax ND HB1
PL 2592/4055	Eavax
PL 1531/4003	Fowl Laryngotracheitis Vaccine (Modified Live Virus)
PL 1598/4055	Fowl Pox Vaccine Poxine
PL 1598/4053	Fowl Pox Vaccine Poxinct
PL 1708/4139	Gumboro Disease Vaccine (Living) Nobilis
PL 2592/4037	Ibvax
PL 0002/4003	IB Vaccine (Living) Massachusetts H120 Strain
PL 0002/4002	IB Vaccine (Living) Massachusetts H52 Strain
PL 1708/4135	Inactivated ND Vaccine (oil emulsion) Newcavac Nobilis
PL 1598/4056	Infectious Laryngotracheitis Vaccine (LT-VAC)
PL 2592/4044	Lentogen HB1
PL 2592/4043	Lentogen La Sota
PL 0086/4004	Marek's Disease Vaccine, Behringwerke
PL 0002/4001	Marek's Disease Vaccine (Living) THV (Marimune)
PL 1531/4002	Marek's Disease Vaccine (Live) THV
PL 1598/4021	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 2592/4033	Newcastle Disease Vaccine (Inactivated) Oil Emulsion (Layer Plus)
PL 1531/4007	Newcastle Disease Vaccine K2C (Inactivated)
PL 1598/4000	NDV (Living) La Sota
PL 0002/4000	ND Vaccine (Living) La Sota
PL 1531/4000	ND Vaccine (Live) La Sota
PL 0039/4000	ND Vaccine (Living) HB1 Strain (Newcadin L)
PL 0039/4029	ND Oil Adjuvant Vaccine (Inactivated) (Newcadin Emulsion)
PL 3318/4000	ND Vaccine (Inactivated) Oil Emulsion
PL 1708/4143	Nobi-Vac Egg Drop Syndrome 76 Vaccine BC14 (Inactivated)
PL 1596/4034	Poulvac AE
PL 1596/4029	Poulvac IB Vaccine H52 (Living)
PL 1596/4030	Poulvac IB Vaccine H120 (Living)
PL 1596/4025	Poulvac Marek THV
PL 1596/4026	Poulvac ND Vaccine (Living) HB1
PL 1596/4027	Poulvac ND Vaccine (Living) La Sota
PL 0002/4005	Tremimune
12. Erysipelas Vaccines	
PL 1345/4004	Swine Erysipelas Vaccine, Inactivated (Oil Adjuvant) Erysivax
13. Salmonella and E. coli Vaccines	
PL 0003/4110	Gletvax—Porcine E. coli Vaccine (Polyvalent) Gletvax—Porcine E. coli Vaccine (Polyvalent) +K88 Gletvax K88—Porcine E. coli Vaccine (Polyvalent)
PL 0086/4113	Porcovac AT

Product Licence No.	Name of Product
14. Other Sheep and Cattle Vaccines	
PL 3520/4000	Inactivated Pasteurella Haemalytica A9 Vaccine
PL 1345/4003	TVL Scabivax
15. Miscellaneous Vaccines	
PL 1531/4011	Pigeon Pox Vaccine (Live Virus—Chicken Embryo Origin)
16. Sulphanilamide Surface Wound Dressings	
PL 2513/4023	Naveline Wound Dressing Powder
17. Local Anaesthetics	
PL 2513/4010	Licaine
PL 2324/4074	Lignocaine Anaesthetic Injection
PL 2000/4029	Lignocaine and Adrenalin Injection
PL 1393/4107	Lignocaine Hydrochloride Injection BP 1973
PL 1599/4005	Ruby Freezaject
18. Others	
PL 0002/4043	Bloat Guard
PL 0002/4051	Bloat Guard Liquid
PL 2613/4000	Cheviot Veterinary Oil
PL 2545/4009	Codifer 10
PL 2545/4014	Codifer 10
PL 0010/4009	Coforta 10
PL 3317/4010	Copavet
PL 2987/4003	Copper (Cupric) Carbonate
PL 1345/4012	Cujec
PL 2987/4002	Cupric Oxide
PL 2987/4001	Cuprous Chloride
PL 1596/4031	Ducrofer
PL 1532/4026	Ferriphor
PL 3317/4041	Ferrofax 10
PL 3026/4009	"Flex Flac" pack for infusion 25% Dextrose Injection BP
PL 0113/4007	Gleptosil
PL 1708/4121	Haemalift
PL 2513/4034	Injection of Vitamin A, D_2 and E
PL 0829/4117	Iron Dextran 10% (Pharmacosmos) Tendex
PL 3058/4002	Iron Dextran Injection 100 mg/ml BVC (Ronidex)
PL 0043/4000	Leodex
PL 0010/4006	Netrosylla
PL 1345/4042	Permaco C
PL 1345/4041	Permaco S
PL 0034/4020	Rehydran
PL 1011/4001	Roscofer 10% Vet
PL 1011/4000	Roscoral Vet
PL 2513/4037	Saltona
PL 1599/4004	Swipoul
PL 2868/4000	Vache Ointment
PL 0829/4121	Vital Multivitamin Solution
PL 1532/4020	Vitamin AD_3E Oral
PL 3317/4069	Vitapol
PL 2969/4001	Vituramol Ura-Mag
PL 1447/4036	Young's Swaycop

Article 4 SCHEDULE 2: PART A

Group/Class	Substance
1. GROWTH PROMOTERS	Avoparcin Bacitracin Zinc Bambermycin Dimetridazole Monensin Sodium Nitrovin Virginiamycin
2. COCCIDIOSTATS	Amprolium hydrochloride Clopidol Decoquinate Diaveridine Dinitolmide Ethopabate Monensin Sodium Nicarbazin Pyrimethamine Robenidine
3. ANTI-BLACKHEAD PREPARATIONS	Acinitrazole Aminonitrothiazole Dimetridazole Nifursol
4. ANTHELMINTICS	Haloxon Mebendazole Parbendazole Phenothiazine Piperazine Carbon Disulphide Complex Tetramisole Thiabendazole
5. OTHERS	Dimetridazole Menaphthone Dimethyl Pyrimidinol Bisulphite Menaphthone 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
PL 0002/4055	Eskalin S 400
PL 0002/4045	Eskalin 500
PL 0006/4052	Romensin (Monensin Sodium) Premix
PL 0095/4007	Payzone 50 MA Nitrovin Milk Replacer Additive
PL 3734/4000	Zinc Bacitracin Dumex Feed Grade

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

SCHEDULE 2: PART B (continued) *Article 4*

Product Licence No.	Name of Product
2. Coccidiostats	
PL 0012/4156	"Deccox" Pure
PL 0109/4000	Dinormix SR 25
PL 1598/4032	DOT Premix 12·5%
PL 1598/4033	DOT Premix 25%
PL 0006/4047	Elancoban
PL 0621/4015	Lerbek
PL 0025/4019	Nicrazin (Premix)
PL 0025/4010	Pancoxin
PL 1598/4036	Salcostat
PL 0025/4003	Supacox
3. Anti-Blackhead Preparations	
4. Anthelmintics	
PL 0025/4022	Equizole 'A' Granules
PL 0002/4004	Helmatac In-Feed Wormer
PL 0829/4113	Mebenvet (1·2%)
PL 0025/4020	Thibenzole Crumbles
PL 0025/4021	Thiprazole Granules
PL 0308/4017	Whithelmin Premix
5. Others	

SCHEDULE 3: PART A *Article 4*

Aklomide
Ampicillin Trihydrate
Arsanilic Acid
Benzylpenicillin
Chlortetracycline
E. coli oral vaccine (Inactivated)
Erythromycin
Framycetin Sulphate
Furazolidone
4 hydroxy-3 nitrophenyl arsonic acid
Lincomycin Hydrochloride
Methyl Benzoquate
Nitrofurazone
Oxytetracycline
Procaine Penicillin
Stilboestrol/Methyltestosterone Mixture
Sulphadimidine
Sulphanitran
Sulphaquinoxaline
Tylosin

Article 4

SCHEDULE 3: PART B

VETERINARY DRUGS

Product Licence No.	Name of Product*
PL 0039/4020	E. coli Polysaccharide Antigens for Pigs
PL 1596/4018	Engemycin 5% Soluble Powder
PL 0057/4068	Fortigro S Premix
PL 1654/4012	Fortracin 100
PL 3317/4031	Framomycin Soluble Powder 25%
PL 3317/4023	Framomycin Feed Additive
PL 0131/4002	Furazolidone BPC 68
PL 3058/4000	Furazolidone NF BVC
PL 2592/4036	Furazolidone Premix
PL 0006/4050	Granulated Tylosin Concentrate
PL 1754/4000	Intagen
PL 0032/4084	Lincocin Premix
PL 0364/4003	Neftin Premix
PL 0364/4004	Neftin Supplement
PL 1598/4037	Nifulidone Premix 11·6% Nifulidone Premix 22·4% Nifulidone Premix 44·8%
PL 0034/4001	Quixalud Feed Additive
PL 0057/4061	Terramycin Concentrate 20%
PL 0057/4031	Terramycin 5% Feed Supplement
PL 0057/4065	Terramycin 20% Feed Supplement
PL 1728/4041	Sermix
PL 0003/4105	Tribrissen Powder
PL 0006/4001	Tylasul Premix Veterinary
PL 3317/4076	Vi-Mycin Soluble Powder
PL 0308/4001	Whitsyn 10

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

EXPLANATORY NOTE

(This Note is not part of the Order.)

The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1978 made some amendments to the articles of the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1977, and substituted new schedules. This Order consolidates the two Orders just mentioned, at the same time again replacing the schedules with up-dated schedules.

This Order provides, as did its predecessor, for certain exemptions from the restrictions imposed by section 52 of the Medicines Act 1968.

Section 52 of the Medicines Act 1968 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. The Order exempts from those restrictions the sale or supply of certain veterinary drugs by certain persons provided that the relevant conditions are complied with.

The Order exempts the sale or supply of any veterinary drug described in article 3 of, and referred to in Schedule 1 to, the Order by the holder of the relevant product licence, by a specially authorised person (as defined in article 2) or by a person carrying on a business comprising the sale or supply by retail of veterinary drugs and agricultural requisites, provided that the conditions set out in that article are fulfilled.

It also exempts the sale or supply of any drug described in article 4 of, and referred to in Schedule 2 or Schedule 3 to, the Order by the product licence holder, by a specially authorised person or by 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 that the conditions set out in that article are fulfilled.

Further, the Order provides for exemption for supply, subsequent to sale, by pharmacists and for exemption in cases involving another's default, and makes certain temporary and transitional exemptions.

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