STATUTORY INSTRUMENTS

1992 No. 33

MEDICINES

The Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) Order 1992

Made - - - -	*9th January 1992*
Laid before Parliament	*13th January 1992*
Coming into force - -	*30th January 1992*

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, and the Secretary of State and the Minister of Agriculture, Fisheries and Food, being Ministers designated(**3**) for the purposes of section 2(2) of the European Communities Act 1972(**4**) in relation to medicinal products and the common agricultural policy of the Economic Community, acting jointly, in exercise of the powers conferred on them by the said section 2(2), hereby make the following Order:

Title and commencement

1. This Order may be cited as the Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) Order 1992 and shall come into force on 30th January 1992.

(**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).

(**3**) S.I. 1972/1811.

(**4**) 1972 c. 68.

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 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(**5**), 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(**6**);

"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;

"placing on the market" means the holding for sale or disposal in any other form whatever to third parties, whether or not for consideration, and actual sale or disposal;

"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(**7**);

"qualifying business" 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 registrar under regulation 3(1) of the Medicines (Medicated Animal Feeding Stuffs) Regulations 1992(**8**);

"the Register of Merchants" means the register of merchants in veterinary drugs or intermediate feed kept respectively by the Society and the Department of Health (N.I.), under articles 5(1) and 8(1) of this Order;

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

(**5**) S.I. 1985/1391.
(**6**) 1966 c. 17 (N.I.), amended by S.I. 1991/1466.
(**7**) S.I. 1991/1392, amended by S.I. 1991/2568.
(**8**) S.I. 1992/32.

"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;

"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(**9**), 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(**10**) ;

"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) (No.2) Order 1989(**11**) , the Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) (No. 2) (Amendment) Order 1990(**12**) and the Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) (No. 2) (Amendment No. 2) Order 1990(**13**) are 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 1 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 a part of his business activities.

(**9**) S.I. 1971/1450, to which there are amendments not relevant to this Order.
(**10**) The current relevant Order is S.I. 1984/768.
(**11**) S.I. 1989/2318.
(**12**) S.I. 1990/568.
(**13**) S.I. 1990/2496.

(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 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 1 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, are complied with; or

(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, are complied with.

(4) No veterinary drug described in paragraph (3) (a) above shall be soldby 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, on the premises of the seller, condition (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, 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 1 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 purposes of article 4(4)

5.—(1) For the purposes of article 4(4) the Society and the Department of Health (N.I.) shall each continue to keep 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 £198 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 £124 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 £174 for each such premises.

(8) The Society or the Department of Health (N.I.) shall refuse to enterin 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 Distributors (Category 1 Agricultural Merchants) Registered for the Sale, Supply and Storage of Licensed Animal Medicines and Medicated Animal Feeding stuffs dated December 1991, 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.

(14) The registrar and the Department shall each furnish to the Minister, on or before 1st April in each year, a copy of the Register kept thereby certified to be a true copy of that Register as at a date specified in the certificate, not being later than 1st March in the year in questionand, pending the furnishing of a further copy of the Register in the following year, shall furnish to the Minister at monthly intervals copies of amendments made to the Register in each month following the date so specified.

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 placing on the market 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 1 or 2 merchant unless that intermediate feed has been manufactured in accordance with these Regulations pursuant to Council Directive 90/167/EEC(**14**) (laying down the conditions governing

(**14**) OJ No. L 92, 7.4.90, p.42.

the preparation, placing on the market and use of medicated feeding stuffs in the Community) and then only if that intermediate feed contains a veterinary drug specified in the second column of either Schedule 2 or Schedule 3 and the conditions contained in paragraphs (2) and (6) below and article 7 are complied with.

(2) No intermediate feed described in paragraph (1) above shall be placed on the market or 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 placing on the market of intermediate feed by a person who is entered in the Register of Merchants as a Category 1 or Category 2 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, are complied with; or

(b) that the intermediate feed—

(i) contains a veterinary drug specified in either 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, are complied with.

(4) No intermediate feed described in paragraph (3) (a) above shall beplaced on the market or 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 placed on the market or 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 placed on the market or 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 placed on the market 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 feed,

(b) in a container which has not been opened since the feed 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, feed on the premises of the seller, condition (c) above shall not apply to the subsequent delivery of that feed 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, date of the sale.

(4) No person shall, in the course of a qualifying business carried on by him, place on the market or sell by retail any intermediate feed described in article 6 unless 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 1 or Category 2 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 purposes of article 7(4)

8.—(1) For the purposes of article 7(4) the Society and the Department of Health (N.I.) shall each continue to keep a register of persons as being persons entitled, in the course of a qualifying business carried on bythem, to place on the market or 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 place on the market or 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 (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 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 (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 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 intermediate feed described in article 6 is to be sold or stored a fee of £115 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 intermediate feed described in article 6 is to be sold or stored a fee of £72 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 intermediate feed described in article 6 is to be sold or stored a fee of £101 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 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 Distributors (Category 2 Agricultural Merchants) Registered for the Sale, Supply and Storage of Medicated Animal Feeding stuffs dated December 1991, and published by the Ministry of Agriculture, Fisheries and Food (being a Code relating tothe sale or supply of intermediate feed containing the veterinary drugs 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 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 intermediate feed described in article 6.

(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 intermediate feed described inarticle 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.

(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 in respect of (a) veterinary drugs to be incorporated in animal feeding stuffs and (b) intermediate feed

9.—(1) The restrictions imposed by section 52 of the Act shall not apply to the placing on the market 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 the reof;

(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 1 merchant; or, in the case of an intermediate feed, a person whose name is entered in the Register of Merchants as a Category 1 or Category 2 merchant;

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

(f) wholesale dealer,

(i) is or contains a veterinary drug specified in Schedule 2 or, in the case of an intermediate feed, specified in either 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, are complied with.

(2) No veterinary drug nor intermediate feed described in paragraph (1) above shall be placed on the market by any of the persons—

(a) specified in paragraph (1) (a) to (d), or, in the case of an intermediate feed, specified in paragraph (1) (e), except to a personwhose 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 education purposes only.

(3) The restrictions imposed by section 52 of the Act shall not apply to the placing on the market 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 the reof;

(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 1 merchant; or, in the case of an intermediate feed, a person whose name is entered in the Register of Merchants as a Category 1 or Category 2 merchant;

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

(f) wholesale dealer,

(i) is or contains a veterinary drug specified in Schedule 2 or, in the case of an intermediate feed, specified in either 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, are complied with.

(4) No veterinary drug nor intermediate feed described in paragraph (3) above shall be placed on the market by any of the persons—

(a) specified in paragraph (3) (a) to (d), or, in the case of an 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 nameis 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 placing on the market 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) a wholesale dealer, veterinary drug specified in Schedule 3 and the conditions contained inparagraph (6) below and article 10 are complied with.

(6) No veterinary drug nor any intermediate feed described in paragraph (5) above shall be placed on the market or 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 oreducational purposes only, or

(ii) whose name is entered in Part B of the Register of Manufacturers and that veterinary drug or intermediate feed is intended for incorporation in animal feeding stuffs at a rate of at least 2 kilograms per tonne of the final medicated feeding stuff, or

(iii) who is a fish farmer,

Further conditions for exemption under article 9

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

(2) In respect of any placing on the market 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, date of the sale.

(3) No person, other than a person whose name is entered in Part A of the Register of Manufacturers shall, in the course of a business carried on by him, place on the market or sell by retail any veterinary drug or intermediate feed such as is described in article 9(1), (3) or (5) unless—

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

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

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

(4) In paragraph (3) above "the relevant particulars", in relation to a business, means the name of the business and the address, or, where appropriate, the location of every premises on which, during the course of the carrying on of that business, veterinary drugs described in article 9(1), (3) or (5) are being, or are during the next twelve months to be, sold or stored.

Exemptions for merchants in horse wormers

11.—(1) The restrictictions 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 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, 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, 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) For the purposes of article 11(5), the Society and the Department of Health (N.I.) shall each continue to keep a register of persons as being persons entitled, in the course of qualifying businesses or saddlery businesses carried on by the m, 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 anapplication 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) isto 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 £99 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 £62 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 £87 for each such premises;whose name is for the time being entered in, or in the course of beingrestored to, the Register of Merchants in respect of those premises asbeing a person entitled to sell or store the reon, 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 enterin 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 December 1991 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 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 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 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 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 inarticles 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 orintermediate feed described in article 9(1), but which, due to the actor 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, places on the market or 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.

Signed by authority of the Secretary of State for Health

Virginia Bottomley
Minister of State,
6th January 1992 Department of Health

Strathclyde
Parliamentary Under Secretary of State, Scottish
8th January 1992 Office

David Hunt
8th January 1992 Secretary of State for Wales

In witness whereof the Official Seal of the Minister of Agriculture, Fisheries and Food is hereunto affixed on 9th January 1992.

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 6th day of January 1992.

L.S.

F. A. Elliott
Permanent Secretary

Sealed with the Official Seal of the Department of Agriculture for Northern Ireland this 7th day of January 1992.

L.S.

W. J. Hodges
Permanent Secretary

We consent.

Sydney Chapman
Thomas Sackville
Two of the Lords Commissioners of Her Majesty's Treasury

7th January 1992

SCHEDULE 1 Article 3(1)

VETERINARY DRUGS

Product Licence No.	*Name of Product*
1. Growth Promoters	
PL 0006/4070	Romensin RDD
2. Coccidiostats	
PL 0025/4009	Amprol Plus Solution
3. Anti-Blackhead Preparations	
PL 0012/4175	“Emtryl” Soluble
4. Sheep Dips and Ectoparasiticides	
PL 0676/4081	Aerosol Poultry Spray
PL 5653/4028	Auriplak
PL 1300/4010	Barricade
PL 1300/4015	Barricade 5 Pour On
PL 0676/4029	Battles Lice and Mange Liquid Dressing
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 0038/4096	Canovel Insecticidal Spray
PL 1300/4011	C Tag 97 Fly Tag/Flectron Fly Tag Attach a Tag
PL 5869/4095	Coopers Border Winter Dip-Scab Approved
PL 0805/4020	Coopers Fly Dip
PL 0038/4119	Coopers Green Label Scab and Tick Dip
PL 5869/4005	Coopers Powerpack Summer Dip
PL 5869/4002	Coopers Powerpack Winter Dip
PL 5869/4104	Coopers Scab Approved Dip (Border Type)
PL 5869/4007	Coopers Spoton Insecticide
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 Fly Tag

Product Licence No.	*Name of Product*
PL 0010/4071	Diazadip All Seasons Scab Approved Sheep Dip
PL 0676/4030	Dog Mange Cure
PL 1978/4001	Ectoral Tablets No. 1, 2 and 3
PL 4055/4031	Farmers Fly and Tick Dip
PL 1447/4109	Flyte 1250
PL 5869/4000140141	**1.** Grenade Emulsifiable Concentrate 20 **2.** Stomoxin or Liquid Concentrate
PL 5869/4132	Grenade 2 Pour-On
PL 0676/4019	Improved Sheep Dip
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 1728/4029	Nuvan Top
PL 1826/4027	Osmonds Northern Fly Dip
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 1728/4070	Parasol Pour-On
PL 2428/4018	Pharmacide
PL 0038/4068	Porect
PL 0038/4093	Ridect
PL 1447/4106	Ryposect
PL 1300/4012	Shell Tirade Fly Tags
PL 1300/4014	Shell Tirade Spray
PL 0095/4041	Stockguard Insecticide Cattle Ear Tags
PL 5869/4009	Stomoxin Fly Tags
PL 8566/4001	Taktic
PL 1345/4040	Taskill
PL 1728/4006	Topclip Gold Shield Scab Approved Sheep Dip
PL 8566/4003	Topline
PL 1728/4072	Vetrazin Dip
PL 1728/4080	Vetrazin Pour-On

Product Licence No.	*Name of Product*
PL 1826/4025	Viper Dip
PL 4055/4000	Viper Winter Dip
PL 1447/4081	Young's Scab Approved Blotic Sheep Dip
PL 1447/4096	Young's Cypor
PL 1447/4016	Young's HCC Pour-On
PL 1447/4118	Young's Rycoben Sheep (Mineralised)
PL 1447/4119	Young's Rycoben Cattle (Mineralised)
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 1447/4120	Young's Seraphos Spray
5. Anthelmintics	
PL 0095/4037	Actelmintic Injectable Wormer
PL 1447/4092	Anthelpor
PL 0010/4089	Armadose Breakwormer
PL 5869/4128140141	**1.** Autoworm Mark III (Pulse Release Cattle Wormer) **2.** Repidose Big 5
PL 5869/4129140141	**1.** Autoworm 6 (Pulse Release Cattle Wormer) **2.** Repidose Mid Season
PL 5869/4123140141	**1.** Autoworm Mark II (Pulse Release Cattle Wormer) **2.** Repidose Mark II
PL 0010/4048	Bayverm Armadose
PL 0010/4058	Bayverm pellets 1.9
PL 0010/4064	Bayverm SC 2.5% Suspension Worm Drench
PL 0010/4047	Bayverm Suspension 2.5
PL 0010/4048	Bayverm Suspension 10
PL 8749/4000	Chaneverm
PL 1728/4082	Combinex Cattle
PL 1728/4081	Combinex Sheep
PL 1861/4055	Day's Worm Drench
PL 2676/4121	Decazole
PL 8751/4011	Dio Horse and Pony Wormer Paste

Product Licence No.	*Name of Product*
PL 8669/4000	Downland Fluke and Worm Drench
PL 0010/4074	Drontal Plus
PL 5151/4001	Equidin Paste
PL 3832/4013	Equitac
PL 0025/4027	Equizole Pony Paste
PL 0025/4005	Equizole Powder
PL 0025/4042	Eqvalan Paste for Horses
PL 0242/4018	Flubenol Pellets
PL 0010/4055	Flukombin
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 0201/4003	Hapadex Soluble Powder for Sheep
PL 0025/4040	Ivomec Injection for Cattle
PL 0025/4043	Ivomec Injection for Pigs
PL 2676/4131	Klomisole
PL 2000/4081	Levacide Low Volume
PL 2000/4049	Levacide Injection
PL 2000/4060	Levacide Worm Drench
PL 8007/4011	Levadin Drench
PL 8007/4010	Levadin Injection
PL 8007/4014	Levadox
PL 8669/4001	Levadren
PL 2000/4080	Levafas Diamond
PL 2000/4068	Levafas Fluke and Worm Drench
PL 3832/4013	Loditac
PL 3832/4066	Loditac 3% Wormer Pellets
PL 0242/4016	Mebenvet (1.2)
PL 0844/4055	Multiwurma
PL 0012/4003	Nemafax Drench
PL 0201/4006	Netobimin
PL 5869/4051	Nilvax
PL 5869/4033	Nilverm Cattle Special/Nemicide Cattle Drench

Product Licence No.	*Name of Product*
PL 5869/4086	Nilverm Gold
PL 0029/4040	Nilverm Injection/Bionem
PL 5869/4032	Nilverm Plus Drench
PL 5869/4022	Nilverm Super
PL 0029/4014	Nilzan Drench
PL 5869/4031	“Nilzan” Drench Plus
PL 5869/4030	“Nilzan” Drench Super
PL 5869/4085	Nilzan Gold
PL 2000/4054	Noroverm Worm Drench
PL 0025/4045	Oramec Drench for Cattle
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
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 3832/4073	Powacide
PL 0242/4015	Ripercol Pour-On
PL 0242/4005	Ripercol 3.2% Oral
PL 0242/4003	Ripercol 7.5% Injection
PL 0242/4019	Ripercol S 036C
PL 1447/4094	Rycovet Horse and Pony Wormer
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

Product Licence No.	*Name of Product*
PL 0057/4073	Strongid Tablets 35 mg
PL 0057/4074	Strongid Tablets 125 mg
PL 0242/4025	Supaverm
PL 0286/4032	Synanthic
PL 0286/4034	Synanthic DC
PL 0286/4039	Synanthic Horse Paste
PL 0286/4049	Synanthic I/R
PL 0286/4050	Synanthic Multidose 130
PL 0286/4058	Synanthic Multidose Extra
PL 0286/4066	Synanthic Multidose Plus
PL 0286/4047	Synanthic Sel/Co
PL 5869/4061	Systamex Paste 18.5% Horse and Pony Wormer
PL 0003/4127	Systamex Paste 18.5% Horse and Pony Wormer
PL 5869/4092	Systamex Plus Fluke
PL 5869/4156	Systamex Plus Fluke SC
PL 5869/4084140141	**1.** Systamex Repidose **2.** Autoworm with Systamex
PL 5869/4014	Systamex SC
PL 5869/4060140141	**1.** Systamex 906 Concentrated Cattle Wormer **2.** Systamex Handipack
PL 5869/4059	Systamex Worm Drench for Cattle and Sheep
PL 0242/4013	Telmin
PL 0242/4001	Telmin KH
PL 0242/4014	Telmin Paste
PL 4462/4002	Tetramisole Hydrochloride BP (Vet)
PL 0025/4015	Thibenzole Drench
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

Product Licence No.	*Name of Product*
PL 3832/4016	Valbazen Cattle Wormer Pellets
PL 8476/4002	Verdisol
PL 0012/4172	Vermadax
PL 2676/4120	Vermaject
PL 2676/4155	Vermisole Forte Drench
PL 2676/4120	Vermisole Injection
PL 2676/4121	Vermisole Worm Drench
PL 2676/4131	Vermofas
PL 1728/4080	Vetrazin R Pour-On
PL 3832/4076	Wormguard Injection
PL 0086/4139	Wormex
PL 1447/4091	Young's Anthelpor 20
PL 1447/4107	Young's Sure/Rycovet Bental 2.5
PL 1447/4117	Young's Sure/Rycovet Bental 7.5
PL 1447/4121	Young's 4% Ricobendazole Drench
PL 1447/4122	Young's 15
Ricobendazole Drench	
6. Milk Fever Preparations	
PL 5271/4008	Calciflex 20
PL 5271/4009	Calciflex 40
PL 0829/4167	Calcitad 50
PL 0123/4034	Calcibor CBG 20
PL 0123/4035	Calcibor CBG 40
PL 0123/4037	Calcibor CMP 30
PL 0123/4038	Calcibor CMP 40
PL 0123/4039	Calcibor CMP and D
PL 2000/4065	Calciject 20
PL 2000/4069	Calciject New Formula 40
PL 2000/4013	Calciject PMD
PL 1826/4020	Calcium Borogluconate 40
PL 1861/4009	Calcium Borogluconate 40 MP
PL 2676/4041	Calcium Borogluconate 20 PM
PL 2676/4035	Calcium Borogluconate 25 PMD
PL 2676/4042	Calcium Borogluconate 40 PM

Product Licence No.	*Name of Product*
PL 2428/4042140141	**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 2324/4076	Calcium Borogluconate Solution CBG 20
PL 2428/4004	Dextrose Injection
PL 2676/4042	Dextrose 40
PL 1596/4010	Duphafral D 3 1000
PL 2324/4078	Injection of Calcium Borogluconate 40 and Magnesium Hypophosphite 2.2 Solution CMP 40
PL 2324/4003	Injection of Calcium Borogluconate 40 w/v CBG-40 No. 2
PL 2324/4002	Injection of Calcium Borogluconate 20 w/v and Magnesium Hypophosphite 3.5 w/v CMP-20 No. 3
PL 2324/4001	Injection of Calcium Borogluconate 20 w/v and Magnesium Hypophosphite 5 w/v and Dextrose 20 w/v CMPD-20 No. 6
PL 2428/4024	Pharmacal 25 PMD
PL 2428/4027	Pharmacal 30 PM
PL 2428/4028	Pharmacal 40
PL 2428/4004	Pharmadex 50
PL 1345/4007	TVL Calcium Borogluconate “Borocal”
7. Warble Fly Treatments	
PL 0025/4046	Invomec-F Injection for Cattle
PL 0025/4040	Ivomec Injection for Cattle
PL 0025/4050	Ivomec Pour-On for Cattle
PL 0010/4004	Tiguvon
PL 1447/4077	Young’s Poron 20
Liver Fluke Remedies	
PL 1728/4065	Fasinex 5
PL 1728/4067	Fasinex 10
PL 0242/4023	Flukiver
PL 3832/4073	Powacide
PL 0012/4017	Trodax 20

Product Licence No.	*Name of Product*
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 0029/4020	Zanil Drench
9. Sheep and Cattle Clostridial Vaccines and Antisera	
PL 0003/4019	Blackleg Vaccine
PL 0086/4018	Blackleg Vaccine
PL 0086/4016	Braxy Blackleg Black Disease Vaccine
PL 0086/4017	Braxy/Blackleg Vaccine
PL 0003/4021	Covexin
PL 0086/4023	Heptavac
PL 0086/4132	Heptavac P
PL 5869/4101	Lamb Dysentery and Pulpy Kidney Antiserum BP (Vet)
PL 0086/4027	Lambisan
PL 0086/4022	Lambivac
PL 5869/4051	Nilvax
PL 0086/4020	Ovivac
PL 0086/4129	Ovivac P
PL 0086/4091	Perfrivac 8
PL 0086/4092	Perfrivac B
PL 0086/4093	Perfrivac T
PL 0086/4028	Pulpy Kidney Antiserum
PL 0003/4079	Pulpy Kidney Plus Tetanus Vaccine
PL 0086/4029	Pulpy Kidney and Tetanus Vaccine
PL 0003/4067	Quadrivexin
PL 5869/4053	Tasvax 8
PL 1345/4023	Tasvax Gold
PL 0003/4094	Tribovax-T
PL 0003/4069	Trivexin-T
10. Poultry Vaccines	
PL 5654/4017	AE

Product Licence No.	*Name of Product*
PL 1598/4075	AE Vaccine (Lyophilised)
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 3832/4033	Bronchimune
PL 3832/4041	Combimune
PL 1598/4029	Combined ND (HB 1) and IB (Massachusetts MM) Vaccine (Living)
PL 1598/4028	Combined ND (La Sota) and IB (MM) Vaccine (Living)
PL 3359/4114	Delsuvac AR-Tox
PL 3359/4075	Delvax 1BH Emulsion
PL 3359/4004	Delvax 1B H 52
PL 3359/4003	Delvax 1B H 120
PL 3359/4001	Delvax Marek THV Freeze-dried
PL 3359/4077	Delvax ND Emulsion
PL 3359/4005	Delvax ND HB 1
PL 3359/4035	Delvax ND Hitchner
PL 3359/4006	Delvax ND La Sota
PL 1708/4188	Duck Plague Vaccine Nobilis
PL 5654/4064	Edsilin
PL 1598/4055	Fowl Pox Vaccine (Poxine)
PL 1598/4053	Fowl Pox Vaccine (Poxinet)
PL 1708/4139	Gumboro Disease Vaccine (Living) Nobilis
PL 1598/4077	Ibinac
PL 1598/4076	Ibinac ND
PL 5654/4000	Iblin
PL 5654/4024	Iblin Bivalent
PL 5654/4013	Iblin Live
PL 3832/4056	IB Vaccine (Living) Massachusetts H 52 Strain
PL 3832/4036	IB Vaccine (Living) Massachusetts H 120 Strain
PL 1708/4225	IB Vaccine Nobilis MA 5

Product Licence No.	*Name of Product*
PL 1708/4135	Inactivated ND Vaccine (oil emulsion) Newcavac Nobilis
PL 1708/4066	Infectious Bronchitis Vaccine (Living) Nobilis H 52
PL 1708/4065	Infectious Bronchitis Vaccine (Living) Nobilis H 120
PL 1598/4056	Infectious Larygotracheitis Vaccine (LT VAC)
PL 5654/4018	Marek's
PL 3832/4039	Marek’s Disease Vaccine (Living) THV (Strain FC 126) Freeze-dried (Marimune)
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 1708/4149	Marexine HVT Vaccine
PL 1708/4141	Marexine MD
PL 1442/4000	Marexine THV
PL 1942/4000	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 5654/4029	Myxilin EDS
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
PL 3832/4057	Newcastle Disease Vaccine (Living) Hitchner B 1 Strain
PL 1598/4000	Newcastle Disease Vaccine (Living) La Sota Strain
PL 3832/4053	Newcastle Disease Vaccine (Living) La Sota Strain

Product Licence No.	*Name of Product*
PL 1708/4067	Newcastle Disease Vaccine Hitchner B 1 (Living) Nobilis
PL 1708/4142	Newcastle Disease Vaccine Living (Nobilis Clone 30)
PL 1708/4068	Newcastle Disease Vaccine (Living) La Sota
PL 1708/4150	Newcavac & EDS `76 Vaccine
PL 1708/4143	Nobi-Vac Egg Drop Syndrome `76 Vaccine B 14 (Inactivated)
PL 1708/4167	Nobi-Vac Gumboro + EDS `76
PL 1708/4155	Nobi-Vac Gumboro Inactivated
PL 1708/4158	Nobi-Vac Gumboro + ND
PL 1708/4184	Nobi-Vac IB + G + ND
PL 1708/4185	Nobi-Vac IB + ND
PL 1708/4187	Nobi-Vac IB + ND + EDS
PL 1708/4157	Nobi-Vac Triple GNE
PL 1598/4008	Pabac
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 H 52 (Living)
PL 1596/4030	Poulvac IB Vaccine H 120 (Living)
PL 1596/4027	Poulvac La Sota
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) HB 1
PL 0002/4025	Tremimune
PL 3832/4024	Tremimune
PL 5654/4019	Ultravac
11. Erysipelas Vaccines and Antisera	
PL 0086/4164	Colisorb
PL 0086/4152	Erysorb Plus
PL 0086/4054	Erysorb ST
PL 1531/4012	Ferrovac Ery Vaccine

Product Licence No.	*Name of Product*
PL 1596/4078	Suvaxyn Erysipelas Vaccine
PL 0086/4007	Swine Erysipelas Antiserum
PL 3317/4110	Swine Erysipelas Vaccine (Inactivated)
PL 0003/4037	Swine Erysipelas Vaccine
12. Salmonella and E. Coli Vaccines and Antisera	
PL 0086/4010	Bovisan DPS
PL 0086/4013	Bovivac
PL 0086/4056	Bovivac Plus
PL 0086/4134	Coliovac
PL 3832/4009	Ecopig
PL 0086/4026	Ecosan
PL 5869/4050	Gletvax 5
PL 0086/4049	Grovax
PL 8327/4063	Imocolibov
PL 1708/4157	Nobi-Vac K 99
PL 1708/4186	Nobi-Vac Porcol 5
PL 0086/4012	Porcovac
PL 0086/4113	Porcovac AT
PL 0086/4163	Porcovac Plus
PL 3832/4004	Scourguard I
PL 0086/4048	Serovax
PL 5811/4000	Sow Intagen O/I
PL 1596/4076	Suvaxyn E. Coli P 4
13. Other Sheep and Cattle Vaccines and Antisera	
PL 0003/4022	Clovax
PL 5869/4047	Footvax
PL 0003/4004	Louping I 11 Vaccine
PL 5869/4041	Ovine Enzootic Abortion Vaccine (Inactivated)
PL 0086/4133	Ovipast
PL 0086/4094	Pastacidin
PL 3317/4021	Pneumovac
PL 3317/4012	Pneumovac Plus
PL 6078/4007	Websters Vaxall Norot Vaccine

Product Licence No.	*Name of Product*
14. Fish Vaccines	
PL 4964/4001	Aeromonas Salmonicide Vaccine
PL 4964/4000	Aeromonas Salmonocide Vibrio A
PL 4964/4005	Aquavac Cyprivac CE
PL 4964/4003	Aquavac-Erm
PL 4964/4004	Aquavac Furovac/Immersion
PL 4964/4002	Aquavac-Vibrio
PL 6149/4000	Ermogen
PL 5869/4106	Fiskevac ERM
PL 5869/4102	Fiskevax-V
PL 6149/4002	Furogen B (Immersion) Vaccine
15. Miscellaneous Vaccines and Antisera	
PL 0201/4010	AR-PAC P
PL 0003/4017	Carovax
PL 3359/4044	Delsuvac RP
PL 2460/4000	Duck Hepatitis Virus Vaccine Living Attenuated
PL 0086/4021	Haemosan
PL 1708/4195	Nobi-Vac AR-T
PL 1708/4152	Nobi-Vac L.T. K 88
PL 1708/4163	Pigeon Pox Vaccine (Living) Nobilis
16. Local Anaesthetics	
PL 3317/4049	Lignavet Plus Injection
PL 2324/4074	Lignocaine Anaesthetic Injection
PL 2000/4029	Lignocaine and Adrenaline Injection
PL 0123/4054	Lignol
PL 2674/4099	Locovetic
PL 0101/4001	MS 222 Sandoz
PL 2428/4021	Pharmacaine
PL 0123/4068	Willcain
7. Others	
PL 2324/4043	Acet Ade
PL 3893/4092	Ash-fer 100
PL 6988/4000	Bloat Guard Premix
PL 6988/4000	Bloat Guard Drench

Product Licence No.	*Name of Product*
PL 3832/4065	Bloat Guard Liquid
PL 1861/4048	Cocom
PL 8751/4007	Codifer 8
PL 0010/4009	Coforta 10
PL 5869/4159	Coopercare 1
PL 0012/4184	Copacaps
PL 3317/4010	Copper Methionine Injection
PL 1345/4066	Copagro
PL 0038/4088	Copporal 2 g
PL 0038/4089	Copporal 4 g
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 0832/4002	Curacho Embrocation
PL 0123/4088	Dalophylline Gel
PL 0086/4035	Defungit
PL 2324/4006	Injection of Dextrose 50 D-50 No. 8
PL 2676/4127	Dextrose 20
PL 0100/4047	Dicarocide Fortie Injection
PL 3656/4012	Dio-Iron
PL 1596/4019	Duphafral Ade Forte
PL 4543/4000	Ferrodown 10
PL 3317/4044	Ferrofax 10 Plus
PL 4543/4001	Ferrodown 20
PL 0208/4003	Ferrowade
PL 0113/4005	Fisons Multivitamins Injection
PL 0113/4006	Fisons Vitamin A, D & E Injection
PL 5764/4000	Footrite
PL 1937/4015	Formaldehyde Solution
PL 0113/4007	Gleptosil
PL 1937/4011	Granulated Copper Sulphate

Product Licence No.	*Name of Product*
PL 0113/4012	Hemofer
PL 0113/4000	Imposil
PL 1826/4015	Improved Medicine for Slow Fever
PL 2000/4017	Intravit 12
PL 0829/4117	Iron Dextran 10 (Pharmacosmos)
PL 2676/4049	IRT Concentrated
PL 0829/4011	Kopertox Aerosol
PL 0043/4000	Leodex 10
PL 0043/4036	Leodex 10 Plus
PL 0043/4042	Leodex 20
PL 5271/4006	Magnaflex
PL 5271/4002	Magniflex
PL 2324/4008	Injection of Magnesium Sulphate 25 w/v MS-25 No. 9
PL 2676/4047	Magnesium Sulphate
PL 0123/4054	Magnesium Sulphate Injection
PL 2000/4043	Magnesium Sulphate Injection 25 w/v
PL 2592/4059	Microdex
PL 1861/4020	Morion
PL 3317/4020	Multivet Injection
PL 2676/4013	Multivitamin Injection
PL 2000/4023	Multivitamin Injection
PL 1861/4052	Nedasol
PL 0010/4070	Negasunt
PL 3832/4082	Nordalyte HE
PL 1861/4019	Nuphasol E
PL 1861/4053	Nuphasol 4:1
PL 0676/4090	Orfoids-Capsules for Orf
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 3832/4044	Pragmatar Shampoo
PL 8476/4001	Ridect Fly Tags/Debantic Ear Tags
PL 1011/4001	Roscofer 10 Vet

Product Licence No.	*Name of Product*
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 1861/4005	Scour Mixture
PL 3317/4077	Sildex
PL 1598/4007	Spray Diluent
PL 2676/4009	Supercal 20
PL 2676/4041	Supercal 20 PM
PL 2676/4035	Supercal 25 PMD
PL 2676/4040	Supercal 30 PM
PL 2676/4010	Supercal 40
PL 2676/4042	Supercal 40 PM
PL 2100/4035	Surefoot
PL 1598/4065	Suvaxyn Iron Dextran 20 Injection
PL 3862/4000	Synthite Foursure Liquid
PL 0829/4117	Tendex
PL 3317/4128	Tensolvet
PL 0676/4062	Terebene Sheep Balsam
PL 2676/4011	Triave
PL 1393/4023	Veterinary Iron Injection
PL 3317/4047	Vetrivite Plus
PL 0829/4121	Vital Multivitamin Solution
PL 3317/4015	Multivet Vitamin B 12 Injection 1000 054g/ml
PL 0829/4032	Vitamin B 12 Injection
PL 2592/4010	Vitasol
PL 0208/4002	Wade Iron 10
PL 0208/4001	Wade Iron 20
PL 0012/4122	Water for Injections
PL 1861/4010	Wound Powder
PL 1447/4037	Young’s Bovicoppa

Items shown in italics did not appear in the Schedule to S.I. 1990/2496.

Product Licence No.	*Name of Product*
	Alternative product names used by specially authorised persons are not shown.

SCHEDULE 2

Articles 3(3) (a) and (b), 6(1), (3) (a) and (b), 9(1) and (3)

VETERINARY DRUGS (OTHER THAN PRESCRIPTION ONLYMEDICINES) FOR INCORPORATION IN ANIMAL FEEDING STUFFS

Product Licence No.	*Name of Product*
1. Growth Promoters	
PL 2805/4005	Albac Feed Supplement 22
PL 2805/4003	Albac Feed Supplement 55
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 0095/4053	Avotan Super G
PL 0095/4039	Avotan Farm Mix
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 5869/4099	Fedan 10% Premix
PL 0086/4031	Flavomycin 5
PL 0086/4148	Flavomycin 80
PL 10101/4003	FPL 50 “ABCHEM”

Product Licence No.	*Name of Product*
1. Growth Promoters	
PL 2805/4005	
PL 5811/4001	Intagen Premix
PL 0006/4077	Maxus 25 Premix Poultry
PL 0006/4068	Maxus G 100
PL 10101/4004	Monensin 100 “ABCHEM”
PL 3405/4031	Monensin-20 Ruminant
PL 3405/4022	Monensin-100 Ruminant
PL 0006/4052	Romensin G 100
PL 2411/4000	Rumenox-20
PL 2411/4001	Rumenox-100
PL 0086/4141	Salocin 120
PL 6051/4000	Spira 200
PL 6051/4001	Spira 200L
PL 0006/4051	Tylamix 20 g/kg
PL 0006/4055	Tylamix G 100
PL 0006/4062	Tylamix G 250
PL 3405/4028	Tylosin-20
PL 3405/4007	Tylosin 100 Premix
PL 10101/4002	Tylosin 250 “ABCHEM” Premix
PL 3405/4027	Virginiamycin 20
PL 10101/4001	Virginiamycin 250 “ABCHEM”
PL 3405/4015	ZB-100
PL 3405/4005	ZB 150
PL 0109/4001	Zinc Bacitracin Premix
2. Coccidiostats	
PL 0025/4008	Amprolmix
PL 0031/4011	Avatec Premix
PL 0006/4075	Carbigran Premix
PL 3405/4017	Clopidol
PL 3405/4025	Clopidol 250
PL 0621/4001	Coyden 25
PL 0095/4000	Cycostat 66
PL 0095/4042	Cygro Premix
PL 8327/4038	Deccox Pure

Product Licence No.	*Name of Product*
1. Growth Promoters	
PL 2805/4005	
PL 0012/4052	Deccox Premix Decoquinate 6
PL 8327/4066	Deccox Sheep Premix
PL 10101/4000	Dinitolmide
PL 0109/4000	Dinormix SR 25
PL 0109/4002	DOT (Dinitolmide)
PL 0006/4047	Elancoban G 200
PL 3405/4006	Monensin 200
PL 3405/4034	Halofuginone-3
PL 8327/4049	Lerbek
PL 0006/4078	Maxiban G 160
PL 3405/4032	Monensin-50 Poultry
PL 3405-4031	Monensin-20 Ruminant
PL 3405/4021	Monensin-100 Poultry
PL 3405/4022	Monensin-100 Ruminant
PL 3405/4006	Monensin-200
PL 10101/4004	Monensin 100 “ABCHEM” Premix
PL 0006/4061	Monteban G 100
PL 3405/4050	Nicarbazin-50
PL 3405/4044	Nicarbazin-250
PL 0025/4019	Nicrazin (Premix)
PL 0086/4135	Sacox 120
PL 1598/4036	Salcostat
PL 1598/4033	Salcostat (DOT) Premix 25
PL 3405/4033	Salgain-30
PL 3405/4053	Salgain-60
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 8327/4034	“Emtryl” Premix
PL 8327/4030	“Emtryl” Pure
PL 3636/4001	Lutrizol — PML Turkeys

Product Licence No.	*Name of Product*
1. Growth Promoters	
PL 2805/4005	
PL 2592/4081	Microvet Premix
4. Anthelmintics	
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 3832/4066	Loditac 3% Wormer Pellets
PL 3636/4001	Lutrizole
PL 0242/4016	Mebenvet (1.2%)
PL 0242/4020	Mebenvet (5%)
PL 8327/4020	Nemafax 14
PL 0086/4110	Penacur 4% Powder
PL 8476/4000	Verdipor
PL 8476/4002	Verdisol
5. Others	
PL 6988/4000	Bloat Guard Premix
PL 2987/4003	Copper (Cupric) Carbonate
PL 2987/4000	Copper Sulphate
PL 2987/4002	Cupric Oxide
	Items shown in italics did not appear in the Schedule to S.I. 1990/2496.
	Alternative product names used by specially authorised persons are not shown.

SCHEDULE 3

Articles 6 (1), (3) (a) and (b), (6) and 9(1), (3) and (5)

PRESCRIPTION ONLY MEDICINES FOR INCORPORATION IN ANIMAL FEEDINGSTUFFS

Product Licence No.	*Name of Product*
PL 8007/4019	Amoxinsol 50
PL 3832/4078	Antibac 100

Product Licence No.	*Name of Product*
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 0012/4191	Apralan 100 Premix
PL 3405/4059	Aquacil
PL 3405/4056	Aquinox
PL 0095/4046	Aureomycin Soluble Powder
PL 2592/4088	Aureosup 100
PL 0095/4048	Aurofac 100 Feed Additive
PL 0095/4047	Aurofac 200 MA Milk Replacer Additive
PL 4188/4018	Auromix 100
PL 1728/4076	Cosumix Plus
PL 3405/4024	CTC-100
PL 0095/4052	Cyfac HS Feed Additive
PL 3405/4010	Dazole Prescription Premix
PL 1596/4113	Duphatrim Poultry Formula
PL 0201/4019	Euphyllin
PL 0012/4159	“Emtryl” Prescription Premix
PL 0012/4160	“Emtryl” Prescription Pure
PL 0012/4161	“Emtryl” Prescription Soluble
PL 1596/4018	Engemycin 5 Soluble Powder
PL 3317/4023	Framomycin Feed Additive
PL 5869/4125	Fulcin Feed Additive
PL 3405/4023	Furazolidone-200
PL 3405/4012	Furazolidone BP
PL 0131/4002	Furazolidone BPC 68
PL 6687/4000	Furazolidone BV
PL 3058/4000	Furazolidone NF BVC
PL 2592/4036	Furazolidone Premix
PL 8007/4022	Grisol-V-Granules
PL 5654/4071	Grisovin Powder
PL 0032/4084	Lincocin Premix
PL 3636/4002	Lutrizol — POM Swine and Turkeys

Product Licence No.	*Name of Product*
PL 2592/4083	Micro-Bio Sulphadimidine Premix
PL 2592/4084	Micro-Bio Sulphadimidine Pure
PL 2592/4085	Microfac H P
PL 8007/4026	Navilox
PL 0032/4111	Neobiotic Soluble Powder 70
PL 1598/4037	Nifulidone Premix 11.6
PL 1598/4037	Nifulidone Premix 22.4
PL 1598/4037	Nifulidone Premix 44.8
PL 2000/4084	Norofulvin Granules
PL 6128/4002	Pharmsure Dimetridazole 20
PL 3405/4054	Potencil
PL 3405/4058	Sulfatrim
PL 3405/4055	Synutrim 300
PL 3405/4061	Synutrim 30 Soluble
PL 0057/4089	Terramycin Feed Supplement 20
PL 0057/4080	Terramycin Feed Supplement 10
PL 0057/4083	Terramycin Soluble Powder Concentrate 20
PL 0057/4084	Terramycin Soluble Powder
PL 8007/4018	Tetcin Premix
PL 4188/4020	Tetramin 100
PL 4188/4023	Tetramin 200
PL 3405/4057	Tetraplex
PL 0043/4045	Tiamutin 2 Premix
PL 0043/4057	Tiamutin 25 Premix
PL 5869/4119	Tribrissen SQX Poultry Formula
PL 8007/4023	Trimediazine 15 Premix
PL 8007/4024	Trimediazine 30 Premix
PL 0006/4056	Tylan Premix
PL 0006/4045	Tylan Premix 20 g/kg
PL 0006/4056	Tylan Premix 100 g/kg
PL 0006/4088	Tylasul Premix 50
PL 0006/4073	Tylasul Premix 100
PL 4188/4016	Uniprim 150
PL 4188/4011	Unizole S-For Pigs and Poultry

Product Licence No.	*Name of Product*
PL 2987/4004	Zinc Oxide Items shown in italics did not appear in the Schedule to S.I. 1990/2496. Alternative product names used by specially authorised persons are not shown.

SCHEDULE 4

Article 11(1) (a)

HORSE WORMERS

Product Licence No.	*Name of Product*
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 3832/4012	Equitac
PL 0025/4027	Equizole Pony Paste
PL 0025/4005	Equizole Powder
PL 0025/4042	Eqvalan Paste for Horses
PL 0844/4055	Multiwurma (Horses)
PL 0086/4107	Panacur 22% Granules
PL 0086/4119	Panacur Paste
PL 0086/4106	Panacur 10 Suspension
PL 1447/4094	Rycovet Horse and Pony Wormer Paste
PL 0057/4060	*Strongid-P Granules*
PL 0057/4062	*Strongid-P Paste*
PL 0286/4039	Synanthic Horse Paste
PL 5869/4061	Systamex Paste
PL 0242/4013	Telmin
PL 0242/4014	Telmin Paste

(a) Items shown in italics did not appear in the Schedule to S.I. 1990/2496.
(b) Alternative product names used by specially authorised persons are not shown.

EXPLANATORY NOTE

(This note is not part of the Order)

This Order revokes the Medicines (Veterinary Drugs) (Pharmacy and Merchants' List) (No. 2) Order 1989, as amended.

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 specified persons subject to certain conditions (article 3).

The Order continues to exempt from section 52 the placing on the market of any veterinary drug incorporated in an intermediate feed by specified persons subject to certain conditions (article 6).

Pursuant to article 2 of Council Directive 90/167/EEC (laying down the conditions governing the preparation, placing on the market and use of medicated feeding stuffs in the Community, OJ No. L 92, 7.4.90 p.42) a definition of "placing on the market" has been inserted in article 2(1) and the consequential amendments made to articles 6, 7, 8, 9 and 10 in respect of provisions governing intermediate feed. This amendment is made under the provisions of section 2(2) of the European Community Act 1972.

The Order also continues to exempt from section 52 the retail sale or supply of specified veterinary drugs for incorporation in animal feeding stuffs where the sale or supply is by specified persons subject to certain conditions (article 9).

Pursuant to article 4 of Council Directive 90/167/EEC the period of time that the record of any sale of veterinary drugs or intermediate feed is required to be kept pursuant to articles 4(3), 7(3) and 10(2) is extended from two to three years.

The Order also continues to exempt from section 52 the retail sale or supply of any veterinary drug (horse wormers) by specified persons subject to certain conditions (article 11).

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).

The fees for the registration, retention and restoration in the Register of Category 1 agricultural merchants (article 5(7)) and for saddlers (article 12(7)) have been slightly increased; the fees for the registration, retention and restoration in the Register of Category 2 agricultural merchants (article 8(7)) have been slightly decreased.

The Codes of Practice referred to in the Order have been updated and are priced publications available from MAFF Publication, London SE 997TP.