STATUTORY INSTRUMENTS

1985 No. 1823

MEDICINES

The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1985

Made - - - -	*25th November* 1985
Laid before Parliament	*9th December* 1985
Coming into Operation	*1st January* 1986

The Secretary of State concerned with health in England, the Secretaries of State respectively concerned with health and with agriculture in Scotland and in Wales, the Minister of Agriculture, Fisheries and Food, the Department of Health and Social Services for Northern Ireland, and the Department of Agriculture for Northern Ireland, acting jointly, in exercise of the powers conferred by sections 57(1), (2), (2A) and 129(4) of the Medicines Act 1968(**a**) and now vested in them(**b**), and of all the powers enabling them in that behalf, after consulting such organisations as appear to them to be representative of interests likely to be substantially affected by the following order in accordance with section 129(6) of the said Act and with the consent of the Treasury, hereby make the following order:—

Title and commencement

1. This order may be cited as the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1985 and shall come into operation on 1st January 1986.

Interpretation

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

"the Act" means the Medicines Act 1968;

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

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

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

(**b**) In the case of the Secretaries of State concerned with health in England and Wales by virtue of S.I. 1969/388, in the case of the Secretary of State concerned with agriculture in Wales by virtue of S.I. 1978/272 and in the case of the Northern Ireland Departments by virtue of section 40 of, and Schedule 5 to, the Northern Ireland Constitution Act 1973 (c. 36) and section 1(3) of, and paragraph 2(1)(*b*) of Schedule 1 to, the Northern Ireland Act 1974 (c. 28).

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

"dosage unit" means—

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

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

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

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

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

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

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

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

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

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

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

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

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

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

(**a**) S.I. 1971/1450.
(**b**) For the current relevant order *see* S.I. 1984/768.

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

Exemption for merchants in veterinary drugs

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

(*a*) that veterinary drug either—

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

(*b*) Details of premises used for the storage of any veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) above at a different postal address to that of premises used to sell by retail such drug shall be recorded in a register kept under sub-paragraph (*a*) above.

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

(9) Where a person who, whilst carrying on a qualifying business in Northern Ireland, makes an application in writing to the Department for his name to be

entered in the Department's register in respect of any premises on which any veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) above is to be sold or stored by him in the course of that qualifying business, the Department shall, subject to paragraphs (12) and (13) below, enter his name in the Department's register in respect of those premises.

(10)(*a*) Subject to paragraphs (14) and (16) below a person whose name is entered in the Society's register or the Department's register in respect of any premises shall, in order to retain his name on the register in respect of those premises in any year subsequent to the year in which his name is first entered in it, in the month of January in any such year make an application in writing to the Society or the Department (as the case may be) for his name to be retained in the Society's register or the Department's register (as the case may be) in respect of those premises.

(*b*) Subject to paragraphs (15) and (16) below a person whose name is removed from the Society's register or the Department's register 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 sub-paragraph (*a*) above or to pay the fee due in respect of the retention of his name in the register pursuant to paragraph (14) below may, in order to restore his name to the register in respect of those premises, make an application to the Society or the Department (as the case may be) for his name to be restored to the Society's register or the Department's register (as the case may be) in respect of those premises.

(11) There shall be paid to the Society or the Department—

(*a*) in respect of the entry in the Society's register or the Department's register (as the case may be) of the name of any person in respect of one or more premises on which any veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) above is to be sold or stored a fee of £125.00 for each such premises;

(*b*) in respect of the retention in the Society's register or the Department's register (as the case may be) of the name of any person in respect of one or more premises on which any veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) above is to be sold or stored a fee of £95.00 for each such premises;

(*c*) in respect of the restoration to the Society's register or the Department's register (as the case may be) of the name of any person in respect of one or more premises on which any veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) above is to be sold or stored a fee of £145.00 for each such premises.

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

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

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

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

(14) The Society or the Department shall refuse to retain in the Society's register or the Department's register (as the case may be) 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 (as the case may be) on or before 31st January in that year the fee specified in paragraph (11)(*b*) above for the retention of his name in the register.

(15) The Society or the Department shall refuse to restore to the Society's register or the Department's register (as the case may be) the name of any person in respect of any premises unless that person, having made proper application pursuant to paragraph (10)(*b*) above, has paid to the Society or the Department (as the case may be) the fee specified in paragraph (11)(*c*) above for the restoration of his name to the register.

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

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

(ii) the conditions under which any veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) above is sold by retail on that premises or under which it is stored on that premises (whether immediately prior to retail sale or not) are unsuitable for that purpose.

(*b*) In respect of any premises the Society or the Department may remove from the Society's register or the Department's register (as the case may be) the name of any person entered in it, at the request of that person.

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

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

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

(*a*) that veterinary drug either—

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

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

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

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

(*a*) for incorporation in animal feeding stuffs; and

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

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

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

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

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

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

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

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

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

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

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

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

Exemptions for merchants in horse wormers

5.—(1) The restrictions imposed by section 52 of the Act shall not apply during the period of 2 years from the date of coming into operation of this order 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 such as is described in paragraph (1)(*a*) above shall be sold by retail except—

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

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

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

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

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

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

(4)(*a*) In respect of any sale by retail of any veterinary drug such as is described in paragraph (1)(*a*) above the seller shall make a record of

the sale containing the particulars specified in sub-paragraph (*b*) below and shall keep such record for a period of two years from the date of the sale.

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

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

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

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

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

(*b*) Details of premises used for the storage of any veterinary drug such as is described in paragraph (1)(*a*) above at a different postal address to that of premises used to sell by retail such drug shall be recorded in a register kept under sub-paragraph (*a*) above.

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

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

(9)(*a*) Subject to paragraphs (13) and (15) below a person whose name is entered in the register kept by the Society or the Department under paragraph (6) above in respect of any premises shall, in order to retain his name on the register in respect of those premises in any year subsequent to the year in which his name was first entered in it, in the month of January in any such year make an application in writing to

the Society or the Department (as the case may be) for his name to be retained in that register in respect of those premises.

(*b*) Subject to paragraphs (14) and (15) below a person whose name is removed from the register kept by the Society or the Department under paragraph (6) 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 sub-paragraph (*a*) above or to pay the fee due in respect of the retention of his name in the register pursuant to paragraph (13) 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 (as the case may be) for his name to be restored to the register kept by the Society or the Department (as the case may be) under paragraph (6) in respect of those premises.

(10) There shall be paid to the Society or the Department—

(*a*) in respect of the entry in the register kept by the Society or the Department (as the case may be) under paragraph (6) above of the name of any person in respect of one or more premises on which any veterinary drug such as is described in paragraph (1)(*a*) above is to be sold or stored a fee of £30.00 for each such premises;

(*b*) in respect of the retention in the register kept by the Society or the Department (as the case may be) under paragraph (6) above of the name of any person in respect of one or more premises on which any veterinary drug such as is described in paragraph (1)(*a*) above is to be sold or stored a fee of £30.00 for each such premises;

(*c*) in respect of the restoration to the register kept by the Society or the Department (as the case may be) under paragraph (6) above of the name of any person in respect of one or more premises on which any veterinary drug such as is described in paragraph (1)(*a*) above is to be sold or stored a fee of £60.00 for each such premises;

except that no such fees 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 Society's register or the Department's register (as the case may be) in respect of those premises as being a person entitled to sell or store thereon, during the course of a qualifying business carried on by him, any veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) of Article 3.

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

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

(*b*) has given to the Society or the Department (as the case may be) an undertaking in writing that he will comply with the provisions of the Code of Practice for Saddlers Selling or Supplying Horse Wormers dated October 1985 and published by the Ministry of Agriculture, Fisheries and Food (being a code of practice relating to the sale or

supply of such veterinary drugs as are described in paragraph (1)(*a*) above).

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

(13) The Society or the Department shall refuse to retain in the register kept by the Society or the Department (as the case may be) under paragraph (6) above 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 (as the case may be) on or before 31st January in that year the fee specified in paragraph (10)(*b*) above for the retention of his name in the register.

(14) The Society or the Department shall refuse to restore to the register kept by the Society or the Department (as the case may be) under paragraph (6) above the name of any person in respect of any premises unless that person, having made proper application pursuant to paragraph (9)(*b*) above, has paid to the Society or the Department (as the case may be) the fee specified in paragraph (10)(*c*) above for the restoration of his name to the register.

(15)(*a*) The Society, with the approval of the Minister, or the Department, with the approval of the Department of Agriculture for Northern Ireland, may refuse to retain in or to restore to, or may remove from, the register kept by the Society or the Department (as the case may be) under paragraph (6) above the name of any person in respect of any premises if, in the opinion of the Society or the Department (as the case may be)—

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

(ii) the conditions under which any veterinary drug such as is described in paragraph (1)(*a*) above is sold by retail on that premises or under which it is stored on that premises (whether immediately prior to retail sale or not) are unsuitable for that purpose.

(*b*) In respect of any premises the Society or the Department may remove from the register kept by the Society or the Department (as the case may be) under paragraph (6) above the name of any person entered in it, at the request of that person.

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

(17) For the purposes of this Article—

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

(*b*) "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.

Exemptions for pharmacists

6.—(1) The restrictions imposed by section 52(*c*) of the Act on the supply of medicinal products shall not apply to the retail sale of a veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) of Article 3 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 on the supply of medicinal products shall not apply to the supply in circumstances corresponding to retail sale of a veterinary drug such as is described in paragraph (1)(*a*)(i) or (ii) of Article 4 by a pharmacist, or his agent, to the person to whom the pharmacist has, in accordance with the provisions of the said section 52, sold the drug by retail.

Exemption in cases involving another's default

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

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

(3) The restrictions imposed by section 52 of the Act shall not apply to the sale by retail, in compliance with the conditions set out in Article 5, of a veterinary drug by a person for the time being carrying on a qualifying business or a saddlery business (as defined in Article 5(17)), which drug that person, having exercised all due diligence, on reasonable grounds believes to be a veterinary drug such as is described in paragraph (1)(*a*) of Article 5, but which, due to the act or default of another person is not such a veterinary drug.

Revocation

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

Norman Fowler,
Secretary of State for Social Services.

25th November 1985.

George Younger,
Secretary of State for Scotland.

21st November 1985.

Nicholas Edwards,
Secretary of State for Wales.

25th November 1985.

In Witness whereof the Official Seal of the Minister of Agriculture, Fisheries and Food is hereunto affixed on 25th November 1985.

L.S.

Michael Jopling,
Minister of Agriculture, Fisheries and Food.

Sealed with the Official Seal of the Department of Health and Social Services for Northern Ireland this 22nd day of November 1985.

L.S.

M. N. Hayes,
Permanent Secretary.

Sealed with the Official Seal of the Department of Agriculture for Northern Ireland this 15th day of November 1985.

L.S.

W. H. Jack,
Permanent Secretary.

We consent,

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

14th November 1985.

SCHEDULE 1: Article 3

PART A

LICENCE OF RIGHT VETERINARY DRUGS

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
1. GROWTH PROMOTERS	Bacitracin Zinc		Incorporation in feed	
	Bambermycin		Incorporation in feed	
	Copper Salts		Incorporation in feed	
	Nitrovin		Incorporation in feed	
	Spiramycin		Incorporation in feed	
	Tylosin Phosphate		Incorporation in feed	For pigs
	Virginiamycin		Incorporation in feed	
2. COCCIDIO-STATS	Amprolium Hydrochloride			
	Clopidol			
	Decoquinate		Incorporation in feed	
	Diaveridine		Not to be incorporated in feed	
	Dinitolmide			
	Ethopabate			
	Methyl Benzoquate	1.75%	Incorporation in feed	When combined with not more than 20.6% of Clopidol
	Robenidine			
	Sulphaquinoxaline	12%	Incorporation in feed	When combined with not more than 20% of Amprolium Hydrochloride and 1% of Ethopabate
3. ANTI-BLACKHEAD PREP-ARATIONS	Acinitrazole		Not to be incorporated in feed	
	Aminonitrothiazole		Not to be incorporated in feed	
	Dimetridazole		Incorporated in feed	
	Nifursol			
4. SHEEP DIPS AND ECTO-PARASITICIDES	Allethrin			
	Amitraz			
	Benzuldazic Acid, Sodium Salt		External use only	
	Benzyl Benzoate			
	Bromocyclen		External use only	
	Bromophos			
	Bucarpolate			
	Butacarb			
	Carbaryl			
	Carbophenothion			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
4. SHEEP DIPS AND ETCO-PARASITICIDES (continued)	Chlorfenvinphos			
	Chlorpyrifos			
	Coal Tar Phenols		External use only	
	Coumaphos			
	Crotoxyphos			
	Cresol	4%		
	Cresylic Acid		External use only	
	Derris Resins			
	Diazinon			
	Dichlofenthion			
	Dichlorvos			
	Dicophane			
	Dioxathion			
	Dursban			
	Fenchlorphos			
	Fenitrothion			
	Gamma BHC			
	Iodofenphos			
	Lethane			
	Malathion			
	Phosalone			
	Pyractone			
	Pyrimithate			
	Rotenone			
5. ANTHEL-MINTICS	Bephenium and its salts			
	Bunamidine and its salts			
	Cyacetazide			
	Dichlorvos			
	Diethyl-carbamazine and its salts			
	Haloxon			
	Levamisole and its salts			
	Mebendazole			
	Metriphonate			
	Morantel and its salts			
	Naphthalophos			
	Niclosamide			
	Nitroxynil and its salts			
	Parbendazole			
	Phenothiazine			
	Piperazine Carbon Disulphide Complex			
	Pyrantel and its salts			
	Sodium			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
5. ANTHELMINTICS (continued)	Glycarsamate			
	Tetramisole and its salts			
	Thiabendazole			
	Thiophanate			
6. MILK FEVER PREPARATIONS	Calcium Borogluconate		Parenteral use only	Products may or may not contain all or any of the following substances:— Dextrose Magnesium Phosphorus
7. WARBLE FLY DRESSINGS	Crufomate			
	Famphur			
	Fenchlorphos			
	Fenthion			
	Metriphonate			
	Prolate			
8. LIVER FLUKE REMEDIES	Brotianide			
	Diamphenethide			
	Hexachloroethane			
	Hexachlorophane			
	Nitroxynil and its salts			
	Oxyclozanide			
	Rafoxanide			
	Tribromsalan			
9. SHEEP AND CATTLE CLOSTRIDIAL VACCINES AND ANTISERA	Black Disease Antisera and Vaccines			
	Blackleg (Blackquarter) Vaccines and Antisera			
	Braxy Vaccines and Antisera			
	Enterotoxaemia Vaccines and Antisera			
	Lamb Dysentry Antisera and Vaccines			
	Pulpy Kidney Vaccines and Antisera			
	Struck Vaccines and Antisera			
	Tetanus Toxoids			
	Combinations of two or more of Braxy, Blackleg (Blackquarter),			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
9. SHEEP AND CATTLE CLOSTRIDIAL VACCINES AND ANTISERA (continued)	Lamb Dysentry, Pulpy Kidney, Enterotoxaemia, Struck, Tetanus, Black Disease and Pasteurella Vaccines			
10. POULTRY VACCINES	Avian Encephalomyelitis Vaccines (Living and Inactivated)			
	Duck Hepatitis Vaccines (Living)			
	Fowl Pox Vaccines			
	Fowl Typhoid Vaccines (Salmonella gallinarum)			
	Infectious Bronchitis Vaccines (Living and Inactivated)			
	Infectious Bursal Disease Vaccines			
	Infectious Laryngotracheitis Vaccines (Living)			
	Marek's Disease Vaccines			
	Newcastle Disease Vaccine (Living and Inactivated)			
	Combinations of Newcastle Disease Vaccines and Avian Encephalomyelitis Vaccines			
	Combinations of Newcastle Disease Vaccines with Infectious Bronchitis Vaccines			
	Pasteurella Vaccines			
11. ERYSIPELAS VACCINES	Avian Erysipelas Vaccines			
	Swine Erysipelas Vaccines			
12. SALMONELLA AND E. COLI VACCINES	E. Coli Vaccines (Killed)			
	Salmonella Vaccines (Killed)			
	E. Coli and Salmonella Sero Vaccines			
13. OTHER SHEEP AND CATTLE VACCINES	Foot Rot Vaccines			
	Louping I11 Vaccines (Killed)			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
13. OTHER SHEEP AND CATTLE VACCINES (continued)	Ovine Enzootic Abortion Vaccines			
	Pasteurella Vaccines (Killed)			
	Pneumonia Combined Vaccines (Pasteurella)			
14. MISCEL-LANEOUS VACCINES	Botulism Vaccines (Mink)			
	Combinations of E. Coli, Salmonella and Pasteurella Vaccines (Killed)			
	Pigeon Pox Vaccines (Living)			
15. SULPHANILA-MIDE SURFACE WOUND DRESSINGS	Sulphanilamide	5%	Powdered surface wound dressing	Use on farm animals only
16. LOCAL ANAESTHETICS	Products containing not more than 5% of Procaine Hydrochloride, 2% Lignocaine, or 2% Lignocaine Hydrochloride with or without not more than 0.002% of Adrenaline, Adrenaline Acid Tartrate or Noradrenaline		Parenteral use only	
17. OTHERS	Ammonia Solution Conc.	4%		
	Broxyquinoline			
	Butafosfan			
	Butyl Amino Benzoate			
	Clioquinol			Treatment of enteritis in livestock
	Cobalt Carbonate			For use only in anthelmintics
	Cobalt Oxide			For use only in combination with anthelmintics and in ruminal pellets
	Copper its inorganic salts and organic preparations		Non-parenteral use only	
	Creosote		Internal use	
	Dextrose		Parenteral use only	
	Dill, Oil of			

Group/Class	Substance	Maximum strength or concentration	Pharmaceutical form or route of administration	Other restrictions
17. OTHERS (continued)	Ethyl M-amino-benzoate			For use as an anaesthetic and tranquillizer in fish, frogs and other cold blooded organisms
	Halquinol	5%		
	Iron organic complexes with or without Vitamin B_{12}			
	Magnesium salts		(1) Subcutaneous injection	
			(2) Slow intravenous injection	(2) When combined with any of Calcium Borogluconate, Phosphorus or Dextrose
	Menandione Sodium Bisulphite			
	Menandione Dimethyl Pyrimidinol Bisulphite			
	Phenol			
	Poloxalene			
	Turpentine	40%	Internal use	
	Vitamin A			
	Vitamin B group			Excluding:— (1) preparations recommended for administation by intravenous route and (2) those recommended for use for conditions requiring a veterinary diagnosis
	Vitamin D_2			
	Vitamin D_3			
	Vitamin E		Parenteral use only	

SCHEDULE 1

Article 3

PART B

VETERINARY DRUGS

Product Licence No.	Name of Product*
1. Growth Promoters	
PL 4131/4000	Advantage with Romensin[R]
PL 3405/4019	*Avoparcin 50 Premix*
PL 0095/4026	Avotan 50
PL 0095/4028	Avotan 50c Avoparcin
	Avotan 100
PL 0095/4036	Avotan Blockmix
	Avotan Super
PL 0095/4039	Avotan Farm Mix
PL 0010/4038	Bayo-n-ox 10% Premix
PL 0002/4020	Eskalin 20
PL 3832/4020	
PL 3832/4031	Eskalin 100
PL 0002/4045	Eskalin 500
PL 3832/4017	
PL 0002/4055	Eskalin S-400
PL 3832/4021	
PL 4594/4001	FPL 40 "ABCHEM"
PL 0029/4102	Fedan 10% Premix
PL 0086/4137	Flavomycin 50
PL 5811/4001	*Intagen Premix*
PL 3405/4022	Monensin-100 Ruminant
PL 3405/4016	Nitrovin
PL 3405/4018	Nitrovin - 20
PL 2592/4075	Nitrozone 25
PL 4869/4000	Panazone 250-Nitrovin
PL 0777/4002	
PL 0095/4007	Payzone 50 MA Nitrovin Milk Replacer Additive
PL 4188/4008	Pentazone 250
PL 0006/4052	Romensin (Monensin Sodium) Premix
PL 2969/4006	Rumevite with Romensin
PL 0012/4170	SPIRA 200
PL 0006/4055	Tylamix Premix 100 g/kg
PL 0006/4062	Tylamix Premix 250 g/kg
PL 3405/4007	Tylosin 100 Premix
PL 4594/4002	Tylosin 250 "ABCHEM" Premix
PL 3405/4015	ZB-100
PL 0109/4001	Zinc Bacitracin Premix
2. Coccidiostats	
PL 0025/4035	Arpocox
PL 0031/4011	Avatec Premix
PL 3405/4017	Clopidol

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

Items shown in italics did not appear in the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1984 as amended.

Product Licence No.	Name of Product
2. Coccidiostats (continued)	
PL 3405/4025	*Clopidol 250*
PL 0109/4000	Dinormix SR 25
PL 4869/4005 *PL 0777/4001*	D.O.T.
PL 0109/4002	DOT (dinitolmide)
PL 0006/4047	Elancoban Premix
PL 0621/4015	Lerbek
PL 3405/4006	Monensin 200
PL 3405/4021	Monensin 100 Poultry
PL 4594/4000	Monensin 200 "ABCHEM" Premix
PL 0006/4061	Monteban 100 Premix
PL 0025/4019	Nicrazin (Premix)
PL 0086/4135	Sacox 60 Premix
PL 1598/4036	Salcostat
PL 1598/4032	Salcostat (DOT) Premix 12.5%
PL 1598/4033	Salcostat (DOT) Premix 25%
PL 0086/4117	Stenorol
3. Anti-Blackhead Preparations	
PL 3405/4009	Dazole Premix
PL 4869/4003 *PL 0777/4003*	Dimetridazole BP (Vet.)
PL 3636/4001	Dimetridazole - PML Turkeys
PL 0012/4176	'Emtryl' Premix
PL 0012/4174	'Emtryl' Pure
PL 0012/4175	'Emtryl' Soluble
PL 4188/4014	*Unizole T for Poultry*
PL 4188/4013	*Unizole T Soluble for Poultry*
4. Sheep Dips and Ectoparasiticides	
PL 1300/4010	Barricade
PL 0676/4089	Battles Improved Organo-Phosphorus Single-Dipping Fluid Dip
PL 0676/4087	Battles Organo-Phosphorus Single-Dipping Fluid Dip
PL 1300/4011	C Tag 97 Fly Tag/Flectron Fly Tag
PL 1300/4004	Ciodrin Insecticide
PL 0003/4140 *PL 5869/4004*	Coopers Fly and Scab Dip 200
PL 5869/4005	*Coopers Powerpack Summer Dip*
PL 5869/4002	*Coopers Powerpack Winter Dip*
PL 0003/4149	*Coopers Scab Approved Dip (Border Type)*
PL 0003/4150 *PL 5869/4007*	*Coopers Spoton Insecticide*
PL 0003/4124 *PL 5869/4003*	*Coopers Summer Dip 400*
PL 0003/4116 *PL 5869/4006*	*Coopers Winter Dip 200*
PL 4149/4001	Deodorised Malathion Premium Grade
PL 1476/4018	Deosan Dysect
PL 1476/4026	Deosan Flectron
PL 0829/4127	Dermol

Product Licence No.	Name of Product
4. Sheep Dips and Ectoparasiticides (continued)	
PL 1978/4001	Ectoral Tablets No. 1, 2 and 3
PL 4436/4005	Fly Dip
PL 1826/4004	Lice Tick and Mange Dressing (LTM)
PL 2428/4018	*Malacide*
PL 1826/4001	Northern Fly Dip
PL 0025/4044	Ovidip Scab Approved Sheep Dip
PL 2428/4018	*Pharmacide*
PL 0038/4068	Porect
PL 0086/4138	Prodip
PL 5656/4000	*Ridect Fly Tags*
PL 2100/4034	Rodgers No. 10 Neu-Fly and Tick Dip
PL 1447/4106	*Ryposect*
PL 0003/4113	Stomoxin
PL 0003/4148 *PL 5869/4009*	Stomoxin Fly Tags
PL 5869/4010	*Stomoxin Liquid Concentrate*
PL 1300/4005	Supona Sheep Dip
PL 4436/4001	Taktic
PL 1345/4040	Taskill
PL 0086/4140	Tirade Fly Tags
PL 1728/4070	Topclip Parasol
PL 1826/4025	Viper Dip
PL 1447/4052	Young's 200 Liquid Tick Dip
PL 1447/4096	Young's Cypor
PL 1447/4015	Young's Powder Fly Dip
PL 1447/4058	Young's Scab Approved 200 Liquid Tick Dip
PL 1447/4085	Young's Scab Approved Diazinon Winter Dip
PL 1447/4070	Young's Scab Approved Dursban Winter Dip Young's Dursban Winter Dip
PL 1447/4013	Young's Scab Approved Ectomort Summer Dip
PL 1447/4073	Young's Scab Approved Iodofenphos Winter Dip Young's Iodofenphos Winter Dip
PL 1447/4055	Young's Scab Approved Killtick Liquid Tick Dip Young's Killtick Liquid Tick Dip
PL 1447/4056	Young's Scab Approved Powder Fly Dip Young's Powder Fly Dip
PL 1447/4080	*Young's Scab Approved Summer Dip*
PL 1447/4056	Young's Scab Approved Summer Mycotic Dip Young's Summer Mycotic Dip
PL 1447/4060	Young's Sheep Blowfly Spray
PL 1447/4083	Young's SP Fly Spray
PL 1447/4015	Young's Summer Mycotic Dip
PL 3893/4069	Zeprox
5. Anthelmintics	
PL 0029/4103	'Amatron' Cattle Drench
PL 0029/4105	'Amatron' Sheep Drench
PL 1447/4092	Anthelpor
PL 4318/4003	Ashmintic Drench
PL 4318/4013	Ashmintic Injection
PL 0010/4063	*Bayverm Granules 10%*
PL 0010/4054	Bayverm L.V. Paste

Product Licence No.	Name of Product
5. Anthelmintics (continued)	
PL 0010/4058	Bayverm Pellets 1.9%
PL 0010/4049	Bayverm Premix 0.6%
PL 0010/4050	Bayverm Premix 2.4%
PL 0010/4062	*Bayverm Roundwormer*
PL 0010/4065	*Bayverm SC 10% Suspension Worm Drench*
PL 0010/4064	*Bayverm SC 2.5% Suspension Worm Drench*
PL 0010/4047	Bayverm Suspension 2.5%
PL 0010/4048	Bayverm Suspension 10%
PL 3974/4026	*Cevasol C Worm Drench*
PL 3974/4021	Cevasol Injection
PL 3974/4020	Cevasol Worm Drench
PL 0095/4040	Cyverm 11.5% Gel
PL 0095/4038	Cyverm Levamisole 3.2% Drench
PL 0095/4037	Cyverm Levamisole 7.5% Injection
PL 1861/4055	Day's Worm Drench
PL 0010/4046	Droncit
PL 1596/4071	Duphamisole 7.5% Oral
PL 5151/4001	*Equidin Paste*
PL 3832/4012	Equitac
PL 0829/4044	Equivurm Plus
PL 0829/4058	Equivurm Plus Paste
PL 0025/4027	Equizole Pony Paste
PL 0025/4042	Eqvalan Paste for Horses
PL 0829/4131	Flubenol Pellets
PL 0829/4120	Flubenol Premix
PL 0010/4055	Flukombin
PL 3763/4000	Gapex
PL 0002/4004 PL 3832/4038	Helmatac In-feed Wormer
PL 2592/4076	Helminate Sow Wormer Pellets
PL 0025/4041	Ivomec Drench
PL 0025/4040	Ivomec Injection
PL 0025/4043	Ivomec Injection for Pigs
PL 2000/4054	Levacide - C Worm Drench
PL 2000/4049	Levacide Injection
PL 2000/4060	*Levacide SC Worm Drench*
PL 2000/4050	Levacide Worm Drench
PL 3832/4066	Loditac 3% Wormer Pellets
PL 3832/4069	Loditac 200
PL 3832/4070	Loditac 20
PL 0829/4126	Mebatreat
PL 0829/4113	Mebenvet (1.2%)
PL 0829/4123	Mebenvet (5%)
PL 0829/4114	Multispec
PL 0012/4003	Nemafax Drench
PL 0012/4149	Nemafax 5
PL 0012/4149	Nemafax 14
PL 0012/4151	Nemafax Cattle, Sheep and Goat Wormer Pellets
PL 0012/4151	Nemafax Sow
PL 0012/4153	Nemafax Wettable Powder
PL 0029/4101	Nemicide Cattle Drench
PL 0029/4100	Nilverm C. Small Dose Cattle Drench Nilverm C. Cattle Drench

Product Licence No.	Name of Product
5. Anthelmintics (continued)	
PL 0029/4101	Nilverm Cattle Special
PL 0029/4114	Nilverm Plus Drench
PL 0029/4118	Nilverm Super
PL 0029/4098	Nilzan C
PL 0029/4115	'Nilzan' Drench Plus
PL 0029/4117	'Nilzan' Drench Super
PL 0829/4114	Ovitelmin
PL 0829/4162	Ovitelmin Bolus
PL 0829/4163	Ovitelmin SC
PL 0086/4121	Panacur 1.5% Pellets
PL 0086/4105	Panacur 2.5% Sheep Wormer
PL 0086/4105	Panacur 2.5% Suspension
PL 0086/4110	Panacur 4% Paste
PL 0086/4106	Panacur 10% Suspension
PL 0086/4107	Panacur 22% Granules
PL 0086/4119	Panacur Paste
PL 0086/4136	Panacur SC Cattle Wormer
PL 0086/4130	Panacur SC Sheep Wormer
PL 0057/4075	Paratect Sustained Release Bolus
PL 0025/4031	Porcam
PL 3832/4073	*Powacide*
PL 0025/4038	Ranizole Paste
PL 5789/4002	Reid's Worm Drench
PL 0829/4150	Ripercol
PL 0829/4133	Ripercol 3.2% Oral
PL 0829/4140	Ripercol 5% Injectable Solution
PL 0829/4151	Ripercol 7.5% Injectable
PL 0829/4132	Ripercol 15% Injectable Solution
PL 0829/4165	*Ripercol SC*
PL 0086/4115	Rumevite Wormablok with Panacur for Cattle
PL 0086/4114	Rumevite Wormablok with Panacur for Sheep
PL 1447/4094	Rycovet Horse and Pony Wormer
PL 0029/4099	Spectril
PL 0057/4060	Strongid-P (Granules)
PL 0057/4062	Strongid-P Paste
PL 0057/4063	Suiminth (Morantel Tartrate)
PL 0286/4032	Synanthic
PL 0286/4034	Synanthic DC
PL 0286/4039	Synanthic Horse Paste
PL 0286/4035	Synanthic Horse Pellets
PL 0286/4040	*Synanthic I/R*
PL 0003/4127	Systamex Paste 18.5% Cattle and Horse Wormer
PL 0003/4127	Systamex Paste 18.5% Horse and Pony Wormer
PL 0003/4121	Systamex 906 Concentrated Cattle Wormer
PL 5869/4014	*Systamex SC*
PL 0003/4112	Systamex Worm Drench for Cattle and Sheep
PL 0829/4044	Telmin Granules
PL 0829/4112	Telmin KH
PL 0829/4114	Telmin Liquid
PL 4462/4002	Tetramisole Hydrochloride BP (Vet)
PL 0025/4024	Thibenzole Paste

Product Licence No.	Name of Product
5. Anthelmintics (continued)	
PL 0002/4061 PL 3832/4022	Valbazen 2.5% Total Spectrum Wormer
PL 0002/4062 PL 3832/4023	Valbazen 10% Total Spectrum Wormer
PL 3832/4015	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 0012/4172	Vermadax
PL 2676/4120	*Vermisole Injection*
PL 0086/4139	Wormex
PL 1447/4091	Young's Anthelpor 20
PL 1447/4075	Young's Anthelworm Rycovet Widespec
PL 1447/4090	Young's Anthelworm Feed Pellets
PL 1447/4076	Young's Anthelworm L
PL 1447/4100	Young's Endozal
6. Milk Fever Preparations	
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 2428/4024	Calcium Borogluconate Injection 25% with Phosphorus Magnesium and Dextrose
PL 2428/4028	Calcium 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 1345/4007	TVL Calcium Borogluconate "Borocal"
7. Warble Fly Dressings	
PL 0003/4115	Cooper Warble Fly Liquid
PL 0095/4024	Cynamid Systemic Warble Fly Dressing
PL 0829/4127	Dermol
PL 0010/4045	Neguvon Spot-on
PL 0038/4062	Orbisect Warble Fly, Louse and Mange Liquid for Cattle
PL 4436/4000	Warbexol - Ready To Use Systemic Warble Fly Dressing
PL 1447/4074	Young's New Poron
PL 1447/4077	Young's Poron 20

Product Licence No.	Name of Product
8. Liver Fluke Remedies	
PL 0010/4031	Dirian
PL 1728/4065	Fasinex 5%
PL 1728/4067	Fasinex 10%
PL 0025/4036	Flukanide
PL 3832/4073	*Powacide*
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 New Flukol *Young's Flukol*
9. Sheep and Cattle Clostridial Vaccines and Antisera	
PL 0086/4132	Heptavac P
PL 1345/4063	Nilvax
PL 1345/4069	Nilvax under 30 kg
PL 0086/4129	Ovivac P
PL 1345/4062	Tasvax 8
PL 1728/4066	Topclip Ewe Vaccine 8 in 1
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 0002/4053 PL 3832/4033	Bronchimune IB Vaccine Infectious Bronchitis Vaccine (Living) Strain HL Massachusetts type (Bronchimune)
PL 0002/4034 PL 3832/4041	Combimune
PL 1598/4029	Combined ND (HB1) and IB (Massachusetts MM) Vaccine (Living)
PL 3359/4004	Delvax IB H52
PL 3359/4003	Delvax IB H120
PL 3359/4001	Delvax Marek THV Freeze-dried
PL 3359/4035	Delvax ND Hitchner
PL 3359/4005	Delvax ND HB1
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 4978/4004 PL 5654/4000	Iblin
PL 5654/4013	Iblin Live
PL 2592/4037	Ibvax
PL 0002/4002 PL 3832/4056	IB Vaccine (Living) Massachusetts H52 Strain
PL 0002/4003 PL 3832/4036	IB Vaccine (Living) Massachusetts H120 Strain
PL 1708/4135	Inactivated ND Vaccine (oil emulsion) Newcavac Nobilis

Product Licence No.	Name of Product
10. Poultry Vaccines (continued)	
PL 1598/4056	Infectious Laryngotracheitis Vaccine (LT-VAC)
PL 2592/4074	Ivamarek Marek's Disease Vaccine
PL 2592/4044	Lentogen HB1
PL 5654/4018	Marek's
PL 0002/4001	Marek's Disease Vaccine (Living) THV (Marimune)
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 1708/4169	Marexine THV/CA
PL 4978/4005	Maternalin
PL 5654/4001	
PL 5654/4012	Maternalin Plus
PL 4978/4003	Myxilin
PL 5654/4002	
PL 5654/4021	*Myxilin Live*
PL 3318/4000	ND Vaccine (Inactivated) Oil Emulsion
PL 4978/4002	Newcadin Day Old
PL 5654/4004	
PL 4978/4001	*Newcadin*
PL 5654/4006	*Newcadin 25*
PL 4978/4000	Newcadin L
PL 5654/4008	
PL 5654/4020	*Newcadin Live B-1*
PL 3317/4087	Newcastle Disease Vaccine K2C (Inactivated)
PL 2592/4033	Newcastle Disease Vaccine (Inactivated) Oil Emulsion (Layer Plus)
PL 3832/4057	Newcastle Disease Vaccine (Living) Hitchner B1 Strain
PL 1708/4150	Newcavac + EDS '76 Vaccine
PL 1708/4143	Nobi-Vac Egg Drop Syndrome '76 Vaccine BC14 (Inactivated)
PL 1708/4155	Nobi-Vac Gumboro Inactivated
PL 1596/4034	Poulvac AE
PL 1596/4040	Poulvac EDS
PL 1596/4029	Poulvac IB Vaccine H52 (Living)
PL 1596/4030	Poulvac IB Vaccine H120 (Living)
PL 1596/4045	Poulvac Marek HVT Vaccine
PL 1596/4025	Poulvac Marek THV
PL 1596/4026	Poulvac ND Vaccine (Living) HB 1
PL 0002/4005	Tremimune
PL 3832/4024	
PL 5654/4019	*Ultravac*
11. Erysipelas Vaccines	
PL 1531/4012	Ferrovac Ery Vaccine
PL 1596/4078	Suvaxyn Erysipelas Vaccine
PL 3317/4110	Swine Erysipelas Vaccine (Inactivated)
PL 1345/4004	Swine Erysipelas Vaccine, Inactivated (Oil Adjuvant) Erysivax

Product Licence No.	Name of Product
12. Salmonella and E. coli Vaccines	
PL 0086/4134	Coliovac
PL 3832/4009	Ecopig
PL 0003/4110	Gletvax K88-Porcine E. coli Vaccine (Polyvalent)
PL 0003/4110	Gletvax-Porcine E. coli Vaccine (Polyvalent)
PL 0003/4110	Gletvax-Porcine E. coli Vaccine (Polyvalent) + K88
PL 0086/4113	Porcovac AT
PL 3832/4004	Scourguard I
PL 1754/4002	Sow Intagen O/I Injectable
PL 1596/4076	*Suvaxyn E coli*
13. Other Sheep and Cattle Vaccines	
PL 1345/4070	Footvax
PL 0003/4135	Ovine Enzootic Abortion (Improved) Vaccine
PL 0086/4133	Ovipast
14. Miscellaneous Vaccines	
PL 1708/4152	Nobi-Vac L.T. K88
15. Local Anaesthetics	
PL 3317/4049	Lignavet Plus Injection
PL 2324/4074	Lignocaine Anaesthetic Injection
PL 2000/4029	Lignocaine and Adrenalin Injection
PL 2428/4021	Pharmcaine
PL 1599/4005	Ruby Freezaject
16. Others	
PL 4318/4002	Ash-fer 100
PL 2428/4026	Bactasorb Tablets
PL 0002/4043 PL 3832/4034	Bloat Guard
PL 0002/4054 *PL 3832/4064*	Bloat Guard Drench
PL 3832/4065	Bloat Guard Liquid
PL 3514/4002	Boar Mate
PL 2428/4023	Brodin
PL 1754/4003	Calf Intagen Premix
PL 2545/4009	Codifer 10
PL 0676/4091	Colostrene-Watery Mouth Drench for Young Lambs
PL 3317/4010	Copavet
PL 2987/4003	Copper (Cupric) Carbonate
PL 0038/4088	Copporal 2 g
PL 0038/4089	Copporal 4 g
PL 0038/4090	Copporal 24 g
PL 0038/4078	Copprite 2 g
PL 0038/4084	Copprite 4 g
PL 0038/4087	Copprite 24 g
PL 1345/4012	Cujec
PL 2987/4002	Cupric Oxide
PL 2987/4001	Cuprous Chloride

Product Licence No.	Name of Product
16. Others (continued)	
PL 3656/4012	Dio-Iron
PL 1596/4031	Ducrofer
PL 0113/4005	Fisons Multivitamin Injection
PL 0113/4006	Fisons Vitamin A, D & E Injection
PL 0113/4007	Gleptosil
PL 5806/4000	Golden Hoof Zinc Sulphate
PL 2324/4079	Glucose Saline Injection
PL 1754/4009	HI-FAT Baby Calf Food 'Intagen'
PL 1754/4000	Intagen Premix
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 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 0676/4090	Orfoids-Capsules for Orf
PL 1345/4051	Permasel-C
PL 1345/4052	Permasel-S
PL 2428/4017	*Pharmamag 25*
PL 2428/4007	Pharmavit AD_3E
PL 0829/4133	Ripercol 3.2% Oral
PL 0829/4140	Ripercol 5% Injectable Solution
PL 0829/4132	Ripercol 15% Injectable
PL 1011/4001	Roscofer 10% Vet
PL 1011/4000	Roscoral Vet
PL 3317/4077	Sildex
PL 5811/4000	*Sow Intagen O/I*
PL 1754/4002	Sow Intagen O/I
PL 1599/4004	Swipoul
PL 0829/4117	Tendex
PL 5923/4002	Tracerglass C
PL 5923/4001	*Tracerglass L*
PL 5923/4000	Tracerglass S
PL 3317/4047	Vetrivite Plus
PL 2428/4007	Vitamin ADE Solution
PL 2969/4005	Vituramol 60 with Romensin
PL 1447/4036	Young's Swaycop

SCHEDULE 2

Article 4

PART A

LICENCE OF RIGHT VETERINARY DRUGS

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

Article 4

SCHEDULE 2

PART B
VETERINARY DRUGS

Product Licence No.	Name of Product*
1. Growth Promoters	
PL 3405/4019	*Avoparcin 50 Premix*
PL 0095/4026	Avotan 50
PL 0095/4028	Avotan 50c Avoparcin
PL 0095/4036	Avotan 100 Avotan Block Mix Avotan Super
PL 0095/4039	Avotan Farm Mix
PL 0010/4043	Bayo-n-ox 10% Premix
PL 0002/4020 PL 3832/4020	Eskalin 20
PL 3832/4031	Eskalin 100
PL 0002/4045 PL 3832/4017	Eskalin 500
PL 0002/4055 PL 3832/4021	Eskalin S-400
PL 4594/4001	FPL 40 'ABCHEM'
PL 0029/4102	Fedan 10% Premix
PL 0086/4137	Flavomycin 50
PL 5811/4001	*Intagen Premix*
PL 3405/4022	Monesin-100 Ruminant
PL 3405/4016	Nitrovin
PL 3405/4018	Nitrovin - 20
PL 2592/4075	Nitrozone 25
PL 4869/4000 *PL 0777/4002*	Panazone 250 Nitrovin
PL 0095/4007	Payzone 50 MA Nitrovin Milk Replacer Additive
PL 4188/4008	Pentazone 250
PL 2969/4006	Rumevite with Romensin
PL 0006/4052	Romensin (Monensin Sodium) Premix
PL 0012/4170	SPIRA 200
PL 0006/4055	Tylamix Premix 100 g/kg
PL 0006/4062	Tylamix Premix 250 g/kg
PL 3405/4007	Tylosin 100 Premix
PL 4594/4002	Tylosin 250 'ABCHEM' Premix
PL 0109/4001	Zinc Bacitracin Premix
PL 3405/4015	ZB-100
2. Coccidiostats	
PL 0025/4035	Arpocox
PL 0031/4011	Avatec Premix
PL 3405/4017	Clopidol
PL 3405/4025	*Clopidol 250*
PL 0109/4000	Dinormix SR 25

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

Items shown in italics did not appear in the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1984 as amended.

Product Licence No.	Name of Product
2. Coccidiostats (continued)	
PL 4869/4005 *PL 0777/4001*	D.O.T.
PL 0109/4002	DOT (dinitolmide)
PL 0006/4047	Elancoban Premix
PL 3405/4006	Elancoban Premix Monensin 200
PL 0621/4015	Lerbek
PL 3405/4022	Monensin 100 Ruminant
PL 4594/4000	Monensin 200 "ABCHEM" Premix
PL 0006/4061	Monteban 100 Premix
PL 0025/4019	Nicrazin (Premix)
PL 0086/4135	Sacox 60 Premix
PL 1598/4036	Salcostat
PL 1598/4032	Salcostat (DOT) Premix 12.5%
PL 1598/4033	Salcostat (DOT) Premix 25%
PL 0086/4117	Stenorol
3. Anti-Blackhead Preparations	
PL 4869/4003 *PL 0777/4003*	Dimetridazole BP (vet)
PL 3636/4001	Dimetridazole - PML Turkeys
PL 3405/4009	Dazole Premix
PL 0012/4176	'Emtryl' Premix
PL 0012/4174	'Emtryl' Pure
PL 0012/4175	'Emtryl' Soluble
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 0829/4131	Flubenol Pellets
PL 0829/4120	Flubenol Premix
PL 0002/4004 PL 3832/4038	Helmatac In-Feed Wormer
PL 3832/4069	Loditac 200
PL 3832/4070	Loditac 20
PL 0829/4113	Mebenvet (1.2%)
PL 0829/4123	Mebenvet (5%)
PL 0012/4149	Nemafax 5
PL 0012/4149	Nemafax 14
PL 0012/4153	Nemafax Wettable Powder
5. Others	
PL 0002/4043 PL 3832/4034	Bloat Guard
PL 1754/4003	Calf Intagen Premix
PL 1754/4009	HI-FAT Baby Calf Food 'Intagen'
PL 1754/4000	Intagen Premix
PL 4127/4000	Micro Anti-Bloat Premix
PL 5811/4000	*Sow Intagen O/I*
PL 1754/4002	Sow Intagen O/I
PL 2696/4005	Vituramol 60 with Romensin

Article 4

SCHEDULE 3

PART A

LICENCE OF RIGHT VETERINARY DRUGS

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

Article 4

SCHEDULE 3

PART B

VETERINARY DRUGS

Product Licence No.	Name of Product*
PL 0006/4053	Apralan Soluble Powder
PL 0006/4057	Apralan 20 Premix
PL 0006/4058	Apralan 100 Premix
PL 3405/4010	Dazole Prescription Premix
PL 3636/4002	Dimetridazole - POM Swine and Turkeys
PL 0034/4031	Dynamutilin 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 0002/4071 PL 3832/4019	'Eskalin' 500 POM for laying and breeding hens
PL 1654/4012	Fortracin BMD^{R}
PL 3317/4031	Framomycin 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

*Alternative product names used by specially authorised persons are not shown.
Items shown in italics did not appear in the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1984 as amended.

Product Licence No.	Name of Product
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%
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 0057/4031	Terramycin 5% Feed Supplement
PL 0057/4065	Terramycin 20% Feed Supplement
PL 0057/4061	Terramycin Concentrate 20%
PL 0003/4105	Tribrissen 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 5

HORSE WORMERS

Product Licence No.	Name of Product*
PL 1732/4059	Astrobot 5
PL 1732/4060	Astrobot 10
PL 0010/4063	*Bayverm Granules 10%*
PL 0010/4054	Bayverm LV 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 0025/4004	Equizole Feed Pellets
PL 0025/4027	Equizole Pony Paste
PL 0025/4005	Equizole Powder
PL 0025/4042	Eqvalan Paste for Horses
PL 0844/4055	Multiwurma (Horses)

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

Items shown in italics did not appear in the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1984 as amended.

Product Licence No.	Name of Product
PL 0086/4107	Panacur 22% Granules
PL 0086/4119	Panacur Paste
PL 1599/4001	Ruby Horse Wormer
PL 1447/4094	Rycovet Horse and Pony Wormer Paste
PL 0057/4060	Strongid P (Granules)
PL 0057/4062	Strongid P Paste
PL 0286/4039	Synanthic Horse Paste
PL 0286/4035	Synanthic Horse Pellets
PL 0003/4127	Systamex Paste 18.5% Horse and Pony Wormer
PL 0829/4058	Telmin
PL 0829/4044	Telmin Granules

Article 8

SCHEDULE 5

REVOCATION

Orders revoked	References
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1984	S.I. 1984/1861
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) Order 1985	S.I. 1985/310
The Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) (Amendment) (No. 2) Order 1985	S.I. 1985/857

EXPLANATORY NOTE

(This Note is not part of the Order.)

This order re-enacts, with amendments, the provisions of the Medicines (Exemptions from Restrictions on the Retail Sale or Supply of Veterinary Drugs) Order 1984 ("the 1984 Order") and its amending instruments, which are revoked.

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

Article 3 of the order continues to exempt from the restrictions imposed by section 52 the sale or supply of any veterinary drug described in Article 3(1)(*a*) of the order by (1) the product licence holder, (2) a specially authorised person (as defined in the order) or (3) a person carrying on a qualifying business (that is to say, a business involving, at least in part, the retail sale of agricultural requisites) provided, in each case, that the relevant conditions contained in Article 3 are complied with.

These conditions include a requirement that a person who, in the course of carrying on a qualifying business, wishes to sell any veterinary drug described in Article 3(1)(*a*) must have his name and details of all relevant premises entered in a register of merchants in veterinary drugs ("the Register"); details of premises used for the storage only of such drugs at a different postal address to that of premises used for selling are to be recorded separately in the Register. The Register is kept by the Pharmaceutical Society of Great Britain ("the Society") and the Department of Health and Social Services for Northern Ireland ("the Department") (Article 3(6) and (7)).

A fee of £125 is to be paid to the Society or the Department for the initial entry in the Register of the name of a person in respect of each premises and a further fee of £95 is payable annually for the retention of a person's name in the Register in respect of each premises (Article 3(11)(*a*) and (*b*)). A person who fails to make proper application for the retention of his name in the Register within the time allowed shall have his name removed from the Register but is entitled to apply for the restoration of his name to the Register upon payment of a fee of £145 in respect of each premises (Article 3(11)(*c*)).

A person's name will not be entered in the Register unless he has paid the prescribed fee and has given an undertaking that he will comply with the provisions of the Code of Practice for Merchants Selling or Supplying Veterinary Drugs dated 30th October 1984. This Code is published by the Ministry of Agriculture, Fisheries and Food (Article 3(12)).

In addition, the Society (with the prior approval of the Minister of Agriculture, Fisheries and Food) or the Department (with the prior approval of the Department of Agriculture for Northern Ireland) may refuse to retain in or to restore to, or may remove from, its Register, the name of any person in respect of any premises if that person has failed to observe any of the provisions of the Code of Practice referred to above or, if the conditions under which any veterinary

drug prescribed in Article 3(1)(*a*) is sold by retail on the premises, or under which it is stored on those or separate premises prior to retail sale, are unsuitable for that purpose. There is provision, too, for a person to have his name removed from the Register in respect of any premises upon request (Article 3(16)).

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

Article 5 of the order continues to exempt from the restrictions imposed by section 52 the sale by retail of any veterinary drug (being a horse wormer) described in Article 5(1)(*a*) of the order by (1) the product licence holder, (2) a specially authorised person and (3) a person carrying on a qualifying business or a saddlery business (that is to say, a business involving, at least in part, the retail sale of saddlery requisites) provided, in each case, that the conditions contained in Article 5 are complied with.

These conditions are similar to those contained in Article 3 except that horse wormers may only be sold by retail to keepers of horses and ponies. The code of practice to be complied with is the Code of Practice for Saddlers Selling or Supplying Horse Wormers dated October 1985 (published by the Ministry of Agriculture, Fisheries and Food) and the registration and retention fee is £30 for each premises in respect of which a person is registered (Article 5(10) and (11)(*a*) and (*b*)). There is provision for the restoration of the registration of a person's name where he fails to make proper application for retention for which the fee is £60 for each premises in respect of which he is registered (Article 5(11)(*c*)). (These fees are not payable by a person carrying on a qualifying business on premises in respect of which his name is entered in the Register for the purposes of the exemption contained in Article 3).

The exemption contained in Article 5 applied for a period of three years from the commencement of the 1984 Order and has two years still to run from 1st January 1986.

Article 6 of the order provides an exemption to that part of the requirement under section 52(*c*) of the Act (that the retail sale, or supply in circumstances corresponding to retail sale, of a medicinal product which is not on a general sale list shall, where the transaction is carried out on behalf of a pharmacist by a person not being himself a pharmacist, be under the supervision of a pharmacist) where the sale is of a veterinary drug described in Article 3(1)(*a*) and is carried out in a registered pharmacy by a person acting on behalf of a pharmacist.

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

The changes of substance made by this order are:—

(1) the definitions of "qualifying business" and "saddlery business" now cover those businesses in which, respectively, any agricultural requisites

or saddlery requisites are sold by retail whereas previously turnover in such requisites had to exceed one half of total sales in the last accounting period (Articles 2 and 5);

(2) fees for registration are now calculated according to the number of premises in respect of which a person's name is registered (Articles 3(11) and 5(10)); there is provision for restoration of a person's name to the respective registers where proper application for retention has not been made (Articles 3(10)(*b*) and 5(9)(*b*)); there are revised fees for registration and retention in the registers and the introduction of a fee for restoration; there is provision for the removal of a person's name in respect of any premises from the relevant registers at his request (Articles 3(16)(*b*) and 5(15)(*b*));

(3) in Article 6, there is a relaxation of the requirement under section 52(*c*) of the Act which requires a pharmacist to supervise the sale of veterinary drugs as are described in Article 3(1)(*a*).

(The Codes of Practice referred to in the order (Articles 3(12)(*b*) and 5(11)(*b*)) are priced publications and are available from MAFF Publications Unit, Willowburn Estate, Alnwick, Northumberland, NE66 2PF).

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