STATUTORY INSTRUMENTS

1977 No. 2167

MEDICINES

The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1977

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

The Secretaries of State respectively concerned with health in England and Wales, the Secretary of State concerned with health and with agriculture in Scotland, the Minister of Agriculture, Fisheries and Food, the Department of Health and Social Services for Northern Ireland, and the Department of Agriculture for Northern Ireland, acting jointly, in exercise of powers conferred by section 57(1) and (2) 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 1977, and shall come into operation on 1st February 1978.

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

(*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;

(a) 1968 c. 67.

(b) In the case of the Secretaries of State concerned with health in England and Wales by virtue of Articles 2(2) of, and Schedule 1 to, the Transfer of Functions (Wales) Order 1969 (S.I. 1969/388 (1969 I, p. 1070)) and in the case of the Northern Ireland Departments by virtue of section 40 of, and Schedule 5 to, the Northern Ireland Constitution Act 1973 (c. 36) and 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(1977 III, p. 5818)).

"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;

"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.

(3) The Interpretation Act 1889**(a)** shall apply for the interpretation of this order as it applies for the interpretation of an Act of Parliament.

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 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 first column of Part B of Schedule 1, and

(*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;

(a) 1889 c. 63.

(*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.

(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) No veterinary drug containing one or more of the substances classified in the first column and specified in the second column of Part A of Schedule 1 shall be sold by retail except—

(*a*) in the pharmaceutical form or for use by the form of administration specified in relation to that drug in the fourth column of the said Part A,

(*b*) in containers or packages labelled with the maximum strength or concentration specified in relation to that drug in the third column, and with the maximum daily dose specified in relation to that drug in the fifth column, of the said Part A, and

(*c*) in compliance with such other restrictions (if any) as are specified in relation to that drug in the sixth column of the said Part A.

(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,

(*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.

(a) S.I. 1977/2132 (1977 III, p. 6030).

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 or by a person for the time being carrying on a business wholly or mainly comprising 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 first column of Part B of Schedule 2 or the first 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 in cases involving another's default

5.—(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

6.—(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 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 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 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 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 on which the product licence is granted or varied and ending one year from the date on which a notice either that the licence has been granted, or, as the case may be, varied is published in the Gazette.

Transitional exemptions

7.—(1) The restrictions imposed by section 52 of the Act shall not apply for a period of two years from the appointed day 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.

David Ennals,
Secretary of State for Social Services.

15th December 1977.

John Morris,
Secretary of State for Wales.

15th December 1977.

Bruce Millan,
Secretary of State for Scotland.

15th December 1977.

In Witness whereof the Official Seal of the Minister of Agriculture, Fisheries and Food is hereunto affixed on 14th December 1977.

(L.S.)

John Silkin,
Minister of Agriculture, Fisheries and Food.

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

(L.S.)

N. Dugdale,
Permanent Secretary.

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

(L.S.)

J. A. Young,
Permanent Secretary.

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
1. GROWTH PROMOTERS	Avoparcin	40g/kg	Incorporation in feed		
	Bacitracin Methylene disalicylate	18,500,000 i.u/kg	Incorporation in feed		
	Bacitracin Zinc	4,600,000 i.u/kg	Incorporation in feed		
	Bambermycin	5g/kg	Incorporation in feed		
	Dimetridazole	225g/kg	Incorporation in feed		For pigs, turkeys and chickens
	Monensin Sodium		Incorporation in feed		
	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				
	Monensin Sodium		Incorporation in feed		
	Nicarbazin				
	Pyrimethamine				
	Robenidine				
	Sulphaquinoxaline	110g/kg 225g/kg for Poultry	Incorporation in feed		

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
4. ANTI-BLACKHEAD PREPARATIONS	Acinitrazole Aminonitrothiazole Dimetridazole Nifursol	225g/kg	Incorporation in feed		
5. SHEEP DIPS AND ECTOPARASITICIDES	Allethrin Amitraz Benzyl Benzoate Bromophos Bucarpolate Butacarb Carbaryl Carbophenothion Chlorfenvinphos Chlorpyrifos Coal Tar Phenols Coumaphos Crotoxyphos Cresol Cresylic Acid Derris Resins Diazinon Dichlofenthion Dicophane Dioxathion Dursban Fenchlorphos Fenitrothion Gamma BHC Iodofenphos Lethane Malathion Phosalone Pyractone Pyrimethate Rotenone	4%	External use only External use only		

6. ANTHELMINTICS	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 Niciosamide 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		With calcium borogluconate as the only active ingredient
8. WARBLE FLY DRESSINGS	Crufomate Famphur Fenchlorphos Fenthion Metriphonate Prolate			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
9. LIVER FLUKE REMEDIES	Brotianide Carbon Tetrachloride Diamphenethide Hexachloroethane Hexachlorophane Nitroxynil and its salts Oxyclozanide Rafoxanide Tribromsalan				
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 Vaccines (Living) Newcastle Disease Vaccines (Inactivated) Combinations of Newcastle Disease Vaccines and Avian Encephalomyelitis Vaccines Combinations of Newcastle Disease Vaccines with Infectious Bronchitis Vaccines
12. ERYSIPELAS VACCINES AND ANTISERA	Avian Erysipelas Vaccines Swine Erysipelas Vaccines and Antisera
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 I 11 Vaccines (Killed) Orf Vaccine (Live attenuated) Ovine Enzootic Abortion Vaccines Pasteurella Vaccines (Killed) Pneumonia Combined Vaccine (Pasteurella)

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Maximum daily dose	Other restrictions
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 for application to farm animals containing not more than 5% of sulphanilamide				
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		Injection		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 and its salts except Copper Sulphate				
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		
Halquinol	5%			
Iron Dextran Complex				
Iron Dextrin				
Iron Galactan				
Iron Injection				
Iron Organic Complexes				
Magnesium				
Menaphthone Sodium Bisulphite				
Methimazole				
1-(n-butylamino)-1-methylethyl phosphorus acid				
Phenol				
Poloxalene				
Triclosan		External use only, as teat dip		
Turpentine	40%	Internal use		
Vitamin A		Non-parenteral use only		
Vitamin D_2		Non-parenteral use only		
Vitamin D_3		Non-parenteral use only		

Article 3

SCHEDULE 1: PART B

VETERINARY DRUGS

Name of Product	*Product Licence No.*
1. Growth Promoters	
Avotan 20 Avoparcin	PL 0095/4026
Fortracin-100	PL 1654/4012
Romensin (Monensin Sodium) Premix	PL 0006/4052
Payzone 50 MA Nitrovin Milk Replacer Additive	PL 0095/4007
2. Implants	
Ralgro	PL 0829/4119
3. Coccidiostats	
DOT Premix 12.5%	PL 1598/4032
DOT Premix 25%	PL 1598/4033
Elancoban	PL 0006/4047
Lerbek	PL 0621/4015
Nicrazin (Premix)	PL 0025/4019
Supacox	PL 0025/4003
4. Anti-Blackhead Preparations	
Dimetridazole	PL 0131/4006
5. Sheep Dips and Ectoparasiticides	
Asuntol Scab Dip	PL 0010/4041
Battle's Organo-Phosphorus Single Dipping Fluid Dip	PL 0676/4087
Battle's Supona Based Summer Fly Dip (Scab Approved)	PL 0676/4086
Boots Fly Dip	PL 0014/4064
Cheviot Sheep Head Ointment	PL 2613/4003
Ciodrin Insecticide	PL 1300/4004
Flymort 24	
Cooper MB Powder Dip (BHC)	PL 0805/4015
Ectoral Emulsifiable Concentrate	PL 1978/4000
Ectoral Tablets No. 1, 2 and 3	PL 1978/4001
Lice Tick and Mange Dressing (LTM)	PL 1826/4004
Lorexane Medicated Shampoo	PL 0430/4001
Northern Fly Dip	PL 1826/4001
Northern Fly Dip	PL 2513/4003
Supona Fly and Tick Dip	PL 2513/4001
Supona Sheep Dip	PL 1300/4005
Taskil	PL 1345/4040
Top Clip Dri-Dress	PL 1728/4050
Viper Winter Dip	PL 2513/4000
Vipex 200 Liquid Dip	PL 2513/4002
Winter Dip	PL 0014/4066
Young's Scab Approved Liquid Fly Dip	PL 1447/4041
Young's 200 Liquid Tick Dip	PL 1447/4052
Young's Scab Approved Killtick Liquid Tick Dip	PL 1447/4055
Young's Scab Approved 200 Liquid Tick Dip	PL 1447/4058
Young's Sheep Blowfly Spray	PL 1447/4060
Young's Scab Approved 400 Fly Dip	PL 1447/4063
Young's Scab Approved 200 Winter Dip	PL 1447/4050
Young's Scab Approved Dursban 400 Winter Dip	PL 1447/4071
Young's Scab Approved Dursban Winter Dip	PL 1447/4070
Young's Scab Approved Bromophos Winter Dip	PL 1447/4068
Young's Scab Approved Iodofenphos Winter Dip	PL 1447/4073
Young's Powder Fly Dip	PL 1447/4015
Young's Summer Mycotic Dip	
Young's Scab Approved Powder Fly Dip	PL 0447/4056
Young's Scab Approved Summer Mycotic Dip	

Name of Product	*Product Licence No.*
6. Anthelmintics	
Anthelworm L.	PL 1447/4072
Bonlam	PL 0025/4032
Cambendazole Paste (Bovicam)	PL 0025/4029
Equizole A Granules	PL 0025/4022
Equizole Pony Paste	PL 0025/4027
Equivurm Plus	PL 0829/4044
Equivurm Plus Paste	PL 0829/4058
Helmatac In-Feed Wormer	PL 0002/4004
Mebenvet	PL 0829/4113
Neguvon	PL 0010/4026
Nemafax	PL 0012/4003
Nemafax P Wormer Pellets	PL 0012/4150
Nemafax Premix	PL 0012/4149
Nemafax Sow Wormer Pellets Nemafax Cattle, Sheep and Goat Wormer Pellets	PL 0012/4151
Nemafax Wormer Drench for Cattle, Sheep and Goats Nemafax Wettable Powder for feed medication of cattle and sheep	PL 0012/4153
Ovitelmin Telmin Liquid	PL 0829/4114
Porcam	PL 0025/4031
Strongid—P (Granules)	PL 0057/4060
Strongid—P Paste	PL 0057/4062
Suiminth (Morantel Tartrate)	PL 0057/4063
Task	PL 1300/4002
Telmon KH	PL 0829/4112
Thibenzole Crumbles	PL 0025/4020
Thibenzole 50% Paste	PL 0025/4024
Thiprazole Granules	PL 0025/4021
Whithelmin Premix	PL 0308/4017
Young's Anthelworm	PL 1447/4064
Young's Nemtrem Cattle	PL 1447/4066
Young's Nemtrem Sheep	PL 1447/4067
7. Milk Fever Preparations	
Calcium Borogluconate Injection B Vet C 20%	PL 2848/4018
Calcium Borogluconate Injection B Vet C 30%	PL 2848/4019
Calcium Borogluconate Injection B Vet C 40%	PL 2848/4020
TVL Calcium Borogluconate "Borocal"	PL 1345/4007
8. Warble Fly Dressings	
Improved Poron	PL 1447/4054
Ruelene Ready to Use (New Formulation)	PL 0621/4017
Trolene 18	PL 0621/4014
Trolene FM	PL 0621/4013
Warbex 16.7% Famphur Pour-on	PL 0095/4024
Young's Concentrated Poron	PL 1447/4059
Young's New Poron Rycovet Warblecide Pour-on	PL 1447/4074
9. Liver Fluke Remedies	
Carbon Tetrachloride Capsules BPC	PL 1937/4012
Dirian	PL 0010/4031
Hexol	PL 1826/4000
Osmond's Fluke Drench	PL 1826/4002
Young's Nemtrem Cattle	PL 1447/4066
Young's Nemtrem Sheep	PL 1447/4067
Young's Flukol	PL 1447/4065

Name of Product	*Product Licence No.*
10. Sheep and Cattle Clostridial Vaccines	
11. Poultry Vaccines	
Addervax ND Vaccine (Living) HB1	PL 1531/4001
Avian Encephalomyelitis Vaccine (Living)	PL 1531/4005
Avian Encephalomyelitis Vaccine (Living) Calnek Strain	PL 1598/4001
Avian Encephalomyelitis Vaccine (Living) Nobilis	PL 1708/4133
Bronchimune IB Vaccine	PL 0002/4053
Combimune	PL 0002/4034
Combined ND (HB1) and IB (Massachusetts MM) Vaccine (Living)	PL 1598/4029
Combined ND La Sota and IB Vaccine (Living)	PL 1598/4028
Delvax IB H52	PL 3359/4004
Delvax IB H120	PL 3359/4003
Delvax Marek THV Freeze-dried	PL 3359/4001
Delvax ND La Sota	PL 3359/4006
Delvax ND HB1	PL 3359/4005
Eavax	PL 2592/4055
Fowl Pox Vaccine Poxine	PL 1598/4055
Fowl Pox Vaccine Poxinct	PL 1598/4053
Gumboro Disease Vaccine (Living) Nobilis	PL 1708/4139
Ibvax	PL 2592/4037
IB Vaccine (Living) Massachusetts H120 Strain	PL 0002/4003
IB Vaccine (Living) Massachusetts H52 Strain	PL 0002/4002
Inactivated ND Vaccine (oil emulsion) Newcavac Nobilis	PL 1708/4135
Lentogen HB1	PL 2592/4044
Lentogen La Sota	PL 2592/4043
Marek's Disease Vaccine, Behringwerke	PL 0086/4004
Marek's Disease Vaccine (Living) THV (Marimune)	PL 0002/4001
Marek's Disease Vaccine (Live) THV	PL 1531/4002
Marek's Disease Vaccine MD-VAC (Living) THV (Witter Strain) Frozen (Wet)	PL 1598/4021
Marek's Disease Vaccine (Lyophilised) MD-VAC	PL 1598/4027
Newcastle Disease Vaccine (Inactivated) Oil Emulsion (Layer Plus)	PL 2592/4033
Newcastle Disease Vaccine K2C (Inactivated)	PL 1531/4007
NDV (Living) La Sota	PL 1598/4000
ND Vaccine (Living) La Sota	PL 0002/4000
ND Vaccine (Live) La Sota	PL 1531/4000
ND Vaccine (Living) HB1 Strain (Newcadin L)	PL 0039/4000
ND Oil Adjuvant Vaccine (Inactivated) (Newcadin Emulsion)	PL 0039/4029
ND Vaccine (Inactivated) Oil Emulsion	PL 3318/4000
Poulvac AE	PL 1596/4034
Poulvac IB Vaccine H52 (Living)	PL 1596/4029
Poulvac IB Vaccine H120 (Living)	PL 1596/4030
Poulvac Marek THV	PL 1596/4025
Poulvac ND Vaccine (Living) HB1	PL 1596/4026
Poulvac ND Vaccine (Living) La Sota	PL 1596/4027
Tremimune	PL 0002/4005
12. Erysipelas Vaccines	
Swine Erysipelas Vaccine, Inactivated (Oil Adjuvant) Erysivax	PL 1345/4004
13. Salmonella and E. coli Vaccines	
14. Other Sheep and Cattle Vaccines	
Inactivated Pasteurella Haemalytica A9 Vaccine	PL 3520/4000
TVL Scabivax	PL 1345/4003
15. Miscellaneous Vaccines	
Pigeon Pox Vaccine (Living) Nobilis	PL 1708/4138

Name of Product	*Product Licence No.*
16. Sulphanilamide Surface Wound Dressings	
17. Local Anaesthetics	
Lignocaine Anaesthetic Injection	PL 2324/4074
Lignocaine and Adrenalin Injection	PL 2000/4029
Lignocaine Hydrochloride Injection BP 1973	PL 1393/4107
Ruby Freezaject	PL 1599/4005
18. Others	
Bloat Guard	PL 0002/4043
Bloat Guard Liquid	PL 0002/4051
Bloat Guard Drench/Therabloat	PL 0002/4054
Cheviot Veterinary Oil	PL 2613/4000
Codifer 10	PL 2545/4009
Codifer 10	PL 2545/4014
Coforta 10	PL 0010/4009
Copper (Cupric) Carbonate	PL 2987/4003
Cupric Oxide	PL 2987/4002
Cuprous Chloride	PL 2987/4001
Ferriphor	PL 1532/4026
"Flex Flac" pack for infusion 25% Dextrose Injection BP	PL 3026/4009
Gleptosil	PL 0113/4007
Haemalift	PL 1708/4121
Iron Dextran 10% (Pharmacosmos)	PL 0829/4117
Iron Dextran Injection 100 mg/ml BVC (Ronidex)	PL 3058/4002
Leodex	PL 0043/4000
Netrosylla	PL 0010/4006
Permaco C	PL 1345/4042
Permaco S	PL 1345/4041
Roscofer 10% Vet	PL 1011/4001
Roscoral Vet	PL 1011/4000
Swipoul	PL 1599/4004
Teat and Udder Cream	PL 0014/4059
Udsal	PL 0014/4060
Vache Ointment	PL 2868/4000
Vitamin AD_3E Oral	PL 1532/4020

Article 4 SCHEDULE 2: PART A

Group/Class	Substance
1. GROWTH PROMOTERS	Avoparcin Bacitracin Methylene disalicylate Bacitracin Zinc Bambermycin Dimetridazole Monensin Sodium Nitrovin Virginiamycin
2. COCCIDIOSTATS	Amprolium hydrochloride Clopidol Decoquinate Diaveridine Dinitolmide Ethopabate Methyl benzoquate Monensin Sodium Nicarbazin Pyrimethamine Robenidine Sulphaquinoxaline
3. ANTI-BLACKHEAD PREPARATIONS	Acinitrazole Aminonitrothiazole Dimetridazole Nifursol
4. ANTHELMINTICS	Haloxon Parbendazole Phenothiazine Piperazine Carbon Disulphide Complex Tetramisole Thiabendazole
5. OTHERS	Dimetridazole Menaphthone Sodium Bisulphite

SCHEDULE 2: PART B

Article 4

VETERINARY DRUGS

Name of Product	*Product Licence No.*
1. Growth Promoters	
Avotan 20 Avoparcin	PL 0095/4026
Eskalin S 400	PL 0002/4055
Eskalin 500	PL 0002/4045
Fortracin-100	PL 1654/4012
Romensin (Monensin Sodium) Premix	PL 0006/4052
Payzone 50 MA Nitrovin Milk Replacer Additive	PL 0095/4007
2. Coccidiostats	
Dinormix SR 25	PL 0109/4000
DOT Premix 12.5%	PL 1598/4032
DOT Premix 25%	PL 1598/4033
Elancoban	PL 0006/4047
Lerbek	PL 0621/4015
Nicrazin (Premix)	PL 0025/4019
Salcostat	PL 1598/4036
Supacox	PL 0025/4003
3. Anti-Blackhead Preparations	
4. Anthelmintics	
Equizole 'A' Granules	PL 0025/4022
Helmatac In-Feed Wormer	PL 0002/4004
Thibenzole Crumbles	PL 0025/4020
Thiprazole Granules	PL 0025/4021
Whithelmin Premix	PL 0308/4017
5. Others	

Article 4

SCHEDULE 3: PART A

Aklomide
Ampicillin Trihydrate
Arsanilic Acid
Benzylpenicillin
Chlortetracycline
E coli oral vaccine (Inactivated)
Erythromycin
Framycetin Sulphate
Furazolidone
4 hydroxy-3 nitrophenyl arsonic acid
Oxytetracycline
Procaine penicillin
Stilboestrol/Methyltestosterone Mixture
Sulphadimidine
Sulphanitran
Tylosin

SCHEDULE 3: PART B

Article 4

VETERINARY DRUGS

Name of Product	*Product Licence No.*
E. coli Polysaccharide Antigens for Pigs	PL 0039/4020
Engemycin 5% Soluble Powder	PL 1596/4018
Framomycin Soluble Powder 25%	PL 3317/4031
Framomycin Feed Additive	PL 3317/4023
Furazolidone BPC 68	PL 0131/4002
Furazolidone NF BVC	PL 3058/4000
Furazolidone Premix	PL 2592/4036
Granulated Tylosin Concentrate	PL 0006/4050
Intagen	PL 1754/4000
Neftin Premix	PL 0364/4003
Neftin Supplement	PL 0364/4004
Nifulidone Premix 11.6% Nifulidone Premix 22.4% Nifulidone Premix 44.8%	PL 1598/4037
Terramycin concentrate 20%	PL 0057/4061
Terramycin 5% Feed Supplement	PL 0057/4031
Terramycin TM 100	PL 0057/4065
Tylasul Premix Veterinary	PL 0006/4001

EXPLANATORY NOTE

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

On and from the appointed day, 1st February 1978, 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. This order provides for exemptions from those restrictions in respect of certain specified veterinary drugs.

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 product licence in respect of that drug, 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 complied with.

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 in respect of that drug, or by a person carrying on a business comprising the sale or supply in bulk of veterinary drugs, provided that the conditions set out in that article are complied with.

Further, the order provides for exemption in cases involving another's default, and makes certain temporary and transitional exemptions.

SI 1977/2167
ISBN 0-11-072167-5

