STATUTORY INSTRUMENTS

1989 No. 2318

MEDICINES

The Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) (No. 2) Order 1989

Made - - - -	*8th December 1989*
Laid before Parliament	*11th December 1989*
Coming into force - -	*1st January 1990*

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 (2A) and 129(4) of the Medicines Act 1968(**1**) and now vested in them(**2**), 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 in accordance with section 129(6) of that Act and with the consent of the Treasury in accordance with section 57(2A) of that Act, hereby makes the following Order:

Title and commencement

1. This Order may be cited as the Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) (No. 2) Order 1989 and shall come into force on 1st January 1990.

Interpretation and revocation

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 for the collection of produce from animals kept for the production of food and things used for the maintenance of such

(**1**) 1968 c. 67; "the appropriate Ministers" referred to in section 57 is defined in section 1 (*see also* the following footnote); section 57(2A) was inserted by the Animal Health and Welfare Act 1984 (c. 40), section 14.

(**2**) In the case of the Secretaries of State concerned with health in England and in 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 the Northern Ireland Constitution Act 1973 (c. 36), section 40 and Schedule 5, and the Northern Ireland Act 1974 (c. 28), section 1(3) and Schedule 1, paragraph 2(1)(b).

equipment, and includes any protective clothing but does not include any other kind of human apparel;

"the Department of Agriculture" means the Department of Agriculture for Northern Ireland;

"the Department of Health (N.I.)" means the Department of Health and Social Services for Northern Ireland;

"final medicated feeding stuff" means any substance, not being a medicinal product, which is for use wholly or mainly by being fed to one or more animals for a medicinal purpose, or for purposes that include that purpose, without further processing;

"fish farmer" means–

(a) a person carrying on a business of fish farming or shellfish farming which is registered in a register kept by the Minister or the Secretary of State (as the case may be) pursuant to the Registration of Fish Farming and Shellfish Farming Businesses Order 1985(**3**), or

(b) a person to whom a licence has been granted by the Department of Agriculture under section 11 of the Fisheries Act (Northern Ireland) 1966(**4**);

"intermediate feed" means a medicated feeding stuff sold, supplied or imported for use wholly or mainly as an ingredient in the preparation of a substance which is to be fed to one or more animals for a medicinal purpose or for purposes that include that purpose, with or without further processing;

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

"prescription only medicine" means a medicinal product falling within a description or class for the time being specified for the purposes of section 58 of the Act in an Order made under that section(**5**);

"qualifying businesses" means a business involving in whole or in part the retail sale of agricultural requisites;

"the Register of Manufacturers" means the register of persons entitled to incorporate medicinal products in animal feeding stuffs kept respectively by the Department of Agriculture, and the Society under regulation 6(1) of the Medicines (Medicated Animal Feeding Stuffs) Regulations 1988(**6**) and regulation 6(1) of the Medicines (Medicated Animal Feeding Stuffs) Regulations 1989(**7**);

"the Register of Merchants" means the register of merchants in veterinary drugs kept respectively by the Department of Health (N.I.), and the Society under article 3(7) of the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1985(**8**), article 5(1) of the Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) Order 1989(**9**) and article 5(1) of this Order;

"saddlery business" means a business involving in whole or in part the retail sale of saddlery requisites;

"saddlery requisites" means products and 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;

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

(**3**) S.I. 1985/1391.
(**4**) 1966 c. 17 (N.I.).
(**5**) S.I. 1989/2319.
(**6**) S.I. 1988/976.
(**7**) S.I. 1989/2320.
(**8**) S.I. 1985/1823.
(**9**) S.I. 1989/1056.

"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 Royal Pharmaceutical Society of Great Britain;

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

(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(**10**), 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" includes a veterinary drug in respect of which a product licence is granted, after the date of coming into force of this Order, containing a provision to the effect that it may be sold by retail only in accordance with a prescription by an appropriate practitioner or by a person referred to in article 3(1), 6(1), 9(1), 11(1) or 13(1) of this Order

"veterinary drug not on a general sale list" means a veterinary drug which is not 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(**11**);

"wholesale dealer" means a person for the time being carrying on a business wholly or mainly comprising the sale or supply in bulk of veterinary drugs.

(2) Unless the context otherwise requires, any reference in this Order to a numbered article or Schedule is to the article of, or Schedule to, this Order which bears that number.

(3) The Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) Order 1989(**12**) is hereby revoked.

Exemptions 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 holder of the product licence in respect of that veterinary drug, by a specially authorised person or by a person entered in the Register of Merchants as a Category I merchant if that veterinary drug is specified in the second column of Schedule 1 and the conditions contained in paragraph (2) below and article 4 are complied with.

(2) No veterinary drug described in paragraph (1) above shall be sold by retail except to a person whom the seller knows, or has reasonable cause to believe, to be a person who has animals under his control for the purposes of, and in the course of carrying on, a business, either as his sole business activity or as part of his business activities.

(3) 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 and not being a prescription only medicine by a person entered in the Register of Merchants as a Category I merchant, if–

(a) that veterinary drug–

(i) is a veterinary drug specified in Schedule 2, and

(ii) is intended for incorporation in animal feeding stuffs at a rate below 2 kilograms per tonne of the final medicated feeding stuff,

and the conditions contained in paragraph (4) below and article 4 are complied with; or

(**10**) S.I. 1971/1450, to which there are amendments not relevant to this Order.
(**11**) The current relevant Order is S.I. 1984/768.
(**12**) S.I. 1989/1056.

(b) that veterinary drug–

(i) is a veterinary drug specified in Schedule 2, and

(ii) is intended for incorporation in animal feeding stuffs at a rate of at least 2 kilograms per tonne of the final medicated feeding stuff,

and the conditions contained in paragraph (5) below and article 4 are complied with.

(4) No veterinary drug described in paragraph (3)(a) above shall be sold by retail except–

(a) to a person whose name is entered in Part A of the Register of Manufacturers, or

(b) to a fish farmer.

(5) No veterinary drug described in paragraph (3)(b) above shall be sold by retail except–

(a) to a person whose name is entered in Part A or Part B of the Register of Manufacturers, or

(b) to a fish farmer.

Further conditions for exemption under article 3

4.—(1) No veterinary drug described in article 3(1) or (3)(a) or (b) 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, and

(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,

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

(2) No veterinary drug described in article 3(1) or (3)(a) or (b) shall be sold by retail by self-service methods.

(3) In respect of any sale by retail of any veterinary drug described in article 3(1) or (3)(a) or (b) the seller shall make a record of the sale containing particulars of–

(a) the date on which the veterinary drug was sold,

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

(c) the name and address of the person to whom the veterinary drug was sold,

and shall keep such record for a period of two years from the date of the sale.

(4) No person shall, in the course of a qualifying business carried on by him, sell by retail any veterinary drug described in article 3(1) or (3)(a) or (b) unless his name is entered in the Register of Merchants as a Category I merchant in respect of each premises on which the drug is sold or stored.

(5) In paragraph (1)(c) above "premises" includes a stall of a permanent nature situated at a market or an agricultural showground.

Register of Merchants for the purpose of article 4(4)

5.—(1) The Society and the Department of Health (N.I.) shall each keep, for the purposes of article 4(4), 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 described in article 3(1) or (3)(a) or (b) free from the restrictions imposed by section 52 of the Act, if and so long as the conditions contained in articles 3 and 4 are complied with.

(2) Details of premises used for the storage of any veterinary drug described in article 3(1) or (3)(a) or (b) at a different postal address from that of premises used to sell by retail such drug shall be recorded in a register kept under paragraph (1) above.

(3) 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 of Merchants in respect of any premises on which any veterinary drug described in article 3(1) or (3)(a) or (b) is to be sold or stored by him in the course of that qualifying business, the Society shall, subject to paragraphs (8) and (9) below, enter his name in that Register in respect of those premises.

(4) Where a person who, whilst carrying on a qualifying business in Northern Ireland, makes an application in writing to the Department of Health (N.I.) for his name to be entered in the Department of Health's (N.I.) Register of Merchants in respect of any premises on which any veterinary drug described in article 3(1) or (3)(a) or (b) is to be sold or stored by him in the course of that qualifying business, the Department of Health (N.I.) shall, subject to paragraphs (8) and (9) below, enter his name in that Register in respect of those premises.

(5) Subject to paragraphs (10) and (12) below, a person whose name is entered in the Register of Merchants in respect of any premises shall, in order to retain his name in that 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 of Health (N.I.) (as the case may be) for his name to be retained in the Register of Merchants in respect of those premises.

(6) Subject to paragraphs (11) and (12) below, a person whose name is removed from the Register of Merchants in respect of any premises by reason only that he failed either to make proper application for the retention of his name in that Register pursuant to paragraph (5) above or to pay the fee due in respect of the retention of his name in that Register pursuant to paragraph (10) below may, in order to restore his name to that Register in respect of those premises, make an application to the Society or the Department of Health (N.I.) (as the case may be) for his name to be restored to the Register of Merchants in respect of those premises.

(7) There shall be paid to the Society or the Department of Health (N.I.)–

(a) in respect of the entry in the Register of Merchants of the name of any person in respect of any premises on which any veterinary drug described in article 3(1) or (3)(a) or (b) is to be sold or stored a fee of £155.00 for each such premises;

(b) in respect of the retention in the Register of Merchants of the name of any person in respect of any premises on which any veterinary drug described in article 3(1) or (3)(a) or (b) is to be sold or stored a fee of £108.00 for each such premises;

(c) in respect of the restoration to the Register of Merchants of the name of any person in respect of any premises on which any veterinary drug described in article 3(1) or (3)(a) or (b) is to be sold or stored a fee of £155.00 for each such premises.

(8) The Society or the Department of Health (N.I.) shall refuse to enter in its respective Register of Merchants the name of any person in respect of any premises unless that person–

(a) has paid to the Society or the Department of Health (N.I.) (as the case may be) the fee specified in paragraph (7)(a) above for the entry of his name in that Register; and

(b) has given to the Society or the Department of Health (N.I.) (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, as amended on 22nd November 1988, and published by the Ministry of Agriculture, Fisheries and Food (being a code relating to the sale or supply of the veterinary drugs described in article 3(1) or (3)(a) or (b)).

(9) The Society, with the approval of the Minister, or the Department of Health (N.I.), with the approval of the Department of Agriculture, may refuse to enter in its respective Register of Merchants the name of any person in respect of any premises if, in the opinion of the Society or the Department of Health (N.I.) (as the case may be), the premises are unsuitable for the storage or safekeeping of any veterinary drug described in article 3(1) or (3)(a) or (b).

(10) The Society or the Department of Health (N.I.) shall refuse to retain in its respective Register of Merchants in any year subsequent to the year in which his name is first entered in it the name of any person in respect of any premises unless that person has paid to the Society or the Department of Health (N.I.) (as the case may be) on or before 31st January in that year the fee specified in paragraph (7)(b) above for the retention of his name in that Register.

(11) The Society or the Department of Health (N.I.) shall refuse to restore to its respective Register of Merchants the name of any person in respect of any premises unless that person, having made proper application pursuant to paragraph (6) above, has paid to the Society or the Department of Health (N.I.) (as the case may be) the fee specified in paragraph (7)(c) above for the restoration of his name to that Register.

(12) The Society, with the approval of the Minister, or the Department of Health (N.I.), with the approval of the Department of Agriculture, may refuse to retain in or to restore to, or may remove from its respective Register of Merchants the name of any person in respect of any premises if, in the opinion of the Society or the Department of Health (N.I.) (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 (8)(b) above; or

(b) the conditions under which any veterinary drug described in article 3(1) or (3)(a) or (b) is sold by retail on those premises or under which it is stored on those premises (whether immediately prior to retail sale or not) are unsuitable for that purpose.

(13) In respect of any premises the Society or the Department of Health (N.I.) may remove from its respective Register of Merchants the name of any person entered in it, at the request of that person.

Exemptions for merchants in intermediate feed

6.—(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 intermediate feed by the holder of a product licence in respect of that intermediate feed, by a specially authorised person or by a person entered in the Register of Merchants as a Category I or II merchant if that intermediate feed contains a veterinary drug specified in the second column of Schedule 2 or 3 and the conditions contained in paragraphs (2) and (6) below and article 7 are complied with.

(2) No intermediate feed prescribed in paragraph (1) above shall be sold by retail except to a person whom the seller knows, or has reasonable cause to believe, to be a person who has animals under his control for the purposes of, and in the course of carrying on, a business, either as his sole business activity or as part of his business activities.

(3) The restrictions imposed by section 52 of the Act shall not apply to the sale by retail of intermediate feed by a person who is entered in the Register of Merchants as a Category I or II merchant if–

(a) that intermediate feed–

(i) contains a veterinary drug specified in Schedule 2 or 3, and

(ii) is intended for incorporation in animal feeding stuffs at a rate below 2 kilograms per tonne of the final medicated feeding stuff,

and the conditions contained in paragraph (4) below and article 7 are complied with; or

(b) that the intermediate feed–

(i) contains a veterinary drug specified in Schedule 2 or 3, and

(ii) is intended for incorporation in animal feeding stuffs at a rate of at least 2 kilograms per tonne of the final medicated feeding stuff,

and the conditions contained in paragraph (5) below and article 7 are complied with.

(4) No intermediate feed described in paragraph (3)(a) above shall be sold by retail except–

(a) to a person whose name is entered in Part A of the Register of Manufacturers, or

(b) to a fish farmer.

(5) No intermediate feed described in paragraph (3)(b) above shall be sold by retail except–

(a) to a person whose name is entered in Part A or Part B of the Register of Manufacturers, or

(b) to a fish farmer.

(6) No intermediate feed which contains a prescription only veterinary drug specified in Schedule 3 shall be sold by retail to any person (except as provided for in article 9(6)(a)) except on production by him of a veterinary written direction.

Further conditions for exemption under article 6

7.—(1) No intermediate feed described in article 6 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, and

(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,

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

(2) No intermediate feed described in article 6 shall be sold by retail by self-service methods.

(3) In respect of any sale by retail of any intermediate feed described in article 6 the seller shall make a record of the sale containing particulars of–

(a) the date on which the intermediate feed was sold,

(b) the name, identification and quantity of the intermediate feed sold, and

(c) the name and address of the person to whom the intermediate feed was sold,

and shall keep such record for a period of two years from the date of the sale.

(4) No person shall, in the course of a qualifying business carried on by him, sell by retail any intermediate feed described in article 6 unless after 1st April 1990 his name is entered in the Register of Merchants in respect of each premises on which the intermediate feed is sold or stored or he is already entered as a Category I or II merchant in the Register of Merchants kept pursuant to article 5(1).

(5) In paragraph (1)(c) above, "premises" includes a stall of a permanent nature situated at a market or an agricultural showground.

Register of Merchants for the purpose of article 7(4)

8.—(1) From 1st April 1990 the Society and the Department of Health (N.I.) shall each keep, for the purposes of article 7(4), a register of persons as being entitled, in the course of a qualifying business carried on by them, to sell by retail on premises in respect of which their names are entered in the register, any intermediate feed described in article 6 free from the restrictions imposed by section 52 of the Act, if and so long as the conditions contained in articles 6 and 7 are complied with.

(2) Details of premises used for the storage of any intermediate feed described in article 6 at a different postal address from that of premises used to sell by retail such intermediate feed shall be recorded in a register kept under paragraph (1) above.

(3) 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 of Merchants in respect of any premises on which any intermediate feed described in article 6 is to be sold or stored by him in the course of that qualifying business, the Society shall, subject to paragraphs (7) and (8) below, enter his name in that Register in respect of those premises.

(4) Where a person who, whilst carrying on a qualifying business in Northern Ireland, makes an application in writing to the Department of Health (N.I.) for his name to be entered in the Department of Health's (N.I.) Register of Merchants in respect of any premises on which any intermediate feed described in article 6 is to be sold or stored by him in the course of that qualifying business, the Department of Health (N.I.) shall, subject to paragraphs (7) and (8) below, enter his name in that Register in respect of those premises.

(5) Subject to paragraph (9) below, a person whose name is entered in the Register of Merchants in respect of any premises shall, in order to retain his name in that 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 of Health (N.I.) (as the case may be) for his name to be retained in the Register of Merchants in respect of those premises.

(6) Subject to paragraph (9) below, a person whose name is removed from the Register of Merchants in respect of any premises by reason only that he failed either to make proper application for the retention of his name in that Register pursuant to paragraph (5) above may, in order to restore his name to that Register in respect of those premises, make an application to the Society or the Department of Health (N.I.) (as the case may be) for his name to be restored to the Register of Merchants in respect of those premises.

(7) The Society or the Department of Health (N.I.) shall refuse to enter in its respective Register of Merchants the name of any person in respect of any premises unless that person has given the Society or the Department of Health (N.I.) (as the case may be) an undertaking in writing that he will comply with the provisions of the Code of Practice for Category 2 Agricultural Merchants Selling or Supplying Veterinary Drugs dated December 1989, and published by the Ministry of Agriculture, Fisheries and Food (being a code relating to the sale or supply of the intermediate feed containing the veterinary drugs described in article 6).

(8) The Society, with the approval of the Minister, or the Department of Health (N.I.) with the approval of the Department of Agriculture, may refuse to enter in its respective Register of Merchants the name of any person in respect of any premises if, in the opinion of the Society or the Department of Health (N.I.) (as the case may be), the premises are unsuitable for the storage or safe keeping of any intermediate feed described in article 6.

(9) The Society, with the approval of the Minister, or the Department of Health (N.I.) with the approval of the Department of Agriculture, may refuse to retain in or to restore to, or may remove from its respective Register of Merchants the name of any person in respect of any premises if, in the opinion of the Society or the Department of Health (N.I.) (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 (7)(b) above; or

(b) the conditions under which any intermediate feed described in article 6 is sold by retail on those premises or under which it is stored on those premises (whether immediately prior to retail sale or not) are unsuitable for that purpose.

(10) In respect of any premises the Society or the Department of Health (N.I.) may remove from its respective Register of Merchants the name of any person entered in it, at the request of that person.

Exemptions in respect of veterinary drugs to be incorporated in animal feeding stuffs and of, intermediate feed

9.—(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 nor of any intermediate feed by–

(a) the holder of the product licence in respect thereof;

(b) the holder of a product licence in respect of an intermediate feed containing or consisting of a veterinary drug;

(c) a specially authorised person;

(d) a person whose name is entered in the Register of Merchants as a Category I merchant; or, in the case of an intermediate feed, a person whose name is entered in the Register of Merchants as a Category I or II merchant;

(e) a person whose name is entered in Part A of the Register of Manufacturers; or

(f) a wholesale dealer,

if that veterinary drug–

(i) is or contains a veterinary drug specified in Schedule 2 or, in the case of an intermediate feed, specified in Schedule 2 or Schedule 3, and

(ii) is intended for incorporation in animal feeding stuffs at a rate below 2 kilograms per tonne of the final medicated feeding stuff,

and the conditions contained in paragraph (2) below and article 10 are complied with.

(2) No veterinary drug nor intermediate feed described in paragraph (1) above shall be sold by retail by any of the persons–

(a) specified in paragraph (1)(a) to (d), or, in the case of intermediate feed, specified in paragraph (1)(e), except to a person whose name is entered in Part A of the Register of Manufacturers or to a fish farmer;

(b) specified in paragraph (1)(e) or (f), except to a person whose name is entered in Part A of the Register of Manufacturers and who the seller knows, or has reasonable cause to believe, to be a person who does not have animals under his control for the purposes of, or in the course of carrying on, a business, either as his sole business activity or as part of his business activities, except for research or educational purposes only.

(3) 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 nor of any intermediate feed by–

(a) the holder of the product licence in respect thereof;

(b) the holder of a product licence in respect of an intermediate feed containing or consisting of a veterinary drug;

(c) a specially authorised person;

(d) a person whose name is entered in the Register of Merchants as a Category I merchant; or, in the case of an intermediate feed, a person whose name is entered in the Register of Merchants as a Category I or II merchant;

(e) a person whose name is entered in Part A of the Register of Manufacturers; or

(f) a wholesale dealer,

if that veterinary drug or intermediate feed–

(i) is or contains a veterinary drug specified in Schedule 2 or, in the case of an intermediate feed, specified in Schedule 2 or Schedule 3, and

(ii) is intended for incorporation in animal feeding stuffs at a rate of at least 2 kilograms per tonne of the final medicated feeding stuff,

and the conditions contained in paragraph (4) below and article 10 are complied with.

(4) No veterinary drug nor intermediate feed described in paragraph (3) above shall be sold by retail by any of the persons–

(a) specified in paragraph (3)(a) to (d), or, in the case of intermediate feed, specified in paragraph (3)(e), except to a person whose name is entered in Part A or Part B of the Register of Manufacturers or to a fish farmer;

(b) specified in paragraph (3)(e) or (f), except to a person whose name is entered in Part A of the Register of Manufacturers and who the seller knows, or has reasonable cause to believe, to be a person who does not have animals under his control for the purposes of, or in the course of carrying on, a business, either as his sole business activity or as part of his business activities, except for research or educational purposes only.

(5) 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 nor of any intermediate feed by–

(a) the holder of the product licence in respect thereof,

(b) the holder of a product licence in respect of an intermediate feed containing or consisting of such a veterinary drug,

(c) a specially authorised person,

(d) a person whose name is entered in Part A of the Register of Manufacturers and who does not have animals under his control for the purposes of, or in the course of carrying on, a business, either as his sole business activity or as part of his business activities, except for research or educational purposes, or

(e) wholesale dealer,

if that veterinary drug or intermediate feed is or contains a veterinary drug specified in Schedule 3 and the conditions contained in paragraph (6) below and article 10 are complied with.

(6) No veterinary drug nor any intermediate feed described in paragraph (3) above shall be sold by retail except–

(a) to a person whose name is entered in Part A of the Register of Manufacturers and whom the seller knows, or has reasonable cause to believe, to be a person who does not have animals under his control for the purposes of or in the course of carrying on, a business, either as his sole business activity or as part of his business activities, except for research or educational purposes only, or

(b) to a person–

(i) whose name is entered in Part A of the Register of Manufacturers and whom the seller knows, or has reasonable cause to believe, to be a person who has animals under his control for the purposes of, and in the course of carrying on, a business, either as his sole business activity or as part of his business activities but not for research or educational purposes only;

(ii) whose name is entered in Part B of the Register of Manufacturers, or

(iii) who is a fish farmer,

on production by him of a veterinary written direction.

Further conditions for exemption under article 9

10.—(1) No veterinary drug nor intermediate feed such as is described in article 9(1), (3) or (5) shall be sold by retail by self-service methods.

(2) In respect of any sale by retail of any veterinary drug described in article 9(1), (3) or (5) the seller shall make a record of the sale containing particulars of–

(a) the date on which the veterinary drug was sold,

(b) the name, quantity and, except where it is apparent from the name, the pharmaceutical form and strength of the veterinary drug sold, and the name, identification and quantity of the intermediate feed sold, and

(c) the name and address of the person to whom the veterinary drug or intermediate feed was sold,

and shall keep such record for a period of two years from the date of the sale.

Exemptions for merchants in horse wormers

11.—(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 of that veterinary drug, 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 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 the 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 described in paragraph (1)(a) above shall be sold by retail by self-service methods.

(4) In respect of any sale by retail of any veterinary drug described in paragraph (1)(a) above the seller shall make a record of the sale containing particulars of–

(a) the date on which the veterinary drug was sold, and

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

and shall keep such record for a period of two years from the date of the sale.

(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 described in paragraph (1)(a) above unless his name is entered in the register kept by the Society or the Department of Health (N.I.) under article 12(1) in respect of each premises on which the drug is sold or stored.

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

Register for the purposes of article 11(5)

12.—(1) The Society and the Department of Health (N.I.) shall each keep, for the purposes of article 11(5), 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 described in article 11(1)(a) free from the

restrictions imposed by section 52 of the Act, if and so long as the conditions contained in article 11 are complied with.

(2) Details of premises used for the storage of any veterinary drug described in article 11(1)(a) at a different postal address from that of premises used to sell by retail such drug shall be recorded in a register kept under paragraph (1) above.

(3) 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 (1) above in respect of any premises on which any veterinary drug described in article 11(1)(a) is to be sold or stored by him in the course of that qualifying business or saddlery business, the Society shall, subject to paragraphs (8) and (9) below, enter his name in that register in respect of those premises.

(4) 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 of Health (N.I.) for his name to be entered in the register kept by the Department of Health (N.I.) under paragraph (1) above in respect of any premises on which any veterinary drug described in article 11(1)(a) is to be sold or stored by him in the course of that qualifying business or saddlery business, the Department of Health (N.I.) shall, subject to paragraphs (8) and (9) below, enter his name in that register in respect of those premises.

(5) Subject to paragraphs (10) and (12) below, a person whose name is entered in the register kept by the Society or the Department of Health (N.I.) under paragraph (1) above in respect of any premises shall, in order to retain his name in 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 of Health (N.I.) (as the case may be) for his name to be retained in that register in respect of those premises.

(6) Subject to paragraphs (11) and (12) below, a person whose name is removed from the register kept by the Society or the Department of Health (N.I.) under paragraph (1) above in respect of any premises by reason only that he failed either to make proper application for the retention of his name in the register pursuant to paragraph (5) above or to pay the fee due in respect of the retention of his name in the register pursuant to paragraph (10) below may, in order to restore his name to the register in respect of those premises, make an application in writing to the Society or the Department of Health (N.I.) (as the case may be) for his name to be restored to the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) in respect of those premises.

(7) There shall be paid to the Society or the Department of Health (N.I.)–

(a) in respect of the entry in the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above of the name of any person in respect of any premises on which any veterinary drug described in article 11(1)(a) is to be sold or stored a fee of £56.00 for each such premises;

(b) in respect of the retention in the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above of the name of any person in respect of any premises on which any veterinary drug described in article 11(1)(a) is to be sold or stored a fee of £53.00 for each such premises;

(c) in respect of the restoration to the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above of the name of any person in respect of any premises on which any veterinary drug described in article 11(1)(a) is to be sold or stored a fee of £100.00 for each such premises;

except that no such fee shall be payable in respect of a person whose name is for the time being entered in, or in the course of being restored to, the Register of Merchants in respect of those premises as being a person entitled to sell or store thereon, during the course of a qualifying business carried on by him, any veterinary drug described in article 3(1) or (3)(a) or (b).

(8) The Society or the Department of Health (N.I.) shall refuse to enter in the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above the name of any person in respect of any premises unless that person–

(a) has paid to the Society or the Department of Health (N.I.) (as the case may be) the fee specified in paragraph (7)(a) above for the entry of his name in the register; and

(b) has given to the Society or the Department of Health (N.I.) (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 October 1985, as amended on 22nd November 1988, and published by the Ministry of Agriculture, Fisheries and Food (being a Code of Practice relating to the sale or supply of the veterinary drugs described in article 11(1)(a)).

(9) The Society, with the approval of the Minister, or the Department of Health (N.I.), with the approval of the Department of Agriculture, may refuse to enter in the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above the name of any person in repect of any premises if, in the opinion of the Society or the Department of Health (N.I.) (as the case may be), the premises are unsuitable for the storage or safekeeping of any veterinary drug described in article 11(1)(a).

(10) The Society or the Department of Health (N.I.) shall refuse to retain in the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above in any year subsequent to the year in which his name is first entered in it the name of any person in respect of any premises unless that person has paid to the Society or the Department of Health (N.I.) (as the case may be) on or before 31st January in that year the fee specified in paragraph (7)(b) above for the retention of his name in the register.

(11) The Society or the Department of Health (N.I.) shall refuse to restore to the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above the name of any person in respect of any premises unless that person, having made proper application pursuant to paragraph (6) above, has paid to the Society or the Department of Health (N.I.) (as the case may be) the fee specified in paragraph (7)(c) above for the restoration of his name to the register.

(12) The Society, with the approval of the Minister, or the Department of Health (N.I.), with the approval of the Department of Agriculture, may refuse to retain in or to restore to, or may remove from, the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above the name of any person in respect of any premises if, in the opinion of the Society or the Department of Health (N.I.) (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 (8)(b) above; or

(b) the conditions under which any veterinary drug described in article 11(1)(a) is sold by retail on those premises or under which it is stored on those premises (whether immediately prior to retail sale or not) are unsuitable for that purpose.

(13) In respect of any premises the Society or the Department of Health (N.I.) may remove from the register kept by the Society or the Department of Health (N.I.) (as the case may be) under paragraph (1) above the name of any person entered in it, at the request of that person.

Exemptions for pharmacists

13.—(1) The restrictions imposed by section 52(c) of the Act shall not apply to the retail sale of a veterinary drug described in article 3(1) or (3)(a) or (b) where the transaction is carried out in a registered pharmacy by a person acting on behalf of a pharmacist.

(2) The restrictions imposed by section 52 of the Act shall not apply to the supply in circumstances corresponding to retail sale of a veterinary drug such as is described in article 9(1), (3) or (5) 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.

Exemptions in cases involving another's default

14.—(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 articles 3(2), (4) or (5), 4, 6(2), (4) or (5) of a veterinary drug or intermediate feed by a person for the time being carrying on a qualifying business, which drug or intermediate feed that person, having exercised all due diligence, on reasonable grounds believes to be a veterinary drug or intermediate feed described in article 3(1) or (3)(a) or (b) or 6 but which, due to the act or default of another person, is not such a veterinary drug or intermediate feed.

(2) 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 9(2) to (6), of a veterinary drug or intermediate feed by a person for the time being carrying on a business described in article 9(1), which drug or intermediate feed that person, having exercised all due diligence, on reasonable grounds believes to be a veterinary drug or intermediate feed described in article 9(1), but which, due to the act or default of another person, is not such a veterinary drug or intermediate feed.

(3) The restrictions imposed by section 52 of the Act shall not apply to the sale by retail, in compliance with the conditions contained in articles 11(2) to (5) and 12, of a veterinary drug by a person for the time being carrying on a qualifying business or a saddlery business, which drug that person, having exercised all due diligence, on reasonable grounds believes to be a veterinary drug described in article 11(1)(a), but which, due to the act or default of another person is not such a veterinary drug.

Defences

15. Any person who, in the course of a business carried on by him, sells by retail, offers or exposes for sale by retail, or supplies in circumstances corresponding to retail sale, any intermediate feed in accordance with a forged veterinary written direction, shall not be guilty of an offence under this Order if, having exercised all due diligence, he believes on reasonable grounds that the veterinary written direction is genuine.

Kenneth Clarke
8th December 1989 — Secretary of State for Health

Sanderson of Bowden
7th December 1989 — Minister of State, Scottish Office

Peter Walker
8th December 1989 — Secretary of State for Wales

In Witness whereof the Official Seal of the Minister of Agriculture, Fisheries and Food is hereunto affixed on 7th December 1989.

L.S.

John Selwyn Gummer
Minister of Agriculture, Fisheries and Food

Sealed with the Official Seal of the Department of Health and Social Services for Northern Ireland this 8th day of December 1989.

L.S.

J. J. M. Harbison
Under Secretary, on behalf of the Permanent Secretary

Sealed with the Official Seal of the Department of Agriculture for Northern Ireland this 8th day of December 1989.

L.S.

W. J. Hodges
Permanent Secretary

We consent,

John Taylor
David Lightbown
Two of the Lords Commissioners of Her Majesty's Treasury

8th December 1989

SCHEDULE 1

VETERINARY DRUGS(**13**)

Product Licence No.	*Name of Product(**14**)*
1. Growth Promoters	
PL 2805/4002	Albac Feed Supplement 100
PL 2805/4000	Albac Feed Supplement 150
PL 2805/4001	Albac Lactodispersable 100
PL 3405/4030	Avoparcin-20
PL 3405/4019	Avoparcin 50 Premix
PL 0095/4026	Avotan 50
PL 0095/4028	Avotan 50c Avoparcin
PL 0095/4036	Avotan Super
PL 0095/4039	Avotan Farm Mix
PL 3405/4026	Bambermycin-5
PL 3405/4047	*Bambermycin-20*
PL 3405/4046	*Bambermycin-40*
PL 0010/4038	Bayo-n-ox 10% Premix
PL 3832/4020	Eskalin 20
PL 3832/4031	Eskalin 100
PL 3832/4017	Eskalin 500
PL 3832/4021	Eskalin S-400
PL 0029/4102	Fedan 10% Premix
PL 0086/4031	Flavomycin 5
PL 0086/4148	Flavomycin 80
PL 4594/4001	FPL 40 “ABCHEM”
PL 10101/4003	FPL 50 “ABCHEM”
PL 5811/4001	Intagen Premix
PL 3405/4031	Monensin-20 Ruminant
PL 3405/4022	Monensin-100 Ruminant
PL 3405/4016	Nitrovin
PL 3405/4018	Nitrovin-20
PL 0777/4002	Panazone 250-Nitrovin

(**13**) Items shown in italics did not appear in Schedule I to S.I. 1989/1056.
(**14**) Alternative product names used by specially authorised persons are not shown.

Product Licence No.	*Name of Product(**14**)*
PL 0095/4007	Payzone 50MA Nitrovin Milk Replacer Additive
PL 4188/4008	Pentazone 250
PL 0006/4052	Romensin (Monesin Sodium) Premix
PL 0006/4070	Romensin RDD
PL 0012/4170	SPIRA 200
PL 6051/4000	SPIRA 200
PL 0012/4182	SPIRA 200L
PL 6051/4001	SPIRA 200L
PL 0006/4055	Tylamix Premix 100 g/kg
PL 0006/4062	Tylamix Premix 250 g/kg
PL 3405/4028	Tylosin-20
PL 3405/4007	Tylosin 100 Premix
PL 4594/4002	Tylosin 250 “ABCHEM” Premix
PL 10101/4002	Tylosin 250 “ABCHEM” Premix
PL 3405/4027	Virginiamycin 20
PL 4594/4004	Virginiamycin 250 “ABCHEM”
PL 10101/4001	Virginiamycin 250 “ABCHEM”
PL 3405/4015	ZB-100
PL 0109/4001	Zinc Bacitracin Premix
2. Coccidiostats	
PL 0025/4008	Amprolmix
PL 0025/4035	Arpocox
PL 0031/4011	Avatec Premix
PL 0006/4077	Carbigran Premix
PL 3405/4017	Clopidol
PL 3405/4025	Clopidol 250
PL 0095/4000	Cycostat 66
PL 0095/4042	Cygro Premix
PL 0012/4188	Deccox Sheep Premix
PL 4594/4003	Dinitolmide
PL 10101/4000	Dinitolmide
PL 0109/4000	Dinormix SR 25
PL 4869/4005	D.O.T.

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

Product Licence No.	*Name of Product(**14**)*
PL 0777/4001	D.O.T.
PL 0109/4002	DOT (dinitolmide)
PL 0006/4047	Elancoban Premix
PL 3405/4034	Halofuginone-3
PL 0621/4015	Lerbek
PL 2592/4087	*Microvet Premix*
PL 4594/4000	Monensin 200 “ABCHEM” Premix
PL 10101/4004	Monensin 100 “ABCHEM” Premix
PL 3405/4006	Monensin 200
PL 3405/4032	Monensin 50 Poultry
PL 3405/4021	Monensin 100 Poultry
PL 0006/4061	Monteban 100 Premix
PL 3405/4050	*Nicarbazin-50*
PL 3405/4044	*Nicarbazin-250*
PL 0025/4019	Nicrazin (Premix)
PL 0029/4042	Pigwormer Feed Additive
PL 0086/4135	Sacox 60 Premix
	Sacox 120
PL 1598/4036	Salcostat
PL 1598/4032	Salcostat (DOT) Premix 12.5%
PL 1598/4033	Salcostat (DOT) Premix 25%
PL 3405/4033	Salinomycin-30
PL 3405/4053	*Salinomycin-60*
PL 0086/4141	*Salocin 120*
PL 0086/4117	Stenorol
PL 0086/4153	*Stenorol for Pheasants*
PL 4188/4004	Unicox Pure
3. Anti-Blackhead Preparations	
PL 3405/4009	Dazole Premix
PL 0777/4003	Dimetridazole BP (Vet)
PL 0012/4176	“Emtryl” Premix
PL 8327/4034	“Emtryl” Premix
PL 0012/4174	“Emtryl” Pure
PL 8327/4030	“Emtryl” Pure

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

Product Licence No.	*Name of Product(**14**)*
PL 0012/4175	"Emtryl" Soluble
PL 3636/4001	Lutrizole
PL 2592/4087	Microvet Premix
PL 4188/4014	Unizole T for Poultry
PL 4188/4013	Unizole T Soluble for Poultry
4. Sheep Dips and Ectoparasiticides	
PL 0676/4081	Aerosol Poultry Spray
PL 0086/4032	Alugan Aerosol Spray
PL 0086/4090	Alugan Concentrate Powder
PL 0086/4047	Alugan Dusting Spray
PL 0010/4011	Asuntol Sheep Dip
PL 5645/4015	Auriclip
PL 1300/4010	Barricade
PL 1300/4015	Barricade 5% Pour On
PL 0676/4029	Battles Lice and Mange Liquid Dressing
PL 0676/4022	Battles Mange Powder
PL 0676/4087	Battles Organo-Phosphorus Single-Dipping Fluid Dip
PL 0676/4010	Battles Special BHC Maggot Oil
PL 0676/4042	Battles Supona Based Organophosphorous Summer Fly Dip
PL 0010/4067	Bayticol Scab & Tick Dip Scab Approved
PL 0010/4069	Bayticol Pour-on
PL 0805/4018	Border Winter Dip
PL 1300/4011	C Tag 97 Fly Tag/Flectron Fly Tag Attach a Tag
PL 1300/4004	Ciodrin Insecticide
PL 1861/4042	Colydrin
PL 5869/4095	Coopers Border Winter Dip – Scab Approved
PL 5869/4082	Coopers Dipmaster Dip Scab Approved
PL 0805/4020	Cooper Fly Dip
PL 0805/4024	Cooper Lice and Mange Liquid
PL 0805/4014	Cooper Maggot Fly Spray
PL 0805/4015	Cooper MD Powder Dip
PL 5869/4005	Coopers Powerpack Summer Dip

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

Product Licence No.	*Name of Product(**14**)*
PL 5869/4002	Coopers Powerpack Winter Dip
PL 5869/4104	Coopers Scab Approved Dip (Border Type)
PL 5869/4007	Coopers Spoton Insecticide
PL 5869/4003	Coopers Summer Dip 400
PL 5869/4006	Coopers Winter Dip 200
PL 4149/4001	Deodorised Malathion Premium Grade
PL 1476/4018	Deosan Dysect
PL 5654/4020	Deosan Dysect Pour-On
PL 1476/4026	Deosan Flectron
	Deosan Flectron Fly Tag
PL 0829/4127	Dermol
PL 0676/4030	Dog Mange Cure
PL 1978/4001	Ectoral Tablets No. 1, 2 and 3
PL 1861/4030	Farm Fly Spray
PL 0113/4001	Ficare Lice and Mange Wash
PL 0676/4021	Fly and Maggot Oil
PL 5869/4000	1. Grenade Emulsifiable Concentrate 20%
	2. Stomoxin or Liquid Concentrate
PL 5869/4132	Grenade 2% Pour-On
PL 0805/4041	H/T Veterinary Demodectic
PL 0805/4036	H/T Veterinary Pb Dressing
PL 0676/4019	Improved Sheep Dip
PL 1861/4023	Keep Off
PL 4055/4012	Lice and Mange Remedy
PL 1826/4004	Lice Tick and Mange Dressing (LTM)
PL 2428/4018	Malacide
PL 0676/4012	Malathion 50 Concentrate Poultry Spray
PL 2592/4023	Microlin
PL 1826/4028	Osmonds Scab Approved Gold Fleece Sheep Dip
PL 1826/4001	Osmonds Superfleece Scab Approved Fly Dip
PL 0676/4097	Paracide Plus
PL 2428/4018	Pharmacide
PL 0038/4068	Porect

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

Product Licence No.	*Name of Product(**14**)*
PL 0086/4138	Prodip
PL 0038/4093	Ridect
PL 6939/4000	Rodgers No. 11 Scab Dip
PL 2100/4032	Rodgers 1–80 Purl Dip
PL 1447/4106	Ryposect
PL 5869/4104	Scab Approved Dip
PL 5869/4007	Spot On Insecticide
PL 0095/4041	Stockguard Insecticide Cattle Ear Tags
PL 5869/4013	Stomoxin-CY Fly Tapes
PL 5869/4009	Stomoxin Fly Tags
PL 5869/4010	Stomoxin Liquid Concentrate
PL 8566/4001	Taktic
PL 1345/4040	Taskill
PL 0086/4140	Tirade Fly Tags
PL 1728/4070	Topclip Parasol
PL 8566/4003	Topline
PL 1826/4025	Viper Dip
PL 1447/4019	Young’s Anti-Tick Smear
PL 1447/4096	Young’s Cypor
PL 1447/4053	Young’s Headfly Repellant
PL 1447/4073	Young’s Iodofenphos Winter Dip
PL 1447/4055	Young’s Killtick Liquid Tick Dip
PL 1447/4014	Young’s Maggot Oil
PL 1447/4017	Young’s Mycotic Dip
PL 1447/4056	Young’s Powder Fly Dip
PL 1447/4032	Young’s Purl Liquid Dip
PL 1447/4085	Young’s Scab Approved Diazinon Winter Dip
PL 1447/4103	Young’s Scab Approved Ectomort Summer Dip
PL 1447/4105	Young’s Scab Approved Jason Winter Dip
PL 1447/4080	Young’s Scab Approved Summer Dip
PL 1447/4060	Young’s Sheep Blowfly Spray
PL 1447/4083	Young’s SP Fly Spray
PL 3893/4069	Zeprox

5. Anthelmintics

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

Product Licence No.	*Name of Product(**14**)*
PL 0095/4037	Actelmintic Injectable Wormer
PL 1447/4092	Anthelpor
PL 5869/4128	1. Autoworm Big 5
	2. Repidose Big 5
PL 5869/4129	1. Autoworm Big 6
	2. Repidose Mid Season
PL 5869/4123	1. Autoworm Mark II (Pulse Release Cattle Wormer
	2. Repidose Mark II
PL 0010/4063	Bayverm Granules 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%
PP 0010/4062	Bayverm Roundwormer
PL 0010/4064	Bayverm SC 2.5% Suspension Worm Drench
PL 0010/4065	Bayverm SC 10% Suspension Worm Drench
PL 0010/4047	Bayverm Suspension 2.5%
PL 0010/4048	Bayverm Suspension 10%
PL 0095/4038	Cyverm Levamisole 3.2% Drench
PL 0095/4037	Cyverm Levamisole 7.5% Injection
PL 1861/4055	Day’s Worm Drench
PL 3656/4015	Dio Horse and Pony Wormer Paste
PL 8751/4011	Dio Horse and Pony Wormer Paste
PL 8669/4000	Downland Fluke and Worm Drench
PL 0010/4046	Droncit
PL 0010/4074	*Drontal Plus*
PL 5151/4001	Equidin Paste
PL 3832/4012	Equitac
PL 0025/4027	Equizole Pony Paste
PL 0025/4042	Equalan Paste for Horses
PL 0242/4013	Equivurm Plus
PL 0086/4144	Fenbendazole
PL 0242/4018	Flubenol Pellets

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

Product Licence No.	*Name of Product(**14**)*
PL 0242/4017	Flubenol Premix
PL 0010/4055	Flukombin
PL 0805/4009	Frantin Dispersable Powder
PL 0805/4007	Frantin Paste
PL 3763/4000	Gapex
PL 0201/4006	Hapadex Drench for Cattle
PL 0201/4002	Hapadex Drench for Sheep
PL 0201/4007	Hapadex Soluble Powder for Cattle
PL 0201/4008	Hapadex Soluble Powder for Sheep
PL 2592/4078	Helminate
PL 0025/4040	Ivomec Injection for Cattle
PL 0025/4043	Ivomec Injection for Pigs
PL 0025/4050	Ivomec Pour-on for Cattle
PL 2000/4081	Levacide Cattle/Levacide Poultry
PL 2000/4049	Levacide Injection
PL 2000/4050	Levacide Worm Drench
PL 8007/4011	Levadin Drench
PL 8007/4070	Levadin Injection
PL 8007/4014	Levadox
PL 8669/4001	Levadren
PL 2000/4080	Levafas Diamond
PL 2000/4068	Levafas Fluke and Worm Drench
PL 3832/4070	Loditac 20
PL 3832/4069	Loditac 200
PL 3832/4066	Loditac 3% Wormer Pellets
PL 0805/4002	Loxon Liquid
PL 0805/4000	Loxon Plus Paste
PL 0805/4003	Loxon Premix
PL 0242/4016	Mebenvet (1.2%)
PL 0242/4020	Mebenvet (5%)
PL 0829/4114	Multispec
PL 0844/4055	Multiwurma
PL 0029/4029	Nemacide

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

Product Licence No.	*Name of Product(**14**)*
PL 0012/4151	Nemafax Cattle, Sheep and Goat Wormer Pellets
PL 0012/4003	Nemafax Drench
PL 0029/4101	Nemicide Cattle Drench
PL 5869/4051	Nilvax
PL 5869/4052	Nilvax under 30kg
PL 5869/4076	Nilverm C
PL 0029/4101	Nilverm Cattle Special
PL 0029/4100	Nilverm C Cattle Drench
PL 5869/4033	Nilverm Cattle Special/Nemicide Cattle Drench
PL 0029/4013	Nilverm Drench
PL 5869/4086	Nilverm Gold
PL 0029/4039	Nilverm In-Feed
PL 0029/4040	Nilverm Injection/Bionem
PL 0029/4015	Nilverm Pig Wormer
PL 5869/4032	Nilverm Plus Drench
PL 5869/4022	Nilverm Super
PL 5869/4074	Nilzan C
PL 0029/4014	Nilzan Drench
PL 5869/4031	“Nilzan” Drench Plus
PL 5869/4030	“Nilzan” Drench Super
PL 5869/4085	Nilzan Gold
PL 0029/4041	Nilzan In-Feed
PL 5869/4021	Nilzan SC
PL 2000/4060	Norocide Worm Drench
PL 2000/4054	Novoverm Worm Drench
PL 0025/4041	Oramec Drench for Sheep
PL 0242/4008	Ovitelmin
PL 0242/4007	Ovitelmin Bolus
PL 0242/4006	Ovitelmin S&C
PL 0086/4121	Panacur 1.5% Pellets
PL 0086/4105	Panacur 2.5% Suspension
PL 0086/4110	Panacur 4% Powder
PL 0086/4106	Panacur 10% Suspension

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

Product Licence No.	*Name of Product(**14**)*
PL 0086/4107	Panacur 22% Granules
PL 0086/4119	Panacur Paste
PL 0086/4136	Panacur SC Cattle Wormer
PL 0086/4130	Panacur SC Sheep Wormer
PL 0057/4090	Paratect Flex Bolus
PL 0057/4075	Paratect Sustained Release Bolus
PL 1375/4000	Piperazine Salts BP
PL 3832/4073	Powacide
PL 0025/4000	Ranizole
PL 0025/4038	Ranizole Paste
PL 0095/4037	Ridaverm Injectable Wormer
PL 0242/4015	Ripercol pour-on
PL 0242/4005	Ripercol 3.2% Oral
PL 0242/4004	Ripercol 5% Injection
PL 0242/4003	Ripercol 7.5% Injection
PL 0242/4000	Ripercol 15% Injection
PL 0242/4079	Ripercol S+C
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 1447/4075	Rycovet Widespec
PL 5869/4075	Spectril
PL 0057/4060	Strongid-P (Granules)
PL 0057/4076	Strongid Paste for Dogs
PL 0057/4062	Strongid-P Paste
PL 0057/4079	Strongid Suspension for Dogs
PL 0057/4063	Suiminth (Morantel Tartrate)
PL 0242/4025	Supaverm
PL 0286/4032	Synanthic
PL 0286/4034	Synanthic DC
PL 0286/4039	Synanthic Horse Paste
PL 0286/4035	Synanthic Horse Pellets
PL 0286/4049	Synanthic I/R
PL 0286/4050	Synanthic Multidose 130

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

Product Licence No.	*Name of Product(**14**)*
PL 0286/4058	Synanthic Multidose Plus
PL 0286/4047	Synanthic Sel/Co
PL 5869/4061	Systamex Paste 18.5% Cattle and Horse Wormer
PL 0003/4127	Systamax Paste 18.5% Horse and Pony Wormer
PL 4869/4092	Systamex Plus Fluke
PL 5869/4084	1. Systamex Repidose
	2. Autoworm with Systamex
PL 5869/4090	Systamex Rumen Injection Cattle Wormer
PL 5869/4014	Systamex SC
PL 5869/4060	Systamex 906 Concentrated Cattle Wormer
PL 5869/4059	Systamex Worm Drench for Cattle and Sheep
PL 0242/4001	Telmin KH
PL 0242/4014	Telmin Paste
PL 4462/4002	Tetramisole Hydrochloride BP (Vet)
PL 0025/4015	Thibenzole Drench
PL 0025/4024	Thibenzole Paste
PL 3832/4022	Valbazen 2.5% 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 3832/4068	Valbazen SC 10% Total Spectrum Wormer
PL 3832/4016	Valbazen Cattle Wormer Pellets
PL 8476/4002	Verdipor
PL 6041/4002	Verdisol
PL 0012/4172	Vermadax
PL 2676/4120	Vermisole Injection
PL 2676/4121	Vermisole Worm Drench
PL 2676/4131	Vermofos
PL 0086/4139	Wormex
PL 3832/4076	Wormguard Injection
PL 1447/4091	Young’s Anthelpor 20

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

Product Licence No.	*Name of Product(**14**)*
PL 1447/4104	Young's Benafox/Rycovet Duospec
PL 1447/4100	Young's Endozal
PL 1447/4107	Young's Sure/Rycovet Bental 2.5%
PL 1447/4117	Young's Sure/Rycovet Bental 7.5%
6. Milk Fever Preparations	
PL 5271/4008	Calciflex 20
PL 5271/4009	Calciflex 40
PL 0829/4167	Calcitad 50
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 2428/4042	1. Calcium Borogluconate Injection 20% Pharmacol 20% 2. Injection of Calcium Borogluconate 20%
PL 2428/4024	Calcium Borogluconate Injection 25% with Phosphorus Magnesium and Dextrose
PL 2428/4028	Calsium Borogluconate Injection 40%
PL 2428/4027	Calcium Borogluconate Injection 30% with Phosphorus and Magnesium
PL 2324/4076	Calcium Borogluconate Solution CBG 20
PL 2428/4004	Dextrose Injection
PL 2324/4079	Glucose Saline Injection
PL 2324/4078	Injection of Calcium Borogluconate 40% and Magnesium Hypophosphite 2.2% Solution CMP 40
PL 2428/4024	Pharmacal 25 PMD
PL 2428/4027	Pharmacal 30 PM
PL 2428/4028	Pharmacal 40
PL 2428/4004	Pharmadex 50
PL 1861/4016	PMF
PL 1345/4007	TVL Calcium Borogluconate "Borocal"
7. Warble Fly Treatments	
PL 5869/4069	Coopers Warble Fly Liquid (Pour-on)
PL 0829/4127	Dermol
PL 0025/4040	Ivomec Injection

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

Product Licence No.	*Name of Product(**14**)*
PL 0038/4062	Orbisect Warble Fly, Louse and Mange Liquid for Cattle
PL 0010/4004	Tiguvon
PL 1447/4074	Young's New Poron
PL 1447/4077	Young's Poron 20
8. Liver Fluke Remedies	
PL 0010/4031	Dirian
PL 1728/4065	Fasinex 5%
PL 1728/4067	Fasinex 10%
PL 0025/4036	Flukanide
PL 0242/4023	Flukiver
PL 3832/4073	Powacide
PL 0012/4017	Trodax 20%
PL 0012/4135	Trodax 34%
PL 3832/4022	Valbazen 2.5% Total Spectrum Wormer
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 1447/4104	Young's Benafox
PL 1447/4101	Young's Flukol
PL 1447/4108	Young's Flukol (unmineralised)
PL 0029/4020	Zanil Drench
PL 0029/4044	Zanil In-Feed
9. Sheep and Cattle Clostridial Vaccines and Antisera	
PL 0086/4018	*Blackleg Vaccine*
PL 0086/4017	*Braxy/Blackleg Vaccine*
PL 0003/4022	Clovax
PL 0003/4021	Clovexin
PL 0086/4023	*Heptavac*
PL 0086/4132	Heptavac P
PL 5869/4101	Lamb Dysentery and Pulpy Kidney Antiserum BP
PL 0086/4027	*Lambisan*
PL 0086/4022	*Lambivac*
PL 5869/4051	Nilvax

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

Product Licence No.	*Name of Product(**14**)*
PL 5869/4052	Nilvax under 30 kg
PL 0003/4025	Omnivac
PL 0003/4024	Omnivac CVA
PL 0086/4020	*Ovivac*
PL 0086/4129	Ovivac P
PL 0086/4028	*Pulpy Kidney Antiserum*
PL 0003/4079	Pulpy Kidney Plus Tetanus Vaccine
PL 0086/4029	*Pulpy Kidney and Tetanus Vaccine*
PL 5869/4053	Tasvax 8
PL 1345/4023	Tasvax Gold
PL 1728/4066	Topclip 8 Sheep Vaccine
	Topclip Ewe Vaccine
10. Poultry Vaccines	
PL 5654/4017	AE
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 3832/4041	Combimune
PL 1598/4029	Combined ND (HB1) and IB (Massachusetts MM) Vaccine (Living)
PL 3359/4114	Delsuvac AR-Tox
PL 3359/4004	Delvax IB H52
PL 3359/4003	Delvax IB H120
PL 3359/4001	Delvax Marek THV Freeze-dried
PL 3359/4005	Delvax ND HB1
PL 3359/4035	Delvax ND Hitchner
PL 2592/4055	Eavax
PL 1598/4055	Fowl Pox Vaccine (Poxine)
PL 1598/4053	Fowl Pox Vaccine (Poxinet)
PL 1708/4139	Gumboro Disease Vaccine (Living) Nobilis
PL 1598/4076	Ibinac ND
PL 5654/4000	Iblin
PL 5654/4024	Iblin Bivalent

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

Product Licence No.	*Name of Product(**14**)*
PL 5654/4013	Iblin Live
PL 3832/4056	IB Vaccine (Living) Massachusetts H52 Strain
PL 3832/4036	IB Vaccine (Living) Massachusetts H120 Strain
PL 2592/4037	Ibvax
PL 1708/4135	Inactivated ND Vaccine (oil emulsion) Newcavac Nobilis
PL 1708/4066	*Infectious Bronchitis Vaccine (living) Nobilis H52*
PL 1708/4065	*Infectious Bronchitis Vaccine (living) Nobilis H120*
PL 1598/4056	Infectious Larygotracheitis Vaccine (LT-VAC)
PL 2592/4074	Ivamarek Marek’s Disease Vaccine
PL 2592/4044	Lentogen HB1
PL 5654/4018	Marek's
PL 3832/4039	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 (witter Strain) Frozen (Wet)
PL 1598/4027	Marek’s Disease Vaccine (Lyophilised) MD-VAC
PL 1708/4141	Marexine MD
PL 1442/4000	*Marexine THV*
PL 1708/4169	Marexine THV/CA
PL 5654/4023	Maridin
PL 5654/4001	Maternalin
PL 5654/4012	Maternalin Plus
PL 5654/4002	Myxilin
PL 5654/4028	Myxilin Bivalent
PL 5654/4021	Myxilin Live
PL 3318/4000	ND Vaccine (Inactivated) Oil Emulsion
PL 5654/4007	Newcadin
PL 5654/4004	Newcadin Day Old
PL 5654/4006	Newcadin 25
PL 5654/4008	Newcadin L
PL 5654/4020	Newcadin Live B-1

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

Product Licence No.	*Name of Product(**14**)*
PL 2592/4033	Newcastle Disease Vaccine (Inactivated) Oil Emulsion (Layer Plus)
PL 3832/4057	Newcastle Disease Vaccine (Living) Hitchner B1 Strain
PL 1708/4067	*Newcastle Disease Vaccine Hitchner B1 (living) Nobilis*
PL 1708/4150	Newcavac & EDS "76 Vaccine
PL 1708/4143	Nobi-Vac Egg Drop Syndrome "76 Vaccine B14 (Inactivated)
PL 1708/4155	Nobi-Vac Gumboro Inactivated
PL 1708/4184	Nobi-Vac IB+G+ND
PL 1708/4185	Nobi-Vac IB+ND
PL 1708/4187	Nobi-Vac IB+ND+EDS
PL 5654/4022	Paramyxovirus-3 Disease Vaccine
PL 0086/4039	*Pasturella Erysipelas Vaccine*
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 1596/4042	Poulvac ND+EDS
PL 1596/4026	Poulvac ND Vaccine (Living) HB1
PL 0002/4025	Tremimune
PL 3832/4024	Tremimune
PL 5654/4019	Ultravac
11. Erysipelas Vaccines	
PL 0003/4037	Erysipelas
PL 0086/4152	Erysorb Plus
PL 0086/4054	*Erysorb ST*
PL 1531/4012	Ferrovac Ery Vaccine
PL 1596/4078	Suvaxyn Erysipelas Vaccine
PL 3317/4110	Swine Erysipelas Vaccine (Inactivated)
12. Salmonella and E. coli Vaccines	
PL 0086/4013	*Bovivac*
PL 0086/4056	*Bovivac Plus*

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

Product Licence No.	*Name of Product(**14**)*
PL 0086/4134	Coliovac
PL 3832/4009	Ecopig
PL 5869/4048	Gletvax K88
PL 0086/4113	Porcovac AT
PL 3832/4004	Scourguard I
PL 1754/4002	Sow Intagen O/I Injectable
PL 1596/4076	Suvaxyn E. coli
13. Other Sheep and Cattle Vaccines	
PL 0003/4019	Blackleg Vaccine
PL 1345/4070	Footvax
PL 0003/4004	Louping I11 Vaccine
PL 0003/4135	Ovine Enzootic Abortion (Improved) Vaccine
PL 0086/4133	Ovipast
PL 0086/4094	*Pastacidin*
PL 0003/4067	Quadrivexin
PL 0003/4094	Tribovax-T
PL 0003/4069	Trivexin-T
14. Fish Vaccines	
PL 4964/4003	Aquavac – Erm
PL 3964/4004	*Aquavac Furovac/Immersion*
PL 4964/4002	Aquavac – Vibrio
PL 6149/4002	*Furogen B (Immersion) Vaccine*
15. Miscellaneous Vaccines	
PL 5869/4046	Atrovax
PL 3359/4044	Delsuvac RP
PL 1708/4195	*Nobi-Vac AR-T*
PL 1708/4152	Nobi-Vac L.T. K88
16. Local Anaesthetics	
PL 0829/4019	Crown Lignocaine with Adrenaline
PL 3317/4049	Lignavet Plus Injection
PL 2324/4074	Lignocaine Anaesthetic Injection
PL 2000/4029	Lignocaine and Adrenaline Injection
PL 1861/4021	Loconil
PL 0101/4001	M5222 Sandoz
PL 2428/4021	Pharmacaine

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

Product Licence No.	*Name of Product(**14**)*
PL 1599/4055	Ruby Freezaject
17. Others	
PL 1393/4002	Antiseptic Ointment
PL 0201/4010	AR Pac P
PL 3893/4092	Ash-fer 100
PL 5653/4028	*Auriplak*
PL 2428/4026	Bactasorb Tablets
PL 0676/4051	Battles Vitamin D3 Drench
PL 1861/4045	Black Drink
PL 6988/4000	Bloat Guard Premix
PL 0002/4054	Bloat Guard Drench
PL 3832/4064	Bloat Guard Drench
PL 3832/4065	Bloat Guard Liquid
PL 3514/4002	Boar Mate
PL 0086/4010	*Bovisan DPS*
PL 2000/4065	Calciject 20
PL 2000/4069	Calciject New Formula 40
PL 2000/4013	Calciject PMD
PL 2000/4014	Calciject 30+3
PL 2000/4015	Calciject 40
PL 1861/4009	Calcium Borogluconate 40% MP
PL 1754/4003	Calf Intagen Premix
PL 1937/4016	Castor Oil BP
PL 1393/4021	Cetrimide Udder Cream
PL 1861/4048	Cocom
PL 2545/4009	Codifer 10
PL 0010/4009	Coforta 10
PL 3317/4018	Collovet
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

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

Product Licence No.	*Name of Product(**14**)*
PL 0038/4090	Copporal 24 g
PL 0038/4078	Copprite 2 g
PL 0038/4084	Copprite 4 g
PL 0038/4087	Copprite 24 g
PL 1861/4040	Covas
PL 5869/4070	Cujec
PL 2987/4002	Cupric Oxide
PL 0832/4002	Curacho Embrocation
PL 0086/4035	Defungit
PL 2676/4127	Dextrose 20%
PL 3656/4012	Dio-Iron
PL 1596/4019	Dupharal Ade Forte
PL 0086/4026	*Ecosan*
PL 0676/4058	Ewe Drinks
PL 5869/4071	Fecundin
PL 4543/4000	Ferrodown 10
PL 3317/4044	Ferrofax 10 Plus
PL 4543/4001	Ferrodown 20
PL 0208/4003	Ferrowade
PL 0113/4005	Fisons Multivitamin Injection
PL 0113/4006	Fisons Vitamin A, D & E Injection
PL 5764/4000	Footrite
PL 1937/4015	Formaldehyde Solution
PL 2100/4018	Formalin 5
PL 0113/4007	Gleptosil
PL 2324/4079	Glucose Saline Injection
PL 1937/4015	Granulated Copper Sulphate Solution
PL 0676/4059	Green Oil
PL 0086/4029	*Grovax*
PL 0086/4021	*Haemosan*
PL 0113/4009	Hemodex
PL 0113/4012	Hemofer
PL 1754/4009	HI-FAT Baby Calf Food “Intagen”
PL 0113/4000	Imposil

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

Product Licence No.	*Name of Product(**14**)*
PL 2200/4017	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 1937/4006	Liquid Paraffin BP
PL 5271/4006	Magnaflex
PL 2428/4017	Magnesium Sulphate Injection
PL 2000/4043	Magnesium Sulphate Injection 25% w/v
PL 4127/4000	Micro Anti-Bloat Premix
PL 2592/4059	Microdex
PL 1861/4020	Morion
PL 3317/4020	Multivit Injection
PL 1861/4052	Nedasol
PL 0010/4017	Negasunt
PL 3832/4082	Nordalyte HE
PL 1861/4019	Nuphasol E
PL 1816/4053	Nuphasol 4:1
PL 0676/4090	Orfoids-Capsules for Orf
PL 1345/4042	Permaco C
PL 1345/4041	Permaco S
PL 5869/4072	Permasel-C
PL 5869/4073	Permasel-S
PL 3405/4043	Peter Hand Iron Dextran 20%
PL 2428/4017	Pharmamag 25
PL 2428/4007	Pharmavit AD_3E
PL 6128/4010	Pharmsure Iron Dextran 20%
PL 0086/4019	*Porcosan*
PL 8476/4001	Ridect FlyTags/Debantic Ear Tags
PL 0676/4067	Ringworm Ointment
PL 0829/4133	Ripercol 3.2% Oral
PL 0829/4140	Ripercol 5% Injectable Solution
PL 0829/4132	Ripercol 15% Injectable

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

Product Licence No.	*Name of Product(**14**)*
PL 1011/4001	Roscofer 10% Vet
PL 1011/4000	Roscoral Vet
PL 3821/4000	Rumbul Rumen Bullet Cattle
PL 3821/4001	Rumbul Rumen Bullet Sheep
PL 4031/4002	Rycovet Cuvine
PL 1393/4018	Scour Mixture
PL 1861/4005	Scour Mixture
PL 0086/4048	*Serovax*
PL 3317/4077	Sildex
PL 1393/4035	Skin Dressing Sulphur 22%
PL 5811/4000	Sow Intagen O/I
PL 3317/4022	Super-Suntax
PL 2100/4035	Surefoot
PL 1598/4065	Suvaxyn Iron Dextran 20% Injection
PL 1599/4004	Swipoul
PL 3862/4002	Synthite Foursure Liquid
PL 0829/4117	Tendex
PL 3317/4128	Tensolvet
PL 0676/4062	Terebene Sheep Balsam
PL 0676/4076	Trace Element Tablets
PL 5923/4002	Tracerglass C
PL 5923/4001	Tracerglass L
PL 0676/4041	Twin Lamb Remedy
PL 0676/4063	Veterinary Ointment
PL 3317/4047	Vetrivite Plus
PL 3317/4013	Vitamin A Injection
PL 3317/4015	Vitamin B12 1000 i.u.
PL 2428/4007	Vitamin ADE Solution
PL 2592/4010	Vitasol
PL 1861/4010	Wound Powder
PL 1447/4034	Young’s Antiseptic Lambing Balsam
PL 1447/4036	Young’s Swaycop

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

SCHEDULE 2 Articles 3(3)(a) and (b) and 6(1)(a)

VETERINARY DRUGS (OTHER THAN PRESCRIPTION ONLY MEDICINES) FOR INCORPORATION IN ANIMAL FEEDING STUFFS(**15**)

Product Licence No.	*Name of Product(**16**)*
1. Growth Promoters	
PL 2805/4002	Albac Feed Supplement 100
PL 2805/4000	Albac Feed Supplement 150
PL 2805/4001	Albac Lactodispersable 100
PL 3405/4030	Avoparcin-20
PL 3405/4019	Avoparcin 50 Premix
PL 0095/4026	Avotan 50
PL 0095/4028	Avotan 50c Avoparcin
PL 0095/4036	Avotan Super
PL 0095/4039	Avotan Farm Fix
PL 3405/4026	Bambermycin-5
PL 3405/4047	*Bambermycin-20*
PL 3405/4046	*Bambermycin-40*
PL 0010/4043	Bayo-n-ox 10% Premix
PL 3832/4020	Eskalin 20
PL 3832/4031	Eskalin 100
PL 3832/4017	Eskalin 500
PL 3832/4021	Eskalin S-400
PL 0029/4102	Fedan 10% Premix
PL 0086/4031	Flavomycin 5
PL 0086/4148	Flavomycin 80
PL 4594/4001	FPL 40 “ABCHEM”
PL 10101/4002	FPL 50 “ABCHEM”
PL 5811/4001	Intagen Premix
PL 3405/4031	Monensin-20 Ruminant
PL 3405/4022	Monensin-100 Ruminant
PL 3405/4016	Nitrovin
PL 3405/4018	Nitrovin-20
PL 0777/4002	Panazone 250 Nitrovin

(**15**) Items shown in italics did not apper in Schedule 2 to S.I. 1989/1056.
(**16**) Alternative product names used by specially authorised persons are not shown.

Product Licence No.	*Name of Product(**16**)*
PL 0095/4007	Payzone 50 MA Nitrovin Milk Replacer Additive
PL 4188/4008	Pentazone 250
PL 0006/4052	Romensin (Monensin Sodium) Premix
PL 0006/4070	Romensin RDD
PL 0012/4170	SPIRA 200
PL 6051/4000	SPIRA 200
PL 0012/4182	SPIRA 200L
PL 6051/4001	SPIRA 200L
PL 0006/4055	Tylamix Premix 100 g/kg
PL 0006/4062	Tylamix Premix 250 g/kg
PL 3405/4028	Tylosin-20
PL 3405/4007	Tylosin 100 Premix
PL 4594/4002	Tylosin 250 “ABCHEM” Premix
PL 10101/4002	Tylosin 250 “ABCHEM” Premix
PL 3405/4027	Virginiamycin 20
PL 4594/4004	Virginiamycin 250 “ABCHEM”
PL 10101/4001	Virginiamycin 250 “ABCHEM”
PL 3405/4015	ZB-100
PL 0109/4001	Zinc Bacitracin Premix
2. Coccidiostats	
PL 0025/4008	Amprolmix
PL 0025/4035	Arpocox
PL 0031/4011	Avatec Premix
PL 0006/4075	Carbigran Premix
PL 3405/4017	Clopidol
PL 3405/4025	Clopidol 250
PL 0095/4000	Cycostat 66
PL 0095/4042	Cygro Premix
PL 0012/4188	Deccox Sheep Premix
PL 4594/4003	Dinitolmide
PL 10101/4000	Dinitolmide
PL 0109/4000	Dinormix SR 25
PL 4869/4005	D.O.T.

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

Product Licence No.	*Name of Product(**16**)*
PL 0777/4001	D.O.T.
PL 0109/4002	DOT (dinitolmide)
PL 0006/4047	Elancoban Premix
PL 3405/6006	Monensin 200
PL 3405/4034	Halofuginone-3
PL 0621/4015	Lerbek
PL 2592/4087	*Microvet Premix*
PL 3405/4032	Monensin 50 Poultry
PL 3405/4022	Monensin 100 Ruminant
PL 4594/4000	Monensin 200 “ABCHEM” Premix
PL 10101/4004	Monensin 100 “ABCHEM” Premix
PL 0006/4061	Monteban 100 Premix
PL 3405/4050	*Nicarbazin-50*
PL 3405/4044	*Nicarbazin-250*
PL 0025/4019	Nicrazin (Premix)
PL 0029/4042	Pig Wormer Feed Additive
PL 0086/4135	Sacox 60 Premix
	Sacox 120
PL 1598/4036	Salcostat
PL 1598/4032	Salcostat (DOT) Premix 12.5%
PL 1598/4033	Salcostat (DOT) Premix 25%
PL 3405/4033	Salinomycin-30
PL 3405/4053	*Salinomycin-60*
PL 0086/4141	*Salocin 120*
PL 0086/4117	Stenorol
PL 0086/4153	*Stenorol for Pheasants*
PL 4188/4004	Unicox Pure
3. Anti-Blackhead Preparations	
PL 3405/4009	Dazole Premix
PL 0777/4003	Dimetridazole BP (Vet)
PL 0012/4176	“Emtryl” Premix
PL 8327/4034	“Emtryl” Premix
PL 0012/4174	“Emtryl” Pure
PL 8327/4030	“Emtryl” Pure

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

Product Licence No.	*Name of Product(**16**)*
PL 0012/4175	"Emtryl" Soluble
PL 3636/4001	Lutrizole
PL 2592/4087	Microvet Premix
PL 4188/4014	Unizole T for Poultry
PL 4188/4013	Unizole T Soluble for Poultry
4. Anthelmintics	
PL 0010/4049	Bayverm Premix 0.6%
PL 0010/4050	Bayverm Premix 2.4%
PL 0086/4144	Fenbendazole
PL 0242/4018	Flubenol Pellets
PL 0242/4017	Flubenol Premix
PL 3832/4070	Loditac 20
PL 3832/4069	Loditac 200
PL 0805/4003	Loxon Premix
PL 3636/4001	Lutrizole
PL 0242/4016	Mebenvet (1.2%)
PL 0242/4020	Mebenvet (5%)
PL 0029/4015	Nilverm Pig Wormer
PL 0029/4041	Nilzan In-Feed
PL 0086/4110	Panacur 4% Powder
PL 1375/4000	Piperazine Salts BP
PL 8476/4002	Verdipor
PL 6041/4002	Verdisol
PL 0086/4139	Wormex
5. Others	
PL 6988/4000	Bloat Guard Premix
PL 1754/4003	Calf Intagen Premix
PL 2987/4003	Copper (Cupric) Carbonate
PL 2987/4000	Copper Sulphate
PL 2987/4002	Cupric Oxide
PL 1754/4009	HI-FAT Baby Calf Food "Intagen"
PL 4127/4000	Micro Anti-Bloat Premix
PL 5811/4000	Sow Intagen O/I

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

SCHEDULE 3

Article 6(3)

PRESCRIPTION ONLY MEDICINES FOR INCORPORATION IN ANIMAL FEEDING STUFFS

Product Licence No.	*Name of Product(**17**)*
PL 0006/4053	Apralan Soluble Powder
PL 0012/4189	Apralan Soluble Powder
PL 0006/4057	Apralan 20 Premix
PL 0006/4058	Apralan 100 Premix
PL 0095/4014	Aureomycin Soluble Powder
PL 0095/4001	Aurofac 100 Feed Additive
PL 0095/4003	Aurofac 200 MA Milk Replacer Additive
PL 0095/4002	Cyfac HS Feed Additive
PL 3405/4010	Dazole Prescription Premix
PL 3636/4002	Dimetridazole—POM Swine and Turkeys
PL 0034/4031	Dynamutalin 2% Premix
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 3812/4019	Eskalin 500 POM for laying and breeding hens
PL 1654/4012	Fortracin BMDR
PL 3317/4031	Framoycin Soluble Powder 25%
PL 3405/4018	Furazolidone – 200
PL 3405/4012	Furazolidone BP
PL 0131/4002	Furazolidone BPC 68
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 1598/4037	Nifulidone Premix 11.6%
PL 1598/4037	Nifulidone Premix 22.4%

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

Product Licence No.	*Name of Product(**17**)*
PL 1598/4037	Nifulidone Premix 44.8%
PL 6128/4002	Pharmsure Dimetridazole 20%
PL 0034/4001	Quixalud Feed Additive
PL 0034/4026	Quixalud Premix 12%
PL 0025/4028	Ridzol 12% Premix
PL 1728/4041	Sermix
PL 4219/4000	Sulphadimidine
PL 3405/4003	Sulphadimidine
PL 3405/4020	Sulphadimidine – 100
PL 0777/4000	Sulphadimidine BP (Vet)
PL 0057/4089	Terramycin Feed Supplement
PL 0057/4080	Terramycin Feed Supplement 10%
PL 0057/4083	Terramycin Soluble Concentrate 20%
PL 0057/4084	Terramycin Soluble Powder
PL 0057/4065	Terramycin 20% Feed Supplement
PL 0057/4061	Terramycin Concentrate 20%
PL 0003/4105	Tribissen Powder
PL 0006/4045	Tylan Premix 20 g/kg
PL 0006/4001	Tylasul Premix
PL 0006/4064	Tylasul Premix 100
PL 4188/4000	Unidim
PL 4188/4007	Unidim 100
PL 4188/4003	Unidone
PL 4188/4011	Unizole S – For Pigs and Poultry
PL 4188/4012	Unizole S Soluble – For Pigs

SCHEDULE 4

Article 7(1)(a)

HORSE WORMERS

Product Licence No.	*Name of Product(**18**)*
PL 1732/4059	Astrobot 5
PL 1732/4060	Astrobot 10

(**17**) Alternative product names used by specially authorised persons are not shown.
(**18**) Alternative product names used by specially authorised persons are not shown.

Product Licence No.	*Name of Product(**18**)*
PL 0010/4063	Bayverm Granules 10%
PL 0010/4054	Bayverm LV Paste
PL 3636/4015	Dio Horse and Pony Wormer Paste
PL 5151/4001	Equidin Paste
PL 1745/4005	Equigard 5
PL 1745/4006	Equigard 10
PL 0829/4043	Equilox
PL 3832/4012	Equitac
PL 0829/4044	Equivurm Plus
PL 0829/4058	Equivurm Plus Paste
PL 0829/4043	Equivurm Syringe
PL 0025/4027	Equizole Pony Paste
PL 0025/4005	Equizole Powder
PL 0025/4042	Equvalan Paste for Horses
PL 0844/4055	Multiwurma (Horses)
PL 0086/4107	Panacur 22% Granules
PL 0086/4119	Panacur Paste
PL 0086/4106	Panacur 10% Suspension
PL 1599/4001	Ruby Horse Wormer
PL 1447/4094	Rycovet Horse and Pony Wormer Paste

EXPLANATORY NOTE

(This note is not part of the Order)

This Order revokes the Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) Order 1989.

The Order continues to provide for certain exemptions from the restrictions imposed by section 52 of the Medicines Act 1968 (“the Act”). 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 medicinal products which are specified in an Order under section 51 of the Act and which may be freely sold) to sale or supply from a registered pharmacy by or under the supervision of a pharmacist.

The Order continues to exempt from section 52 the retail sale or supply of any veterinary drug described in article 3(1) by product licence holders, by specially authorised persons (as defined in article 2(1)) or by persons carrying on a business involving, at least in part, the retail sale of

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

agricultural requisites (described as Category I merchants in the Register of Merchants) provided that the relevant conditions are satisfied (articles 3(1) and (2) and 4). Additionally the Order exempts merchants selling or supplying veterinary drugs described in article 3(3)(a) or (b) where the drugs are sold or supplied to fish farmers or to persons registered in a Register of Manufacturers kept by the Royal Pharmaceutical Society of Great Britain ("the Society") or the Department of Agriculture for Northern Ireland under regulation 6(1) of the Medicines (Medicated Animal Feeding Stuffs) Regulations 1989, provided that the relevant conditions are satisfied (articles 3(3) to (5) and 4). In the case of veterinary drugs described in article 3(1) or (3)(a) or (b), these conditions include a requirement that merchants be registered in a Register of Merchants in veterinary drugs kept by the Society or the Department of Health and Social Services for Northern Ireland (article 4(4)). Detailed requirements continue to be set out in the Order relating to registration in that Register, including provision for payment of fees, which have been increased, and giving an undertaking to comply with a specified Code of Practice (article 5).

The Order exempts from section 52 the retail sale or supply of any veterinary drug or intermediate feed described in article 6(1) by product licence holders in respect of an intermediate feed containing or consisting of such a veterinary drug, by specially authorised persons or merchants entered as Category I or II merchants in the Register of Merchants provided that the relevant conditions are satisfied (articles 6(1) and (2) and 7). Additionally the Order exempts merchants selling or supplying intermediate feed containing or consisting of veterinary drugs described in article 6 where the intermediate feed containing such drugs is sold or supplied to fish farmers or to persons registered in the Register of Manufacturers provided that the relevant conditions are satisfied (articles 6(3) to (5) and 7). In the case of intermediate feed described in article 6, these conditions include a requirement that merchants be registered from 1st April 1990 in the Register of Merchants in veterinary drugs (article 6(4)). Detailed requirements are set out in the Order relating to registration in that Register and the giving of an undertaking to comply with a specified Code of Practice (article 8). Intermediate feed containing a prescription only veterinary drug can be sold only on production of a veterinary written direction (article 6(6)) or to a person entered in Part A of the Register of Manufacturers who does not have animals under his control by way of his business (article 9(6)(a)).

The Order also continues to exempt from section 52 the retail sale or supply of specified veterinary drugs for incorporation in animal feeding stuffs and intermediate feed where the sale or supply is by product licence holders, by specially authorised persons, by wholesalers of veterinary drugs and, additionally, where the sale is by product licence holders in respect of an intermediate feed, by persons registered in the Register of Merchants as Category I merchants, by Category II merchants in the case of intermediate feed and by persons entered in Part A of the Register of Manufacturers provided that certain conditions are satisfied. These conditions now include a requirement that the sale or supply by product licence holders, specially authorised persons and Category I merchants must be only to persons registered in Part A of the Register of Manufacturers; the sale or supply by Part A manufacturers or wholesalers in veterinary drugs must be only to persons registered in Part A of the Register of Manufacturers who do not have animals under their control by way of their business (article 9(2), (4) and (6)). In the case of intermediate feed, a person entered in Part A of the Register of Manufacturers can sell only to another Part A manufacturer or to a fish farmer (article 9(2) and (4)).

The Order also continues to exempt from section 52 the retail sale or supply of any veterinary drug described in article 11(1)(a) (horse wormers) by product licence holders, by specially authorised persons, by merchants or by persons carrying on a saddlery business (as defined in article 2(1)), provided that specified conditions are complied with (article 11). In the case of merchants and persons carrying on a saddlery business, these conditions include a requirement that they be registered in the appropriate register kept by the Society or the Department of Health and Social Services for Northern Ireland (articles 11(5) and 12). The fee for entry in the register has been increased (article 12(7)(a)).

Further exemptions continue to be given in respect of the retail sale or supply of specified drugs in a registered pharmacy by persons acting on behalf of a pharmacist and to the supply of specified drugs by a pharmacist subsequent to retail sale (article 13) and in cases involving another person's default (article 14). Additionally a defence is available to any person who, having exercised all due diligence, sells or supplies any intermediate feed against a forged veterinary written direction (article 15).

The Codes of Practice referred to in the Order are priced publications and are available from MAFF Publications, London SE99 7TP.