

SCHEDULE 1

Regulation 4(3)

Marketing authorisations

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PART 1

Application for a marketing authorisation

Application for a marketing authorisation

1. An application under these Regulations for a marketing authorisation for a veterinary medicinal product must be made to the Secretary of State.

Information with the application

2.—(1) An application must include all necessary administrative information, and all scientific documentation necessary for demonstrating the safety, quality and efficacy of the product.

(2) In particular, the applicant must provide all the data required in Annex I to [Directive 2001/82/EC](#) of the European Parliament and of the Council on the Community code relating to veterinary medicinal products⁽¹⁾, generated in accordance with that Annex.

(3) The application must contain the following information—

- (a) the name of the person who will hold the marketing authorisation, that person’s address and, if different, the name and address of all the manufacturers involved in each stage of the manufacture, and the sites where the manufacture will take place;
- (b) the name of the veterinary medicinal product, which may be either—
 - (i) an invented name provided that this is not liable to be confused with the common name of the product or the international non-proprietary name (INN) recommended by the World Health Organization; or
 - (ii) a common or scientific name accompanied by a trademark or the name of the marketing authorisation holder;
- (c) the qualitative and quantitative particulars of all the constituents of the veterinary medicinal product, including its INN recommended by the World Health Organization, where an INN exists, or its chemical name;
- (d) a description of the method of manufacture;

(1) OJ No L 211, 28.11.2001, p. 1 as last amended by Regulation [\(EC\) No 470/2009](#) of the European Parliament and of the Council (OJ No L 152, 16.6.2009, p. 11). Annex I was inserted by Commission [Directive 2009/9/EC](#) (OJ No L 44, 14.2.2009, p. 10).

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- (e) all therapeutic indications, contra-indications and adverse reactions;
 - (f) the dosage for each species of animal for which the veterinary medicinal product is intended, its pharmaceutical form, method and route of administration and proposed shelf life;
 - (g) any proposed precautionary and safety measures to be taken when storing the veterinary medicinal product, administering it to animals or disposing of waste, together with an indication of potential risks that the veterinary medicinal product might pose to the environment, to human or animal health or to plants, together with the reasons;
 - (h) in the case of medicinal products intended for food-producing species, the proposed withdrawal period necessary to ensure that the maximum residue limits specified in Regulation (EC) No 470/2009 of the European Parliament and of the Council are not exceeded;
 - (i) a description of the testing methods to be used during manufacture;
 - (j) the results of—
 - (i) pharmaceutical (physico-chemical, biological or microbiological) tests;
 - (ii) safety tests and residue tests;
 - (iii) pre-clinical and clinical trials;
 - (iv) tests assessing the potential risks to the environment from the product;
 - (k) a detailed description of the pharmacovigilance system and, where appropriate, the risk management system that the applicant will put in place;
 - (l) a summary of the product characteristics, mock-ups of all proposed packaging and the proposed package leaflet, if any;
 - (m) a document showing that the manufacturer is authorised to produce veterinary medicinal products;
 - (n) copies (which must be updated if there are any changes while the application is being considered) of—
 - (i) any marketing authorisation obtained in another member State or in a third country for the relevant veterinary medicinal product, and a list of any other member States in which an application for authorisation of the product has been submitted;
 - (ii) if the product is already authorised outside the United Kingdom, the summary of product characteristics for each authorisation;
 - (iii) any decision to refuse authorisation, whether in the Community or a third country and the reasons for that decision;
 - (o) proof that the applicant has the services of a qualified person responsible for pharmacovigilance (referred to in these Regulations as a qualified person (pharmacovigilance)) and has the necessary means for the notification of any adverse reaction suspected of occurring either in the Community or in a third country;
 - (p) if the veterinary medicinal product is intended for food-producing species and contains one or more pharmacologically active substances not yet included for the species in question in Commission Regulation (EU) No 37/2010, a document certifying that a valid application for the establishment of maximum residue limits has been submitted to the Agency in accordance with Regulation (EC) No 470/2009 of the European Parliament and of the Council.
- (4) All documents relating to the results of tests or trials must be accompanied by a detailed and critical expert report that has been drafted and signed by a person with the requisite technical

or professional qualifications and that has a brief curriculum vitae of the person signing the report attached to it.

(5) In the case of immunological products, the applicant must submit a description of the methods used to establish that the manufacturing process will consistently produce a veterinary medicinal product that is in accordance with the marketing authorisation.

Summary of product characteristics

3. The summary of product characteristics required under the preceding paragraph must include the following information, set out in the same format—

Summary of product characteristics	
1	Name of the veterinary medicinal product, followed by its strength and pharmaceutical form.
2	The name and proportion of each active substance, and of any excipient if knowledge of the excipient is needed for safety reasons.
3	Pharmaceutical form.
4	Clinical particulars—
4.1	target species;
4.2	indications for use, specifying the target species;
4.3	contra-indications;
4.4	special warnings for each target species;
4.5	special precautions for use, including special precautions to be taken by the person administering the medicinal product to the animals;
4.6	adverse reactions (frequency and seriousness);
4.7	use during pregnancy, lactation or lay;
4.8	interaction with other medicinal products and other forms of interaction;
4.9	amounts to be administered and administration route;
4.10	overdose (symptoms, emergency procedures, antidotes) if necessary;
4.11	withdrawal periods for the various foodstuffs, including those for which the withdrawal period is zero.
5	Pharmacological properties—
5.1	pharmacodynamic properties;
5.2	pharmacokinetic particulars;
6	Pharmaceutical particulars—
6.1	list of excipients;
6.2	major incompatibilities;
6.3	shelf life, when necessary after reconstitution of the medicinal product or when the immediate packaging is opened for the first time;
6.4	special precautions for storage;

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6.5	nature and contents of immediate packaging;
6.6	special precautions for the disposal of unused veterinary medicinal products or waste materials derived from the use of such products, if appropriate;
7	Marketing authorisation holder;
8	Marketing authorisation number;
9	Date of the first authorisation or date of renewal of the authorisation;
10	Date of any revision of the text;
11	Any other information required by the Secretary of State.

Supply of a copy of the summary of product characteristics

4. A holder of a marketing authorisation must supply a copy of the summary of product characteristics to any person on demand.

Time limits for applications for products for use in food-producing animals

5. In the case of a veterinary medicinal product for food-producing animals, a marketing authorisation may not be applied for until at least six months after a valid application has been made for the establishment of a maximum residue limit in accordance with Regulation (EC) No 470/2009 of the European Parliament and of the Council.

PART 2

Derogations from some of the requirements in Part 1

Scope

6. This Part provides for applications for marketing authorisations in which not all the information required in Part 1 is required, but for the avoidance of doubt any applicant may apply for a marketing authorisation using Part 1 if the applicant wishes to do so.

Bibliographic application

7.—(1) An applicant for a marketing authorisation need not provide the results of safety tests, residue tests, pre-clinical trials or clinical trials if the active substance of the veterinary medicinal product has been in an authorised veterinary medicinal product for that species in the Community for at least ten years, and the applicant provides appropriate scientific literature to demonstrate this.

(2) The applicant may use any publicly available document.

(3) If an applicant makes use of scientific literature to obtain authorisation for a food-producing species, and submits, in respect of the same medicinal product and with a view to obtaining authorisation for another food-producing species, new residue studies, together with further clinical trials, a third party may not use those studies or trials in an application for a pharmacologically equivalent product for a period of three years from the grant of the authorisation for the additional species.

Application for a product using a new combination of active substances

8. If an application is for a veterinary medicinal product containing active substances already used in an authorised veterinary medicinal product but not previously used in that combination in a veterinary medicinal product, the applicant need not provide the safety and efficacy data for the individual active substances.

Application using existing data

9. If the Secretary of State has granted a marketing authorisation, the Secretary of State may, with the permission of the holder, use the data submitted in support of that marketing authorisation when assessing an application for another marketing authorisation.

Application for a pharmacologically equivalent medicinal product

10.—(1) An applicant need not provide the results of safety tests, residue tests, pre-clinical trials or clinical trials if the applicant can demonstrate that the veterinary medicinal product is pharmacologically equivalent to a veterinary medicinal product already authorised in the Community.

(2) For the purposes of this paragraph a product is pharmacologically equivalent to an existing product if—

- (a) it has the same qualitative and quantitative composition in active substances;
- (b) it has the same pharmaceutical form; and
- (c) bioequivalence has been demonstrated by means of appropriate bioavailability studies.

(3) For the purposes of this paragraph—

- (a) the different salts, esters, ethers, isomers, mixtures of isomers, complexes or derivatives of an active substance are considered to be the same active substance, unless they differ significantly in properties with regard to safety or efficacy; and
- (b) if they do differ significantly in properties with regard to efficacy or safety, additional information intended to provide proof of the safety or efficacy of the various salts, esters or derivatives of an authorised active substance must be supplied by the applicant.

(4) Different immediate-release oral pharmaceutical forms are regarded as the same pharmaceutical form.

(5) Bioavailability studies are not required if the bioequivalence guidelines produced by the Agency exempt the product.

(6) In the case of a reference product authorised in another member State but not in the United Kingdom, the Secretary of State must be satisfied that the risk-benefit balance of the original product is appropriate for the product to be placed on the market in the United Kingdom, and if the data provided under Article 13, third paragraph of [Directive 2001/82/EC](#) by the member State in which the product is authorised are insufficient for the Secretary of State to be satisfied of this, the Secretary of State may notify the applicant and require the applicant to provide further data.

Time limits for marketing authorisations granted under the procedure for a pharmacologically equivalent product

11.—(1) This paragraph establishes the time limits relating to granting a marketing authorisation under the procedure for a pharmacologically equivalent product.

(2) An application for a marketing authorisation cannot be made until two years before the product may be placed on the market in accordance with this paragraph.

(3) The product may not be placed on the market until ten years (or, in the case of medicinal products for fish or bees where the application for a marketing authorisation was submitted after 30th October 2005, thirteen years) have elapsed from the initial authorisation of the reference product.

(4) Time limits in this paragraph are calculated from the first grant of the marketing authorisation for the reference product.

Extension of time limits

12.—(1) This paragraph applies in relation to veterinary medicinal products that—

- (a) are intended for administration to food-producing species; and
- (b) contain a new active substance that was not authorised in the Community by 30th April 2004.

(2) If a person submitted an application for a marketing authorisation for a product on or after 30th October 2005, and within 5 years of the original marketing authorisation being granted, the marketing authorisation is extended to include additional food-producing species, the ten-year protection period is extended by one year for each additional food-producing species added to the marketing authorisation.

(3) The total period may not exceed 13 years.

(4) The extension applies only if the marketing authorisation holder originally applied for determination of the maximum residue limits for the active substance.

Parallel imports

13.—(1) The Secretary of State may grant a marketing authorisation in relation to a veterinary medicinal product authorised in another member State and imported into the United Kingdom from that member State in accordance with this paragraph without the data required in Part 1 if the applicant can demonstrate compliance with this paragraph.

(2) If the product is for a food-producing species it must be identical to a product authorised in the United Kingdom.

(3) Other products must be therapeutically the same as a product authorised in the United Kingdom unless the importer can justify any differences.

(4) The member State from which it is imported must have authorised the product in accordance with [Directive 2001/82/EC](#).

(5) The applicant must be established within the Community.

(6) The applicant must hold (or have a contract with the holder of) a wholesale dealer's authorisation in the United Kingdom appropriate to the type of product to be imported.

(7) If re-labelling is to take place in the United Kingdom the applicant must also be (or have a contract with) the holder of a suitable manufacturing authorisation in the United Kingdom.

Specific batch control scheme

14.—(1) Where a veterinary medicinal product (other than a biological veterinary medicinal product) has been granted a marketing authorisation or an animal test certificate, and any starting material (active substance, excipient or packaging) or any batch of the product does not fully meet the requirements of the authorisation or animal test certificate, the holder may apply to the Secretary of State to place one or more batches on the market notwithstanding this.

(2) The Secretary of State may authorise the placing on the market on being satisfied that the safety, quality and efficacy of the product are not compromised, and that in all the circumstances of the case the product should be placed on the market.

(3) This paragraph does not apply in relation to a product recognised in more than one member State.

(4) In this paragraph a biological veterinary medicinal product is a veterinary medicinal product, the active substance of which is a biological substance; and a biological substance is a substance that is produced by or extracted from a biological source and for which a combination of physico-chemical-biological testing and the production process and its control is needed for its characterisation and the determination of its quality.

Similar immunological products

15. Where an immunological veterinary medicinal product is pharmacologically equivalent to a reference product other than differences in raw materials or in the manufacturing process, the results of the appropriate pre-clinical tests or clinical trials must be provided, but the applicant need not provide the results of safety tests or residue tests.

Marketing a product authorised in another country

16. Where the health situation so requires, the Secretary of State may authorise the placing on the market of a veterinary medicinal product that has been authorised by another member State or, if there is no such authorised product, authorised in a third country.

PART 3

Grant of a marketing authorisation

Time limits

17. The Secretary of State must ensure that the procedure for granting an authorisation for a veterinary medicinal product is completed within a maximum of 210 days after the submission of the application.

Place of establishment of applicant

18. Only an applicant established in a member State may be granted a marketing authorisation.

Procedure

19. The Secretary of State may require the applicant to provide additional information or to generate additional data, including laboratory testing, or may require the applicant to provide samples of any medicinal product, its starting materials and intermediate products or other constituent materials for testing in a laboratory.

Products authorised in another member State

20. Where the Secretary of State is informed or discovers that another member State has authorised a veterinary medicinal product that is the subject of an application for authorisation by the Secretary of State, the Secretary of State must reject the application unless it was submitted in accordance with the mutual recognition procedure or the decentralised procedure in Part 6.

Assessment reports

21. The Secretary of State must produce an assessment of the dossier, consisting of an evaluation of the results of the pharmaceutical, safety and residue tests and the pre-clinical and clinical trials of the veterinary medicinal product concerned, and any additional related information.

Grant of a marketing authorisation

22. When granting a marketing authorisation, the Secretary of State must inform the applicant of the summary of product characteristics that has been approved, and the distribution category of the product.

Marketing authorisations for food-producing species

23.—(1) The Secretary of State must not grant a marketing authorisation for a veterinary medicinal product for food-producing species unless all its pharmacologically active substances appear in Table 1 in the Annex to Commission Regulation (EU) No 37/2010.

(2) This does not apply in the case of a marketing authorisation for a veterinary medicinal product for administration to a horse that has been declared on its horse passport as not intended for slaughter for human consumption; but in this case the product must not include an active substance that appears in Table 2 in the Annex to Commission Regulation (EU) No 37/2010 and must not be intended for the treatment of a condition for which a veterinary medicinal product is already authorised for horses.

Refusal of a marketing authorisation

24.—(1) The Secretary of State must refuse to grant a marketing authorisation if the application does not comply with these Regulations.

(2) In addition, the Secretary of State must refuse to grant it if—

- (a) the data submitted with the application are inadequate;
- (b) the risk-benefit balance of the veterinary medicinal product is unfavourable;
- (c) the product has insufficient therapeutic effect;
- (d) the withdrawal period proposed by the applicant is not long enough to ensure that Regulation (EC) No 470/2009 of the European Parliament and of the Council is complied with, or is insufficiently substantiated;
- (e) the veterinary medicinal product is for a prohibited use;
- (f) the way that the product will be used will have an unnecessarily undesirable effect on the environment.

(3) The Secretary of State may refuse to grant a marketing authorisation—

- (a) if there is Community legislation pending that is incompatible with the requested authorisation; or
- (b) if additional data have been requested and those data are not provided within such time limit as may be stipulated.

(4) If the Secretary of State, on the grounds of safety, quality or efficacy, intends to refuse an application, or proposes to grant a marketing authorisation that is different from the one applied for, the Secretary of State must notify the applicant accordingly, and the applicant may appeal to the Veterinary Products Committee.

Publication following the grant of a marketing authorisation

25.—(1) On granting a marketing authorisation the Secretary of State must publish—

- (a) the notice granting the marketing authorisation;
- (b) the summary of the product characteristics;
- (c) the assessment report that has already been prepared but with any commercially confidential or personal information deleted.

(2) The Secretary of State must update the assessment report whenever new information that is of importance and relates to the quality, safety or efficacy of the veterinary medicinal product becomes available.

(3) The Secretary of State must send a copy of the assessment report, and any update, to the holder of the marketing authorisation before publication to enable the holder to make representations concerning any confidential or personal information that may be in it, and may specify a date by which representations must be made.

Marketing authorisations in exceptional circumstances

26.—(1) In exceptional circumstances, and if there is no other product with a full marketing authorisation for the indicated condition in the target species, the Secretary of State may grant an exceptional marketing authorisation consisting of—

- (a) a provisional marketing authorisation subject to a requirement for the applicant to provide further data; or
- (b) a limited marketing authorisation for a product with a limited market.

(2) The Secretary of State must reassess each provisional or limited marketing authorisation annually.

Provisions of samples and expertise

27.—(1) The Secretary of State may require a marketing authorisation holder to provide, at any time and at any stage of the manufacturing process, samples of starting materials or the veterinary medicinal product for testing.

(2) At the request of the Secretary of State, the marketing authorisation holder must provide technical expertise to facilitate any analysis of the product.

Supply of information

28.—(1) A marketing authorisation holder must immediately inform the Secretary of State on receipt of any new information that might adversely affect the risk-benefit balance of the veterinary medicinal product.

(2) The holder must immediately inform the Secretary of State of any prohibition or restriction imposed by the competent authorities of any country in which the veterinary medicinal product is authorised.

(3) The Secretary of State may at any time require the marketing authorisation holder to provide data relating to the risk-benefit balance.

Duties on the holder of a marketing authorisation relating to an immunological product

29.—(1) Before placing an immunological product on the market the holder of the marketing authorisation must either—

- (a) notify the Secretary of State asking for written approval to do so; or
- (b) if the holder has already received written approval from another member State permitting the release of the product, send a copy of that approval to the Secretary of State.

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(2) If notified under sub-paragraph (1)(a) the Secretary of State must give or refuse a written approval as soon as is reasonably practicable.

(3) No person may place an immunological product on the market without a written approval issued by the Secretary of State or (if the approval was issued by another member State) without sending a copy of that approval to the Secretary of State.

Control tests

30. The holder of a marketing authorisation must give to the Secretary of State on demand evidence that the holder has carried out all control tests required under the marketing authorisation, and the results of those tests.

Placing on the market

31.—(1) A holder of a marketing authorisation must notify the Secretary of State when the veterinary medicinal product is first placed on the market in the United Kingdom, and the date on which it was placed on the market.

(2) A holder of a marketing authorisation who removes the veterinary medicinal product from the market in the United Kingdom must notify the Secretary of State at least two months (or a shorter period in exceptional circumstances) before doing so.

(3) Upon request by the Secretary of State, the marketing authorisation holder must provide—

- (a) all data relating to the volume of sales of the veterinary medicinal product by the holder; and
- (b) any data in the holder's possession relating to the number of prescriptions written for the product and the total volume supplied under those prescriptions.

Duration and validity of a marketing authorisation

32.—(1) A marketing authorisation is initially valid for five years.

(2) The authorisation may be renewed after five years on the basis of a re-evaluation of the risk-benefit balance.

(3) An application for renewal must be made at least six months, and not more than nine months, before the marketing authorisation ceases to be valid.

(4) An applicant who applies for the renewal of the marketing authorisation must enclose a list of all documents concerning the product that the applicant has submitted to the Secretary of State since the marketing authorisation was granted.

(5) The Secretary of State may require the applicant to provide a copy of any of the listed documents at any time.

(6) Once renewed, the marketing authorisation is valid indefinitely unless, within five years of the renewal, the Secretary of State notifies the holder, on justified grounds relating to pharmacovigilance, that the authorisation will cease to be valid five years from the first renewal unless the holder applies for a further renewal.

(7) The further renewal is not time-limited.

(8) Any marketing authorisation granted under these Regulations that is not followed within three years of its granting by the actual placing on the market of the authorised veterinary medicinal product in the United Kingdom ceases to be valid.

(9) When a veterinary medicinal product authorised under these Regulations and previously placed on the market in the United Kingdom is not present on the market in the United Kingdom for a period of three consecutive years, its marketing authorisation ceases to be valid.

(10) The Secretary of State may, on human or animal health grounds, grant exemptions from sub-paragraphs (8) and (9).

PART 4

Variations of marketing authorisations on the application of the holder

Variation of a marketing authorisation

33.—(1) The Secretary of State is the competent authority for the purposes of [Commission Regulation \(EC\) No 1234/2008](#)(2).

(2) The holder of a marketing authorisation may apply to the Secretary of State for a variation of that marketing authorisation.

(3) An application for a variation under paragraph (2) may only relate to a “single variation” unless the application is submitted in accordance with—

- (a) Article 7 of [Commission Regulation \(EC\) No 1234/2008](#) (“grouped variations”), or
- (b) Article 20 of [Commission Regulation \(EC\) No 1234/2008](#) (“workshare variations”).

(4) The Secretary of State, when granting a variation of a veterinary medicinal product, may (unless there are exceptional circumstances necessary to protect human or animal health or the environment) specify transitional measures to enable products produced in accordance with the previous authorisation to continue to be marketed for the transitional period.

Refusal of a variation of a marketing authorisation

34.—(1) This paragraph applies in relation to the refusal by the Secretary of State of an application for a variation unless the procedure following the refusal of a variation is one of those set out in Article 13 of Regulation 1234/2008.

(2) The grounds on which the Secretary of State may refuse an application for a variation of a marketing authorisation are those set out in paragraph 24 of this Schedule (refusal of a marketing authorisation).

(3) The Secretary of State must give written reasons for refusing to grant a variation; and if—

- (a) those reasons are on the grounds of safety, quality or efficacy; and
- (b) the variation is Type II or an extension application (whether or not in each case as part of an application for a worksharing or grouped application),

the applicant may appeal to the Veterinary Products Committee.

Administrative variations

35.—(1) The holder of a marketing authorisation may apply for a minor change in a marketing authorisation to be made without the Secretary of State considering any scientific data (an “administrative variation”).

(2) If the Secretary of State grants an administrative variation, and subsequently establishes that this should have been a variation requiring consideration of scientific data, the Secretary of State may notify the marketing authorisation holder, require the holder to submit an application for a variation enabling data to be assessed and revoke the administrative variation.

(2) OJ No L334, 12.12.2008, p. 7.

Changes after a marketing authorisation has been issued

36. After a marketing authorisation has been issued, the holder must take account of scientific and technical progress in manufacturing and control methods, and apply to the Secretary of State for any variation in the marketing authorisation that may be required to enable that veterinary medicinal product to be manufactured and checked by means of generally accepted scientific methods.

Compulsory variation

37.—(1) If the Secretary of State decides, for any of the reasons for suspending a marketing authorisation specified in paragraph 38, or because the classification of a veterinary medicinal product should be changed, that a variation to a marketing authorisation is necessary, the Secretary of State must by a notification in writing to the holder of the marketing authorisation require that person to apply for a variation of the marketing authorisation, giving reasons for requiring the application to be made.

(2) The notification may specify a time limit within which the marketing authorisation holder must apply for the variation.

(3) If the variation is on the grounds of safety, quality or efficacy, the applicant may, within 28 days of the notification, appeal to the Veterinary Products Committee.

(4) If the marketing authorisation holder fails to apply for the variation within that time limit the Secretary of State may suspend or revoke the marketing authorisation.

PART 5

Suspension, etc. of a marketing authorisation

Suspension of a marketing authorisation: grounds

38.—(1) The Secretary of State may suspend a marketing authorisation at any time on being satisfied that —

- (a) this is necessary for the protection of animal or public health or the environment;
- (b) the terms of the marketing authorisation have not been complied with; or
- (c) the veterinary medicinal product has insufficient therapeutic effect.

(2) The Secretary of State may also suspend a marketing authorisation on being satisfied that a marketing authorisation holder has failed to make an application for a variation to take account of scientific and technical progress in manufacturing and control methods to enable the veterinary medicinal product to be manufactured and checked by means of generally accepted scientific methods.

(3) The Secretary of State must suspend a marketing authorisation on being satisfied that—

- (a) the risk-benefit balance is unfavourable;
- (b) the withdrawal period does not ensure that residues in foodstuffs obtained from the treated animal comply with Regulation (EC) No 470/2009 of the European Parliament and of the Council;
- (c) information given in the application documents is incorrect;
- (d) any control tests required have not been carried out;
- (e) changes have been made to the manufacturing process without the authority of the Secretary of State; or
- (f) any information required to be supplied to the Secretary of State has not been so supplied.

Suspension of a marketing authorisation: procedure

39.—(1) If a marketing authorisation is suspended the Secretary of State must notify the holder immediately, and, unless the Secretary of State directs otherwise, the suspension has immediate effect, and continues in effect unless the marketing authorisation is reinstated.

(2) If the suspension is on the grounds of safety, quality or efficacy, the holder may, within 28 days of the notification, appeal to the Veterinary Products Committee.

(3) If the veterinary medicinal product is authorised in more than one member State, the Secretary of State—

(a) must immediately refer the matter to the Agency, and must comply with a decision of the Commission within 30 days of the decision; and

(b) may suspend the marketing and the use of the veterinary medicinal product in the United Kingdom pending a decision of the Agency, but must inform the Commission and the other member States no later than the following working day of the reasons for the action.

(4) When a marketing authorisation is suspended, the Secretary of State may in addition prohibit the supply of the veterinary medicinal product, and if necessary require the marketing authorisation holder to recall the product.

Revocation

40. The Secretary of State may revoke any marketing authorisation that has been suspended for more than 28 days unless there is a current appeal to the Veterinary Products Committee, and may publicise a revocation in such manner as the Secretary of State sees fit.

Prohibiting the supply of veterinary medicinal products

41.—(1) In addition to the powers to suspend a marketing authorisation, the Secretary of State, on being satisfied that a product has not been manufactured in accordance with the marketing authorisation, may prohibit the supply of a veterinary medicinal product, and if necessary require the marketing authorisation holder to recall it.

(2) The prohibition on supply and the requirement for recall may be confined to specific production batches.

(3) In the case of an immunological veterinary medicinal product manufactured outside the United Kingdom, if a batch has had all the tests that were originally carried out by the manufacturer repeated by the competent authority of another member State, the Secretary of State may not prohibit the release of that batch if all the results have been submitted to the Secretary of State and the results demonstrate that the product is within the terms of the authorisation.

PART 6

Mutual recognition and multiple applications

Application for a marketing authorisation where one already exists in another member State

42.—(1) If a veterinary medicinal product has already received a marketing authorisation in another member State at the time of application, and the holder of the marketing authorisation applies for a marketing authorisation in the United Kingdom, the following procedure (“the mutual recognition procedure”) applies.

Status: This is the original version (as it was originally made).

(2) The applicant must submit to the Secretary of State a dossier identical to the one submitted to the competent authority of the member State in which the veterinary medicinal product has been authorised (“the reference member State”).

(3) If there is a marketing authorisation current in more than one member State the applicant must identify which member State is acting as the reference member State.

(4) An applicant applying in more than one member State must supply the Secretary of State with a list of all the States in which the applicant is applying.

(5) The Secretary of State must obtain an assessment report from the reference member State and, where appropriate, an explanation of any extension of the period of data protection.

(6) Within 90 days after receipt of the assessment report, the Secretary of State must, subject to the following provisions, either—

- (a) approve the assessment report, the summary of product characteristics, the labelling and the package leaflet, and inform the reference member State accordingly; or
- (b) notify the reference member State that they have not been approved, and provide the reference member State with a detailed statement of the reasons.

(7) The Secretary of State may only refuse an application on the grounds of serious risk to human or animal health or the environment.

(8) If the assessment report, the summary of product characteristics, the labelling and the package leaflet are approved, the Secretary of State must ensure that a decision whether or not to grant a marketing authorisation can be made within 30 days of the approval.

(9) If the Secretary of State is notified by the reference member State that—

- (a) not all member States concerned have within 90 days approved the assessment report, summary of product characteristics, labelling or package leaflet; and
- (b) the reference member State has sent a detailed statement of the reasons to the other member States involved in the application, the applicant and the coordination group for action in accordance with Article 33(3) of [Directive 2001/82/EC](#),

the Secretary of State must within 30 days comply with the decision of the coordination group or, if the coordination group refers the matter to the Agency, the decision of the Commission.

(10) The Secretary of State may grant the marketing authorisation even though not all member States have agreed to grant it, but must revoke or vary the authorisation if this is necessary to comply with the decision of the Commission when it is received.

Application in another member State

43.—(1) When the Secretary of State has granted a marketing authorisation for a veterinary medicinal product and is notified by the marketing authorisation holder that the marketing authorisation holder has applied to have that veterinary medicinal product authorised in another member State, the Secretary of State must prepare an assessment report for the product within 90 days of the notification and send it to the member State or States concerned.

(2) If the other member State (or, if there is more than one, all of them) agrees with the assessment report, the summary of product characteristics, the labelling and the package leaflet the Secretary of State need take no further action.

(3) If not all the other member States concerned so agree within a further 90 days the Secretary of State must send a detailed statement setting out why they have disagreed to the other member States, the applicant and the coordination group for action in accordance with Article 33(3) of [Directive 2001/82/EC](#).

(4) The Secretary of State must within 30 days comply with the decision of the coordination group or, if the coordination group refers the matter to the Agency, the decision of the Commission.

Application for a marketing authorisation in multiple member States where a marketing authorisation does not exist in any member State

44.—(1) If an applicant wishes to apply for a marketing authorisation in more than one member State, and a marketing authorisation does not exist in any member State for the product (“the decentralised procedure”), the applicant must—

- (a) apply simultaneously in all the relevant member States;
- (b) submit a dossier to the Secretary of State that is identical to the dossier being submitted to all the other member States;
- (c) include a list of all member States in which applications have been made; and
- (d) nominate one of them to act as the reference member State to prepare a draft assessment report and drafts of the summary of product characteristics, labelling and package leaflet for consideration by the other member States (“the concerned member States”).

(2) If the United Kingdom is the reference member State, the Secretary of State must prepare a draft assessment report and drafts of the summary of product characteristics, labelling and package leaflet within 120 days of the receipt of a valid application and must send them to the other concerned member States and to the applicant.

(3) If the United Kingdom is not the reference member State, within 90 days after receipt of the assessment report and drafts of the summary of product characteristics, labelling and package leaflet from the reference member State, the Secretary of State must, subject to the following provisions, either—

- (a) approve the assessment report, the summary of product characteristics, the labelling and the package leaflet, and inform the reference member State accordingly; or
- (b) notify the reference member State that the Secretary of State will not approve it, and provide the reference member State with a detailed statement of the reasons.

(4) The Secretary of State may only refuse an application on the grounds of serious risk to human or animal health or the environment.

(5) If all the member States involved agree the assessment report, the summary of product characteristics, the labelling and the package leaflet within 90 days, the Secretary of State must ensure that a decision whether or not to grant a marketing authorisation can be made within 30 days.

(6) If, within 90 days, not all the member States have agreed the assessment report, summary of product characteristics, labelling and package leaflet on grounds of a potential serious risk to human or animal health or to the environment, the Secretary of State (if the United Kingdom is the reference member State) must send a detailed statement of the reasons to the other member States involved in the application, the applicant, and the coordination group to act in accordance with Article 33(3) of [Directive 2001/82/EC](#).

(7) If reference has been made to the coordination group by any member State, the Secretary of State must within 30 days comply with the decision of the coordination group or, if the coordination group refers the matter to the Agency, the decision of the Commission.

(8) If the Secretary of State wishes to do so, the Secretary of State may grant the marketing authorisation even though not all member States have agreed to grant it, but must revoke or vary the authorisation if this is necessary to comply with the decision of the Commission when it is received.

PART 7

Labelling and package leaflets

Approval by the Secretary of State

45. The Secretary of State, when issuing a marketing authorisation, must approve all containers, packaging, labels and package leaflets.

Reference to being authorised

46. A label and package leaflet of an authorised veterinary medicinal product may contain in legible characters the words “UK authorised veterinary medicinal product” or, if the marketing authorisation provides, other wording specified in the authorisation indicating that the product is authorised in the United Kingdom.

Language

47.—(1) All labels and package leaflets must be in English, but may contain other languages provided that the information given is identical in all the languages.

(2) This requirement does not apply in the case of a product imported by a veterinary surgeon and administered by or under the responsibility of that same veterinary surgeon.

Labelling with all the information on the immediate packaging

48.—(1) If it is reasonably practicable to do so, the following must be provided on the immediate packaging, in legible characters—

- (a) the name, strength and pharmaceutical form of the veterinary medicinal product;
- (b) the name and strength of each active substance, and of any excipient if this is required under paragraph 2 of the summary of product characteristics;
- (c) the route of administration (if not immediately apparent);
- (d) the batch number;
- (e) the expiry date;
- (f) the words “For animal treatment only” and, if appropriate, “To be supplied only on veterinary prescription”;
- (g) the contents by weight, volume or number of dose units;
- (h) the marketing authorisation number;
- (i) the name and address of the marketing authorisation holder or, if there is a distributor authorised in the marketing authorisation, that distributor;
- (j) a suitably labelled space to record discard date (if relevant);
- (k) the target species;
- (l) the distribution category;
- (m) the words “Keep out of reach of children”;
- (n) storage instructions;
- (o) the in-use shelf-life (if appropriate);
- (p) for food-producing species, the withdrawal period for each species or animal product concerned;

- (q) any warning specified in the marketing authorisation;
- (r) disposal advice;
- (s) full indications;
- (t) dosage instructions;
- (u) contra-indications;
- (v) further information required in the marketing authorisation;
- (w) if the product is one that requires a dose to be specified for the animal being treated, a space for this.

(2) If all this is on the immediate packaging, there is no need for any outer packaging or a package leaflet.

Products with immediate and outer packaging

49.—(1) If it is not reasonably practicable to have all the required information on the immediate packaging then this paragraph applies.

(2) The immediate packaging must have at least the following information—

- (a) the name of the veterinary medicinal product, including its strength and pharmaceutical form;
- (b) the name and proportion of each active substance, and of any excipient if knowledge of the excipient is needed for safety reasons;
- (c) the route of administration (if not immediately apparent);
- (d) the batch number;
- (e) the expiry date;
- (f) the words “For animal treatment only” and if appropriate, “To be supplied only on veterinary prescription”;
- (g) the words “Keep the container in the outer carton”.

(3) In addition, the immediate packaging must have as much of the required information as is reasonably practicable.

(4) The outer packaging must contain all the required information if it is reasonably practicable to do this, and if it is not reasonably practicable to do this a package leaflet must be supplied with the product in accordance with the following paragraph.

Package leaflets

50.—(1) If it is not reasonably practicable to have all the required information on the immediate packaging or all of this information on the outer packaging, there must be a package leaflet supplied with the product, containing all the required information except for the batch number and the expiry date, and including the name of both the marketing authorisation holder and, if different, the name of the distributor named in the marketing authorisation.

(2) If there is a package leaflet, the immediate packaging and the outer packaging must both refer the user to it.

(3) A package leaflet must relate solely to the veterinary medicinal product with which it is included.

(4) It must be written in plain English.

(5) Only a package leaflet approved in the marketing authorisation may be included with the veterinary medicinal product.

Status: This is the original version (as it was originally made).

Ampoules

51.—(1) In the case of ampoules or other unit dose forms, where the container cannot bear legibly the required information, only the following information must be shown on the immediate packaging—

- (a) the name of the veterinary medicinal product;
- (b) the name and strength of the active ingredient;
- (c) the route of administration (if not immediately apparent);
- (d) the batch number;
- (e) the expiry date;
- (f) the words “For animal treatment only” and if appropriate, “To be supplied only on veterinary prescription”.

(2) The outer packaging must contain all the required information if it is reasonably practicable to do this, and if it is not reasonably practicable to do this a package leaflet must be supplied with the product, except that the ampoule need not refer to the package leaflet.

Small containers other than ampoules

52. As regards small immediate packaging containing a single dose, other than ampoules, on which it is impossible to give the required information, all the required information must appear on the outer packaging or outer packaging and package leaflet, but the immediate packaging must be labelled with the batch number and the expiry date and, if there is room, the other information in the preceding paragraph.

Homeopathic remedies

53.—(1) A homeopathic remedy registered under these Regulations must be labelled in accordance with this paragraph.

(2) There must be no specific therapeutic indication on the labelling or in any information relating to it.

(3) The labelling (or labelling and package leaflet) must contain the following and no other information—

- (a) the words “homeopathic remedy without approved therapeutic indications for veterinary use”;
- (b) the scientific name of the stock or stocks followed by the degree of dilution, using the symbols of the pharmacopoeia used (if the homeopathic remedy is composed of more than one stock, the labelling may mention an invented name in addition to the scientific names of the stocks);
- (c) the name and address of the registration holder and (on the package leaflet) of the manufacturer;
- (d) the method and, if necessary, route of administration;
- (e) the expiry date;
- (f) the pharmaceutical form;
- (g) the contents of the pack;
- (h) any special storage precautions;
- (i) the target species;
- (j) any necessary special warnings;

- (k) the batch number; and
- (l) the registration number.

Variations

54. The Secretary of State may permit variations in the above in any individual marketing authorisation if this is necessary for public or animal health purposes or the protection of the environment.

PART 8

Pharmacovigilance

Qualified persons responsible for pharmacovigilance

55. A marketing authorisation holder must have permanently and continuously the services of an appropriately qualified person responsible for pharmacovigilance (“a qualified person (pharmacovigilance)”) who resides in a member State.

Duties relating to the qualified person

56. The marketing authorisation holder must ensure that the qualified person (pharmacovigilance)—

- (a) establishes and maintains a system that ensures that information about all suspected adverse reactions reported to the marketing authorisation holder is collected and collated in order to be accessible at least at one point in a member State;
- (b) answers any request from the Secretary of State for the provision of additional information necessary for the evaluation of the benefits and risks afforded by a veterinary medicinal product fully and within any time limit imposed by the Secretary of State when the information was requested, including the volume of sales of the veterinary medicinal product concerned and, if available, details of prescriptions;
- (c) provides to the Secretary of State any other information relevant to the evaluation of the benefits and risks afforded by a veterinary medicinal product, including appropriate information on post-marketing surveillance studies; and in this paragraph “post-marketing surveillance studies” means a pharmacoepidemiological study or a clinical trial carried out in accordance with the terms of the marketing authorisation, conducted with the aim of identifying and investigating a safety hazard relating to an authorised veterinary medicinal product.

Adverse reactions to a veterinary medicinal product administered in the United Kingdom

57.—(1) A marketing authorisation holder must act in accordance with this paragraph on learning of any suspected—

- (a) serious adverse reaction;
- (b) human adverse reaction; or
- (c) unintended transmission of an infectious agent through a veterinary medicinal product,

following the administration of the product in the United Kingdom.

- (2) The holder must make a record of what happened.

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(3) The holder must without delay and in any event within 15 days report it (electronically if this is practicable) to the Secretary of State.

(4) In addition, the holder must supply to the Secretary of State all relevant veterinary pharmacovigilance information that the holder possesses relating to the reaction, giving a full description of the incident and a list of all the symptoms using internationally recognised veterinary and medical terminology, either with the report or, if the information becomes available after the report has been sent, as soon after it becomes available as is reasonably practicable.

(5) In this and the following paragraph—

“human adverse reaction” means a reaction that is noxious and unintended and that occurs in a human being following exposure to a veterinary medicine;

“serious adverse reaction” means an adverse reaction that results in death, is life-threatening, results in significant disability or incapacity, is a congenital anomaly or birth defect, or that results in permanent or prolonged signs in the animals treated.

Adverse reactions to a veterinary medicinal product administered in a third country

58.—(1) A marketing authorisation holder for a veterinary medicinal product authorised in the United Kingdom must act in accordance with this paragraph on learning of any suspected—

- (a) serious, unexpected adverse reaction (for these purposes a reaction is unexpected if its nature, severity or outcome is not consistent with the summary of the product characteristics);
- (b) human adverse reaction; or
- (c) unintended transmission of an infectious agent through a veterinary medicinal product,

following the administration of the product in a third country.

(2) The holder must make a record of what happened.

(3) The holder must without delay and in any event within 15 days report the suspected reaction or transmission (electronically if this is practicable) to the Secretary of State, the competent authorities of all member States in which the product is authorised, and the Agency.

(4) In addition to the report, the holder must supply to the Secretary of State, the competent authorities of all other member States where the product is authorised and the Agency, all relevant veterinary pharmacovigilance information in the holder’s possession relating to the reaction as in the preceding paragraph.

Periodic safety update reports

59.—(1) The marketing authorisation holder must submit to the Secretary of State records of all adverse reactions (including nil reports) in the form of a periodic safety update report for each marketing authorisation in accordance with this paragraph, including a summary of each incident and a list of all the symptoms using internationally recognised veterinary and medical terminology.

(2) A marketing authorisation holder who has not yet placed a product on the market in the United Kingdom must submit a periodic safety update report immediately upon request of the Secretary of State and at least every six months after authorisation.

(3) Following the placing on the market in the United Kingdom, the marketing authorisation holder must submit a periodic safety update report to the Secretary of State immediately upon request and—

- (a) at least every six months during the first two years following the initial placing on the market;
- (b) once a year for the following two years; and

- (c) thereafter, at three-yearly intervals.
- (4) Following the granting of a marketing authorisation, the marketing authorisation holder may apply to the Secretary of State to change the periods of notification.
- (5) The periodic safety update report must include a scientific evaluation of the risk-benefit balance of the veterinary medicinal product.
- (6) The periodic safety update report must include—
 - (a) the volume of the product sold in each year covered by the report, calculated on an annual basis beginning 1st January;
 - (b) the number of adverse reactions for each year of the report;
 - (c) the ratio of adverse reactions to volume of product sold for each year of the report, together with an explanation of the basis of the calculation;
 - (d) differentiation of data based on—
 - (i) target species (if the product is authorised for use in more than one species);
 - (ii) reaction type (such as serious, non-serious, human, suspected lack of efficacy, unauthorised use or other);
 - (iii) the country of origin of the report.
- (7) If the product is indicated for more than one species, the information in sub-paragraph (6)(c) must be based so far as is practicable on the estimated use of the product.
- (8) Data relating to different formulations (either different dosage forms or different strengths) must be provided in separate reports.

Release of information by the marketing authorisation holder

60.—(1) A marketing authorisation holder must not communicate information relating to pharmacovigilance concerns to the general public in relation to its authorised veterinary medicinal product without giving prior or simultaneous notification to the Secretary of State.

(2) The marketing authorisation holder must ensure that such information is presented objectively and is not misleading.

Action taken on account of pharmacovigilance

61.—(1) Where, as a result of the evaluation of veterinary pharmacovigilance data, the Secretary of State considers that a marketing authorisation should be—

- (a) suspended;
- (b) revoked; or
- (c) varied so as to—
 - (i) restrict the indications;
 - (ii) change the distribution category;
 - (iii) amend the dose;
 - (iv) add a contraindication; or
 - (v) add a new precautionary measure,

the Secretary of State must forthwith inform the Agency, all other member States (irrespective of whether the product is authorised in another member State) and the marketing authorisation holder.

(2) If urgent action is necessary for protecting human or animal health, the Secretary of State may suspend the marketing authorisation of a veterinary medicinal product, but must inform the Agency, the Commission and the other member States within one working day.

(3) If, following the opinion of the Agency, the Commission requests the Secretary of State to suspend, withdraw or vary the marketing authorisation, the Secretary of State must comply with that request immediately on a temporary basis.

(4) The Secretary of State must take final measures in accordance with the Decision of the Commission.

PART 9

Homeopathic remedies

Meaning of “homeopathic remedy”

62. For the purposes of these Regulations, a homeopathic remedy is a veterinary medicinal product (which may contain a number of principles) prepared from homeopathic stocks in accordance with a homeopathic manufacturing procedure described in the European Pharmacopoeia⁽³⁾ or, if it is not described there, in a pharmacopoeia published by the British Pharmacopoeial Commission or by the competent authority of any member State.

Placing a homeopathic remedy on the market in accordance with a registration

63.—(1) By way of derogation from the provisions of these Regulations requiring a marketing authorisation for a veterinary medicinal product, a homeopathic remedy may be placed on the market in accordance with a registration by the Secretary of State instead of in accordance with a marketing authorisation if it complies with this paragraph.

(2) It must not be an immunological product.

(3) The route of administration must be as described in the European Pharmacopoeia or, if it is not described there, by a pharmacopoeia currently used officially in any member State.

(4) There must be a sufficient degree of dilution to guarantee the safety of the product, and in any event it must not contain more than one part in 10,000 of the mother tincture.

(5) All other provisions relating to marketing authorisations apply in the same way to registrations of a homeopathic remedy.

Application for registration

64.—(1) An applicant for registration must submit the following to the Secretary of State—

- (a) the scientific name or other name of the homeopathic stock given in a pharmacopoeia, together with a statement of the various routes of administration, pharmaceutical forms and degree of dilution;
- (b) a dossier describing how the homeopathic stock is obtained and controlled, and justifying its homeopathic nature, on the basis of an adequate bibliography;
- (c) in the case of a product containing biological substances, a description of the measures taken to ensure the absence of pathogens;
- (d) the manufacturing and control file for each pharmaceutical form and a description of the method of dilution and potentiation;

(3) ISBN 9287145873.

- (e) a copy of the manufacturing authorisation for the product;
- (f) copies of any registrations or authorisations obtained for the same homeopathic remedy in other member States;
- (g) a mock-up of the outer packaging and immediate packaging;
- (h) stability data;
- (i) the proposed withdrawal period necessary to ensure that the provisions of Regulation (EC) No 470/2009 of the European Parliament and of the Council are complied with together with all necessary justification.

(2) These documents must demonstrate the pharmaceutical quality and the batch-to-batch homogeneity of the products concerned.

(3) In the case of a food-producing animal, if the applicant states in the application that the homeopathic remedy contains an active substance, or has been manufactured using an active substance, that substance must be one that appears in Table 1 in the Annex to Commission Regulation (EU) No 37/2010 and complies with any requirements in that Table relating to that substance.

(4) If a product is registered in another member State, the Secretary of State may waive some or all of the requirements of this paragraph on being satisfied that it is reasonable to do so.

Procedure for registration

65.—(1) The procedure for registration is the same as the procedure for granting a marketing authorisation in accordance with Part 3, except—

- (a) the applicant is not required to provide proof of efficacy;
- (b) the product is not required to have a summary of product characteristics;
- (c) the Secretary of State is not required to publish an assessment report.

(2) The procedure for variation, suspension and revocation is the same as for a marketing authorisation.

Products on the market before 1994

66. A homeopathic remedy that was on the market before 1st January 1994 may be placed on the market without being registered.

Classification

67. The registration must specify the classification of the homeopathic remedy, which must be one of the classifications specified for a veterinary medicinal product in Schedule 3.

Offences

68. It is an offence to fail to comply with—

- (a) a requirement made under paragraph 27(1);
- (b) a request made under paragraph 27(2);
- (c) paragraph 28(1) or (2);
- (d) a requirement made under paragraph 28(3);
- (e) paragraph 29(3);
- (f) paragraph 30;
- (g) paragraph 31(1) or (2);

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- (h) a request made under paragraph 31(3);
- (i) a prohibition or requirement made under paragraph 39(4);
- (j) a prohibition or requirement made under paragraph 41(1);
- (k) paragraph 55;
- (l) paragraph 56;
- (m) paragraph 57;
- (n) paragraph 58;
- (o) paragraph 59; or
- (p) paragraph 60.

SCHEDULE 2

Regulation 5(2)

The manufacture of veterinary medicinal products

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PART 1

Manufacturing authorisations

Application

1. An application for a manufacturing authorisation must be made to the Secretary of State.

Time limits

2.—(1) The Secretary of State must process an application for a manufacturing authorisation within 90 days of receiving it.

(2) The Secretary of State must process an application for a variation of a manufacturing authorisation within 30 days unless the Secretary of State notifies the applicant in writing that the time has been extended to 90 days.

Granting the authorisation

3. The Secretary of State must grant a manufacturing authorisation on being satisfied that the applicant has suitable and sufficient premises, staff, technical equipment and facilities for the manufacture, control and storage of the products, and will comply with these Regulations.

The authorisation

4.—(1) The manufacturing authorisation must specify—

- (a) the types of veterinary medicinal products and pharmaceutical forms that may be manufactured or imported;
- (b) the place where they are to be manufactured or controlled;
- (c) the name and address of the person holding the authorisation;
- (d) the address of the premises to which it relates;
- (e) the names of all qualified persons nominated to act under this Schedule.

(2) It may specify that different activities must be carried out in different premises or parts of premises, and may require the holder of the manufacturing authorisation to restrict access to premises or parts of premises to persons carrying out activities there.

(3) The holder of a manufacturing authorisation must notify the Secretary of State, and if necessary apply for a variation of the authorisation, before making a material alteration to the premises or facilities used under the authorisation, or to the operations for which they are used.

Suspension, variation or revocation of the authorisation

5.—(1) The Secretary of State may suspend, vary or revoke a manufacturing authorisation if the holder—

- (a) has not complied with these Regulations;
- (b) has manufactured a veterinary medicinal product not authorised by the manufacturing authorisation;
- (c) has produced a veterinary medicinal product outside the terms of a marketing authorisation; or
- (d) no longer has suitable premises or equipment.

(2) The Secretary of State may also suspend, vary or revoke it on being satisfied that the qualified person (manufacturer) is not fulfilling their duties under these Regulations.

Inspection of premises

6.—(1) The Secretary of State must, from time to time, inspect premises registered under paragraph 3, basing the frequency of the inspection on the risks associated with each premises' history and the nature of the products handled at the premises.

(2) Within 90 days after an inspection, the Secretary of State must issue a certificate of good manufacturing practice to the manufacturer if the inspection established compliance with the principles and guidelines on good manufacturing practice in accordance with Commission [Directive 91/412/EEC](#) laying down the principles and guidelines of good manufacturing practice for veterinary medicinal products⁽⁴⁾.

(3) If an inspection is carried out at the request of the European Pharmacopoeia to establish compliance with a monograph, the Secretary of State must issue a certificate of compliance with the monograph, if appropriate.

(4) The Secretary of State must provide details of each certificate of good manufacturing practice issued to the Agency for entry into a database.

(4) OJ No L 228, 17.8.91, p. 70.

(5) If the outcome of the inspection is that the manufacturer does not comply with the principles and guidelines of good manufacturing practice, the Secretary of State must provide details to the Agency for entry into the database.

Report following inspection

7.—(1) After each inspection of manufacturing premises, the inspector must make a written report to the Secretary of State on whether the principles and guidelines on good manufacturing practice and the conditions of these Regulations are being complied with.

(2) The Secretary of State must inform the inspected manufacturer of the content of such reports.

Duties on the holder of a manufacturing authorisation

8.—(1) A holder of a manufacturing authorisation must ensure that the veterinary medicinal product is manufactured in accordance with the marketing authorisation.

(2) The holder must have permanently at their disposal the services of at least one qualified person (manufacture) who is on the register of qualified persons (manufacture) maintained by the Secretary of State and must place all necessary facilities at the qualified person's disposal.

(3) The holder must—

- (a) have a current Certificate of Good Manufacturing Practice;
- (b) have in place a system of Quality Assurance and Quality Control; and
- (c) give to the Secretary of State on request proof of all control tests carried out on the veterinary medicinal product or the constituents and intermediate products of the manufacturing process in accordance with the data submitted in support of the application for the marketing authorisation.

(4) A holder who makes up a bulk package of veterinary medicinal products must ensure that the package is labelled, in a way that the label is clearly visible and legible, with—

- (a) the name of the veterinary medicinal product, its strength as shown in the summary of product characteristics and its pharmaceutical form;
- (b) the batch number;
- (c) the expiry date;
- (d) any storage requirements; and
- (e) any other warning necessary for the safe handling of the package.

(5) A holder must keep an adequate number of representative samples of each batch of a veterinary medicinal product in stock at least until the expiry date of the batch, and must submit any such sample to the Secretary of State if required in writing to do so.

Qualified persons for manufacture

9.—(1) The Secretary of State may appoint as a qualified person (manufacture) any person who is—

- (a) a member of the Royal Pharmaceutical Society or registered with the Pharmaceutical Society of Northern Ireland;
- (b) a Chartered Chemist or a Fellow, Member or Associate Member of the Royal Society of Chemistry; or
- (c) a Chartered Biologist or a Fellow, Member or Associate Member of the Society of Biology;

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who qualified on the basis of a formal course of study lasting not less than three years full-time or equivalent and who has sufficient practical experience to carry out the duties under this Schedule.

(2) The Secretary of State may exceptionally appoint a person who is not a member of one of those institutions to act as a qualified person (manufacture) on being satisfied that that person has the educational qualifications or practical experience to carry out the duties under this Schedule.

Refusal or revocation of appointment

10. The Secretary of State may refuse or revoke an appointment if the Secretary of State is not satisfied that a person has fulfilled or will fulfil duties under these Regulations.

Duties on a qualified person

11.—(1) The qualified person (manufacture) must ensure that each batch of veterinary medicinal product manufactured under that person's responsibility is manufactured and checked in compliance with these Regulations and in accordance with the data submitted in support of the application for the marketing authorisation.

(2) If a manufacturer imports a veterinary medicinal product from a third country, including a product manufactured in a member State, the qualified person (manufacture) must ensure that, following importation, each production batch imported is fully tested in a member State, including a full qualitative analysis, a quantitative analysis of at least all the active substances and all the other tests or controls necessary to ensure the quality of a veterinary medicinal product is in accordance with the requirements of the marketing authorisation.

(3) Sub-paragraph (2) does not apply where appropriate arrangements have been made by the European Union with the exporting country to ensure that the manufacturer of the veterinary medicinal product applies standards of good manufacturing practice at least equivalent to those laid down in Commission [Directive 91/412/EEC](#) and to ensure that the controls in sub-paragraph (2) have been carried out in the exporting country.

(4) At each stage of manufacture, including release for sale, the qualified person (manufacture) must certify in writing that all control tests required under the marketing authorisation have been carried out, and that the production batch complies with the marketing authorisation.

Register

12. The Secretary of State must maintain and publish a register of—

- (a) holders of manufacturing authorisations; and
- (b) qualified persons (manufacture) appointed under paragraph 9(2).

Test sites

13.—(1) The Secretary of State may authorise premises to act as a test site to carry out contract testing for a holder of a manufacturing authorisation.

(2) The premises must have a current certificate of good manufacturing practice.

(3) Authorisation and inspection of the premises are the same as for a manufacturing authorisation.

PART 2

Authorisation of manufacturers of autogenous vaccines

Authorisation to manufacture autogenous vaccines

14.—(1) The Secretary of State may authorise a person to manufacture autogenous vaccines and may authorise premises for the purpose of such manufacture by that person.

(2) In order to be authorised the premises must be under the supervision of—

- (a) a veterinary surgeon; or
- (b) a person who the Secretary of State is satisfied has sufficient qualifications and experience to manufacture the product safely.

(3) Before authorising the premises, the Secretary of State must be satisfied that the production process will produce a consistent, safe product.

(4) No person may manufacture an autogenous vaccine other than in accordance with an authorisation under sub-paragraph (1).

Types of authorisation

15.—(1) The authorisation must specify the products that may be manufactured.

(2) It may either be for the production of a single batch of product or for ongoing production of the products specified in the authorisation.

(3) If it is for a single batch it must be time limited.

(4) Only the products specified in the authorisation may be manufactured, and in the case of an authorisation for a single batch the product may only be manufactured before the expiry of the authorisation.

Labelling

16. The operator of the premises must ensure that every container containing autogenous vaccine is labelled with—

- (a) the name of the veterinary surgeon who ordered the vaccine;
- (b) a precise description of the vaccine;
- (c) the date the vaccine was produced;
- (d) the name of the authorisation holder and address of the authorised premises;
- (e) the expiry date;
- (f) any necessary warnings; and
- (g) instructions for use.

Records

17. The operator of the premises must, as soon as is reasonably practicable, record—

- (a) the name and address of the veterinary surgeon who ordered the vaccine;
- (b) the identification of the source animal;
- (c) the expiry date;
- (d) the date of supply to the veterinary surgeon,

and must keep the records for at least five years.

Adverse reactions

18. The authorised person must notify the Secretary of State of any adverse reactions to an autogenous vaccine within 15 days of learning of the reaction.

Inspection of premises

19. The Secretary of State must inspect the authorised premises, basing the frequency of the inspection on the risks associated with each premises' history and the nature of the products handled at the premises.

PART 3

Authorisation of blood banks

Authorisation of blood banks

- 20.—(1) The Secretary of State may authorise blood banks for—
- (a) the collection, storage and supply of blood, or
 - (b) the storage and supply of blood constituents obtained by the physical separation of donor blood into different fractions within a closed-bag system,
- for the treatment of non-food-producing animals.
- (2) The authorisation may be for either or both of these activities.
- (3) In order to be authorised a blood bank must be under the supervision of—
- (a) a veterinary surgeon named in the authorisation; or
 - (b) a person named in the authorisation who the Secretary of State is satisfied is suitably qualified to operate the blood bank.
- (4) Before authorising a blood bank, the Secretary of State must be satisfied—
- (a) that the welfare of animals used in the collection of blood will be respected; and
 - (b) that the production process will produce a consistent, safe product.
- (5) The Secretary of State may suspend, vary or revoke an authorisation of a blood bank if—
- (a) the holder no longer uses fit and proper processes;
 - (b) the premises in which the blood bank is being or is to be operated are not suitable;
 - (c) the equipment is not suitable; or
 - (d) the holder has not complied with these Regulations.
- (6) Blood may only be collected under the responsibility of a veterinary surgeon.
- (7) No person may operate a blood bank for treatment of animals other than in accordance with such an authorisation.

Supply and administration of blood from a blood bank

- 21.—(1) The operator of a blood bank may only supply blood to a veterinary surgeon.
- (2) No person other than a veterinary surgeon or someone acting under a veterinary surgeon's responsibility may administer blood.
- (3) No person may administer blood to a food-producing animal.

Labelling

22.—(1) The operator of a blood bank must ensure that every container used for the blood is labelled with—

- (a) the identification of the donor animal;
- (b) the date of collection;
- (c) the authorisation number of the blood bank;
- (d) any necessary warnings;
- (e) the expiry date.

(2) There must be no specific therapeutic indication on the label or on any information relating to the product.

Records

23. The operator of a blood bank must, as soon as is reasonably practicable, record—

- (a) the date of collection;
- (b) the identification of the donor animal;
- (c) the veterinary surgeon who collected it;
- (d) the expiry date;
- (e) the date the blood was used or, if it was supplied to another veterinary surgeon, the name of that veterinary surgeon and the date it was supplied;

and must keep the records for at least five years.

Inspection of premises

24. The Secretary of State must inspect the authorised premises, basing the frequency of the inspection on the risks associated with each premises' history and the nature of the products handled at the premises.

PART 4

Authorisation of manufacturers of products for administration under the cascade

Authorisation to manufacture products for administration under the cascade

25.—(1) The Secretary of State may authorise a person and premises to manufacture an unauthorised veterinary medicinal product for administration under the cascade.

(2) In order to be authorised the premises must be under the supervision of a person who the Secretary of State is satisfied has sufficient qualifications and experience to manufacture the product safely.

(3) Before authorising the premises, the Secretary of State must be satisfied that the production process will produce a safe product.

(4) The authorisation must specify what types of product it covers.

(5) No person may manufacture an unauthorised veterinary medicinal product other than in accordance with an authorisation under sub-paragraph (1).

Labelling

26. The authorised person must ensure that, before a veterinary medicinal product is supplied, every container is labelled with—

- (a) the name of the veterinary surgeon who ordered the veterinary medicinal product;
- (b) a precise description of the veterinary medicinal product;
- (c) the date of production;
- (d) the name of the authorisation holder and the address of the authorised premises;
- (e) the expiry date;
- (f) any necessary warnings; and
- (g) instructions for use.

Records

27. The authorised person must, as soon as is reasonably practicable, record—

- (a) the name and address of the veterinary surgeon who ordered the veterinary medicinal product;
- (b) a precise description of the veterinary medicinal product;
- (c) the date of production;
- (d) the expiry date; and
- (e) the date of supply to the veterinary surgeon,

and must keep the record for at least five years.

Adverse reactions

28. The authorised person must notify the Secretary of State of any adverse reactions to a product manufactured by that person within 15 days of learning of the reaction.

Inspection of premises

29. The Secretary of State must inspect the authorised premises, basing the frequency of the inspection on the risks associated with each premises' history and the nature of the products handled at the premises.

PART 5

Authorisation of equine stem cell centres

Authorisation of stem cell centres

30.—(1) The Secretary of State may authorise equine stem cell centres for the collection, storage, processing, production and administration of equine stem cells for use as an autologous treatment for horses.

(2) In order to be authorised a centre must be under the supervision of—

- (a) a veterinary surgeon named in the authorisation; or
- (b) a person named in the authorisation who the Secretary of State is satisfied is suitably qualified to operate the centre.

- (3) Before authorising a centre, the Secretary of State must be satisfied—
 - (a) that the welfare of animals used in the collection of equine stem cells will be respected; and
 - (b) that the production process will produce a consistent, safe product.
- (4) Equine stem cells may only be collected under the responsibility of a veterinary surgeon.
- (5) The Secretary of State may suspend, vary or revoke an authorisation of an equine stem cell centre if—
 - (a) the holder no longer uses fit and proper processes;
 - (b) the premises in which the centre is being or is to be operated are not suitable;
 - (c) the equipment of the centre is not suitable; or
 - (d) the holder has not complied with these Regulations.
- (6) No person may operate an equine stem cell centre other than in accordance with such an authorisation.

Supply and administration of stem cells

- 31.**—(1) The operator of an equine stem cell centre may only collect equine stem cells.
- (2) The operator of an equine stem cell centre may not collect stem cells from embryonic tissues.
 - (3) No person other than a veterinary surgeon or someone acting under a veterinary surgeon's responsibility may administer any product grown from collected equine stem cells.
 - (4) No person may administer any product grown from collected equine stem cells to a food-producing horse.

Labelling

- 32.**—(1) The operator of an equine stem cell centre must ensure that every container used for the stem cell product is labelled with—
 - (a) the identification of the donor animal;
 - (b) the date of collection;
 - (c) the authorisation number of the equine stem cell centre;
 - (d) any necessary warnings;
 - (e) the expiry date.
- (2) The operator of an equine stem cell centre must ensure that no specific therapeutic indication is included on the label or on any information relating to the product.

Records

- 33.** The operator of an equine stem cell centre must, as soon as is reasonably practicable, record for each stem cell product—
 - (a) the identification of the donor animal;
 - (b) the date of collection;
 - (c) the veterinary surgeon under whose responsibility the stem cells were collected;
 - (d) the expiry date;
 - (e) the date the product was used or, if it was supplied to another veterinary surgeon, the name of that veterinary surgeon and the date it was supplied,and must keep the records for at least five years.

Inspection of premises

34. The Secretary of State must inspect the authorised premises, basing the frequency of the inspection on the risks associated with each premises' history and the nature of the products handled at the premises.

Offences

35. It is an offence to fail to comply with—

- (a) paragraph 4(3);
- (b) paragraph 11;
- (c) paragraph 14(4);
- (d) paragraph 16;
- (e) paragraph 17;
- (f) paragraph 18;
- (g) paragraph 20(6) or (7);
- (h) paragraph 21;
- (i) paragraph 22;
- (j) paragraph 23;
- (k) paragraph 25(5);
- (l) paragraph 26;
- (m) paragraph 27;
- (n) paragraph 28;
- (o) paragraph 30(4) or (6);
- (p) paragraph 31;
- (q) paragraph 32; or
- (r) paragraph 33.

SCHEDULE 3

Regulation 7

Classification and supply, wholesale dealers and sheep dip

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PART 1

Classification and supply of authorised veterinary medicinal products

Classification of veterinary medicinal products

- 1.—(1) There shall be the following categories of authorised veterinary medicinal products—
 - (a) Prescription Only Medicine–Veterinarian (abbreviated to POM-V);
 - (b) Prescription Only Medicine–Veterinarian, Pharmacist, Suitably Qualified Person (abbreviated to POM-VPS);
 - (c) Non-Food Animal–Veterinarian, Pharmacist, Suitably Qualified Person (abbreviated to NFA-VPS);
 - (d) Authorised Veterinary Medicine–General Sales List (abbreviated to AVm-GSL).
- (2) The Secretary of State must specify the classification of the veterinary medicinal product when granting the initial marketing authorisation.
- (3) The Secretary of State may change the classification after the marketing authorisation has been granted, either at the request of the marketing authorisation holder or in accordance with paragraph 37 of Schedule 1 (compulsory variation).

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(4) When granting the marketing authorisation the Secretary of State must classify the following as POM-V—

- (a) products containing narcotic or psychotropic substances;
- (b) products intended for administration following a diagnosis or clinical assessment by a veterinary surgeon.

(5) When granting the marketing authorisation the Secretary of State must classify the following as POM-V or POM-VPS—

- (a) products for food-producing animals;
- (b) products in respect of which special precautions must be taken in order to avoid any unnecessary risk to—
 - (i) the target species;
 - (ii) the person administering the products to the animal; and
 - (iii) the environment;
- (c) products that may cause effects that impede or interfere with subsequent diagnostic or therapeutic measures; and
- (d) new veterinary medicinal products containing an active substance that has not been included in an authorised veterinary medicinal product for five years.

(6) The requirement in sub-paragraph (5)(a) relating to veterinary medicinal products for food-producing animals does not apply if all the following criteria are met—

- (a) the administration of the veterinary medicinal product is restricted to formulations requiring no particular knowledge or skill in using the product;
- (b) the veterinary medicinal product does not present a direct or indirect risk, even if administered incorrectly, to the animal or animals treated, to the person administering the product or to the environment;
- (c) the summary of product characteristics of the veterinary medicinal product does not contain any warnings of potential serious side effects deriving from its correct use;
- (d) neither the veterinary medicinal product nor any other product containing the same active substance has previously been the subject of frequent serious adverse reaction reporting;
- (e) the summary of product characteristics does not refer to contra-indications related to other veterinary medicinal products commonly used without prescription;
- (f) the veterinary medicinal product is not subject to special storage conditions;
- (g) there is no risk for consumer safety as regards residues in food obtained from treated animals even where the veterinary medicinal products are used incorrectly; and
- (h) there is no risk to human or animal health as regards the development of resistance to antimicrobials or anthelmintic substances even where the veterinary medicinal products containing those substances are used incorrectly.

Wholesale supply of veterinary medicinal products

2.—(1) Only a holder of a marketing authorisation, the holder of a manufacturing authorisation or the holder of a wholesale dealer's authorisation granted by the Secretary of State may supply a veterinary medicinal product wholesale, or be in possession of it for that purpose.

(2) A person mentioned in sub-paragraph (1) may only supply a veterinary medicinal product if—

- (a) the authorisation in question relates to that product, and
- (b) the supply is to another person who is entitled to supply that product under these Regulations, either wholesale or retail.

(3) If the supply is to a suitably qualified person, it must be to the premises approved in accordance with paragraph 14.

(4) It is immaterial whether or not the supply is for profit.

(5) This paragraph does not apply in relation to a retailer of veterinary medicinal products who supplies another retailer with such products for the purpose of alleviating a temporary supply shortage that could be detrimental to animal welfare.

(6) A wholesale dealer may break open any package (other than the immediate packaging) of a veterinary medicinal product.

Retail supply of veterinary medicinal products

3.—(1) This paragraph applies in relation to retail supply of veterinary medicinal products.

(2) A veterinary medicinal product classified as POM-V may only be supplied by a veterinary surgeon or a pharmacist and must be supplied in accordance with a prescription from a veterinary surgeon.

(3) A veterinary medicinal product classified as POM-VPS may only be supplied by—

- (a) a veterinary surgeon;
- (b) a pharmacist; or
- (c) a suitably qualified person in accordance with paragraph 14,

and must be in accordance with a prescription from one of those persons.

(4) A veterinary medicinal product classified as NFA-VPS may be supplied without prescription, but may only be supplied by—

- (a) a veterinary surgeon;
- (b) a pharmacist; or
- (c) a suitably qualified person in accordance with paragraph 14.

(5) There are no restrictions on the supply of AVM-GSL products.

(6) In this paragraph—

- (a) “retail supply” means any supply other than to or from the holder of a wholesale dealer’s authorisation, and whether or not for payment; and
- (b) a person may supply a product irrespective of who owns it.

Prescriptions by a veterinary surgeon

4.—(1) A veterinary surgeon who prescribes a veterinary medicinal product classified as POM-V must first carry out a clinical assessment of the animal, and the animal must be under that veterinary surgeon’s care.

(2) This does not apply in relation to the administration of such a product to a wild animal where the administration is authorised by the Secretary of State.

Prescriptions

5.—(1) A prescription may be oral or written, but a veterinary medicinal product classified as POM-V or POM-VPS may only be supplied—

- (a) by the person who prescribed it;
- (b) under a written prescription that complies with paragraph 6; or

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- (c) (in the case of POM-VPS) by a suitably qualified person in accordance with paragraph 14(5).
- (2) A person supplying such a product under a written prescription—
 - (a) may only supply the product specified in that prescription;
 - (b) must take all reasonable steps to be satisfied that the prescription has been written and signed by a person entitled to prescribe the product; and
 - (c) must take all reasonable steps to ensure that it is supplied to the person named in the prescription.
- (3) No person may alter a written prescription unless authorised to do so by the person who signed it.

Written prescriptions

- 6.—(1) A written prescription must include—
 - (a) the name, address and telephone number of the person prescribing the product;
 - (b) the qualifications enabling the person to prescribe the product;
 - (c) the name and address of the owner or keeper;
 - (d) the identification (including the species) of the animal or group of animals to be treated;
 - (e) the premises at which the animals are kept if this is different from the address of the owner or keeper;
 - (f) the date of the prescription;
 - (g) the signature or other authentication of the person prescribing the product;
 - (h) the name and amount of the product prescribed;
 - (i) the dosage and administration instructions;
 - (j) any necessary warnings;
 - (k) the withdrawal period if relevant; and
 - (l) if it is prescribed under the cascade, a statement to that effect.
- (2) A written prescription for a controlled drug as specified in Schedules 2 to 4 of the Misuse of Drugs Regulations 2001⁽⁵⁾ is valid for 28 days.
- (3) A written prescription for any other drug is valid for six months or such shorter period as may be specified in the prescription.
- (4) If the prescription is repeatable it must specify the number of times the veterinary medicinal product may be supplied.

Duties when a product is prescribed or supplied

- 7. A person who prescribes a product classified as POM-V or POM-VPS, or supplies a product classified as NFA-VPS—
 - (a) before doing so, must be satisfied that the person who will use the product is competent to do so safely, and intends to use it for a purpose for which it is authorised;
 - (b) when doing so, must advise on its safe administration and on any warnings or contraindications on the label or package leaflet; and

(5) [S. I. 2001/3998](#); relevant amending instruments are [S. I. 2003/1432](#) and [2005/1653](#).

- (c) must not prescribe (or, in the case of a NFA-VPS product, supply) more than the minimum amount required for the treatment; but it is a defence to a charge of failing to comply with this paragraph to show that—
 - (i) the product prescribed or supplied was in a container specified in the marketing authorisation;
 - (ii) the manufacturer does not supply that veterinary medicinal product in a smaller container; and
 - (iii) the person prescribing or supplying is not a person authorised to break open the package before supply.

Supply by a veterinary surgeon from registered premises

8.—(1) A veterinary surgeon may only supply a veterinary medicinal product from practice premises registered with the Royal College of Veterinary Surgeons as veterinary practice premises at which veterinary medicinal products are stored or supplied.

(2) This paragraph does not apply in relation to a veterinary medicinal product classified as AVM-GSL.

(3) The Royal College of Veterinary Surgeons must, on request, supply the Secretary of State with a copy of the register of veterinary practice premises.

(4) The Secretary of State must, from time to time, inspect premises registered under sub-paragraph (1), basing the frequency of the inspection on the risks associated with each premises' history and the nature of the products handled at the premises.

(5) Where an inspection under sub-paragraph (4) reveals significant breaches of these Regulations the Secretary of State may require the Royal College of Veterinary Surgeons to remove the premises from the register maintained under sub-paragraph (1).

(6) Where the Secretary of State requires the removal of premises from the register the veterinary surgeon concerned may appeal using the procedure in regulation 30.

(7) Where premises have been removed from the register under sub-paragraph (5) they may not be re-registered without the approval of the Secretary of State.

(8) The Secretary of State may only grant approval under sub-paragraph (7) after a further inspection of the premises.

Supply by a veterinary surgeon

9.—(1) A veterinary surgeon supplying a veterinary medicinal product (other than one classified as AVM-GSL) must be present when it is handed over unless the veterinary surgeon—

- (a) authorises each transaction individually before the product is supplied; and
- (b) is satisfied that the person handing it over is competent to do so.

(2) A veterinary surgeon or a person acting under a veterinary surgeon's responsibility may open any package containing a veterinary medicinal product.

Supply by a pharmacist

10.—(1) A pharmacist may only supply a veterinary medicinal product classified as POM-V, POM-VPS or NFA-VPS from—

- (a) premises registered as a pharmacy with the General Pharmaceutical Council or with the Pharmaceutical Society of Northern Ireland;

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- (b) premises registered with the Royal College of Veterinary Surgeons as being premises from which a veterinary surgeon supplies veterinary medicinal products; or
 - (c) (in the case of a veterinary medicinal product classified as POM-VPS or NFA-VPS) from premises approved under paragraph 14.
- (2) A pharmacist supplying a veterinary medicinal product (other than one classified as AVM-GSL) must be present when it is handed over unless the pharmacist—
- (a) authorises each transaction individually before the product is supplied; and
 - (b) is satisfied that the person handing it over is competent to do so.
- (3) A pharmacist may supply any veterinary medicinal product prepared in a pharmacy in accordance with the prescriptions of a pharmacopoeia and intended to be supplied directly to the end-user.
- (4) A pharmacist may supply a homeopathic remedy prepared extemporaneously by a pharmacist in a registered pharmacy (as well as any other homeopathic remedy permitted to be supplied by a pharmacist under these Regulations) provided that it is prepared in accordance with paragraph 63 of Schedule 1 and intended to be supplied directly to the end user.
- (5) A pharmacist may break open any package containing a veterinary medicinal product for the purposes of supply other than the immediate packaging of an injectable product.

Supply of a veterinary medicinal product for incorporation into feedingstuffs

- 11.**—(1) This paragraph applies in relation to the supply of a veterinary medicinal product intended for incorporation into feedingstuffs.
- (2) The marketing authorisation holder, an authorised manufacturer of the product or an authorised wholesale dealer may only supply such a veterinary medicinal product to—
- (a) a veterinary surgeon, pharmacist or, in the case of a product classified as POM-VPS, a suitably qualified person;
 - (b) an approved premixture manufacturer; or
 - (c) an approved feedingstuffs manufacturer if the approval permits the rate of incorporation specified on the label of that veterinary medicinal product (if the manufacturer is the end-user the supply must be in accordance with a prescription).
- (3) A veterinary surgeon, pharmacist or, in the case of a product classified as POM-VPS, a suitably qualified person may only supply such a veterinary medicinal product to—
- (a) an approved premixture manufacturer; or
 - (b) an approved feedingstuffs manufacturer if the approval permits the rate of incorporation specified on the label of that veterinary medicinal product (if the manufacturer is the end user the supply must be in accordance with a prescription).
- (4) An approved premixture manufacturer or an approved feedingstuffs manufacturer may only supply such a veterinary medicinal product to another approved premixture manufacturer or approved feedingstuff manufacturer if the amount supplied does not exceed five per cent in terms of value of veterinary medicinal product incorporated annually by the person supplying the veterinary medicinal product.

Labelling at the time of retail supply

- 12.**—(1) If a veterinary medicinal product is supplied in a container specified in the marketing authorisation, it must not be supplied if any information on the outer packaging (or, if there is no outer packaging, the immediate packaging) is not clearly visible at the time of supply or has been changed in any way.

(2) Sub-paragraph (1) does not apply to a veterinary surgeon who amends a label, or a pharmacist who amends it in accordance with a prescription from a veterinary surgeon, provided that the unamended information remains clearly visible.

(3) If a veterinary medicinal product is supplied in a container other than that specified in the marketing authorisation, the person supplying the veterinary medicinal product must ensure that the container is suitably labelled and must supply sufficient written information (which may include a copy of the summary of product characteristics or the package leaflet) to enable the product to be used safely.

Supply of veterinary medicinal products for use under the cascade

13.—(1) A veterinary medicinal product supplied for administration under the cascade may only be supplied in accordance with a prescription from a veterinary surgeon.

(2) Unless the veterinary surgeon who prescribed the veterinary medicinal product both supplies the product and administers it to the animal in person, the person supplying it must label it (or ensure that it is labelled) with at least the following information—

- (a) the name and address of the pharmacy, veterinary surgery or approved premises supplying the veterinary medicinal product;
- (b) the name of the veterinary surgeon who has prescribed the product;
- (c) the name and address of the animal owner;
- (d) the identification (including the species) of the animal or group of animals;
- (e) the date of supply;
- (f) the expiry date of the product, if applicable;
- (g) the name or description of the product, which should include at least the name and quantity of active ingredients;
- (h) dosage and administration instructions;
- (i) any special storage precautions;
- (j) any necessary warnings for the user, target species, administration or disposal of the product;
- (k) the withdrawal period, if relevant; and
- (l) the words “Keep out of reach of children” and “For animal treatment only”.

Supply by a suitably qualified person

14.—(1) The Secretary of State may recognise bodies that are suitable to maintain a register for suitably qualified persons to prescribe and supply veterinary medicinal products classified as POM-VPS and NFA-VPS.

(2) In order to recognise such a body, the Secretary of State must be satisfied that the body—

- (a) has in place a system for ensuring that persons applying for registration have adequate training to act as a suitably qualified person under these Regulations;
- (b) has adequate standards in deciding whether or not to register someone as a suitably qualified person;
- (c) maintains a programme of continuing professional development for persons registered with it;
- (d) operates an adequate appeal system if it intends to refuse to register anyone with appropriate qualifications or to remove anyone from the register.

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(3) For the purposes of these Regulations, a suitably qualified person is a person who has passed examinations specified by such a body, and is registered with such a body as a suitably qualified person.

(4) A suitably qualified person may only supply a veterinary medicinal product classified as POM-VPS, NFA-VPS or AVM-GSL, and may only supply it from—

- (a) premises approved by the Secretary of State as being suitable for the storage and supply of veterinary medicinal products by a suitably qualified person;
- (b) premises registered as a pharmacy with the General Pharmaceutical Council or with the Pharmaceutical Society of Northern Ireland; or
- (c) practice premises registered under these Regulations as being premises from which a veterinary surgeon supplies veterinary medicinal products.

(5) A suitably qualified person who supplies a product classified as POM-VPS or NFA-VPS must either—

- (a) hand over or despatch the product personally;
- (b) ensure that, when the product is handed over or despatched, the suitably qualified person is in a position to intervene if necessary; or
- (c) check the product after it has been allocated for supply to a customer, and be satisfied that the person handing over or dispatching it is competent to do so.

(6) A suitably qualified person supplying products from premises approved under this regulation by the Secretary of State who considers that the premises no longer comply with the approval must notify the Secretary of State without unreasonable delay.

(7) The Secretary of State may issue a Code of Practice for suitably qualified persons, and a body recognised under this paragraph must take appropriate action in accordance with any disciplinary code that applies to that body if a suitably qualified person registered with it does not comply with the Code of Practice.

(8) The Secretary of State must publish a list of—

- (a) suitably qualified persons; and
- (b) the trading names and the addresses of premises approved under this paragraph(6).

(9) A suitably qualified person may break open any package (other than the immediate packaging) of a veterinary medicinal product.

(10) The Secretary of State may suspend or revoke the approval of approved premises on being satisfied that they are no longer suitable for the storage and supply of veterinary medicinal products.

Annual audit

15. At least once a year every person entitled to supply a veterinary medicinal product on prescription must carry out a detailed audit, and incoming and outgoing veterinary medicinal products must be reconciled with products currently held in stock, any discrepancies being recorded.

PART 2

Requirements for a wholesale dealer's authorisation

Application

16. An application for a wholesale dealer's authorisation must be made to the Secretary of State.

Time limits

17. The Secretary of State must process an application for a wholesale dealer's authorisation within 90 days of receiving it.

Granting the authorisation

18.—(1) The Secretary of State must grant a wholesale dealer's authorisation on being satisfied that this paragraph is complied with.

(2) The authorised site must be—

- (a) weatherproof;
- (b) secure and lockable;
- (c) clean; and
- (d) free from contaminants.

(3) If the veterinary medicinal products covered by the authorisation are subject to specific storage conditions, the site must be capable of fulfilling those requirements.

(4) The authorisation holder must—

- (a) have the services of technically competent staff; and
- (b) have an effective emergency recall plan.

The authorisation

19.—(1) The wholesale dealer's authorisation must specify—

- (a) the types of veterinary medicinal products and pharmaceutical forms that may be dealt in;
- (b) the place where they are to be stored;
- (c) the name and address of the person holding the authorisation;
- (d) the address of the premises to which it relates;
- (e) the name of the qualified person nominated to act under the Guidelines on Good Distribution Practice for Human Use⁽⁷⁾.

(2) It may cover more than one site.

(3) It lapses if the holder does not deal in veterinary medicinal products for five years.

(4) The holder of a wholesale dealer's authorisation must notify the Secretary of State, and if necessary apply for a variation of the authorisation, before making a material alteration to the premises or facilities used under the authorisation, or in the operations for which they are used.

(7) OJ No C 63, 1.3.94, p. 4.

Suspension, variation or revocation of the authorisation

20. The Secretary of State may suspend, vary or revoke a wholesale dealer's authorisation if the holder—

- (a) has not complied with these Regulations; or
- (b) no longer has suitable premises or equipment.

Duties on the holder of a wholesale dealer's authorisation

21. The holder of a wholesale dealer's authorisation must—

- (a) store veterinary medicinal products in accordance with the terms of the marketing authorisation for each product;
- (b) comply with the Guidelines on Good Distribution Practice of Medicinal Products for Human Use as if the veterinary medicinal products were authorised human medicinal products;
- (c) carry out a detailed stock audit at least once a year; and
- (d) supply information and samples to the Secretary of State on demand.

PART 3

Sheep dip

Supply of sheep dip

22.—(1) A person who supplies by retail sheep dip which contains a veterinary medicinal product must supply it in accordance with this paragraph.

(2) The supply must be to a person (or a person acting on that person's behalf) who is qualified to use it in accordance with paragraph 23.

(3) The supplier must make a record of that person's certificate or award number as soon as is reasonably practicable, and keep it for at least three years.

(4) If the active ingredient of the veterinary medicinal product is an organophosphorus compound, the supplier must give to the buyer—

- (a) a double-sided laminated notice meeting the specifications in the following subparagraph (unless the notice has been provided to the buyer within the previous twelve months and the supplier knows or has reasonable cause to believe that the buyer still has it available for use); and
- (b) two pairs of gloves either as described in the notice or providing demonstrably superior protection to the proposed user against exposure to the dip than would be provided by gloves as so described.

(5) The notice must be at least A4 size with a laminated transparent cover and must tell the user of the sheep dip—

- (a) to read and act in accordance with the label, including instructions on measuring and diluting concentrate;
- (b) that sheep dip is absorbed through the skin;
- (c) always to wear the recommended protective clothing, including gloves, and have spare protective clothing available;
- (d) always to wash protective clothing before taking it off; and

- (e) to direct any questions to the supplier or manufacturer.
- (6) The notice must contain a diagram showing recommended protective clothing.

Use of sheep dip

23.—(1) No person may use sheep dip which contains a veterinary medicinal product unless the person is acting under the supervision and in the presence of, a person who holds either—

- (a) a Certificate of Competence in the Safe Use of Sheep Dips showing that Parts 1 and 2 or units 1 and 2 of the assessment referred to in the Certificate have been satisfactorily completed; or
 - (b) NPTC Level 2 Award in the Safe Use of Sheep Dip (QCF).
- (2) The certificate must be issued—
- (a) in England, Wales and Northern Ireland; by—
 - (i) the National Proficiency Tests Council;
 - (ii) NPTC Part of the City & Guilds Group; or
 - (iii) City and Guilds NPTC;
 - (b) in Scotland, by one of those organisations or the Scottish Skills Testing Service.

Offences

24. It is an offence to fail to comply with—

- (a) paragraph 2;
- (b) paragraph 3;
- (c) paragraph 4(1);
- (d) paragraph 5;
- (e) paragraph 7;
- (f) paragraph 8(1);
- (g) paragraph 9(1);
- (h) paragraph 10;
- (i) paragraph 11;
- (j) paragraph 12(1) or (3);
- (k) paragraph 13;
- (l) paragraph 14(4), (5) or (6);
- (m) paragraph 15;
- (n) paragraph 19(4);
- (o) paragraph 21;
- (p) paragraph 22; or
- (q) paragraph 23(1).

SCHEDULE 4

Regulation 8

Administration of a veterinary medicinal product outside the terms of a marketing authorisation

CONTENTS

1. Administration under the cascade
2. Withdrawal periods
3. Administration to food-producing horses
4. Immunological products for serious epizootic disease
5. Immunological products for an imported or exported animal
6. Administration by veterinary surgeons from other member States
7. Treatment in exceptional circumstances
8. Administration of a homeopathic remedy
9. Administration under an animal test certificate
10. Offences
 - Signature
 - Explanatory Note

Administration under the cascade

1.—(1) A veterinary surgeon acting under this paragraph who prescribes a veterinary medicinal product may either administer it personally or may direct another person to do so under the responsibility of the veterinary surgeon.

(2) If there is no authorised veterinary medicinal product in the United Kingdom for a condition the veterinary surgeon responsible for the animal may, in particular to avoid unacceptable suffering, treat the animal concerned with the following (“the cascade”), cascaded in the following order—

- (a) a veterinary medicinal product authorised in the United Kingdom for use with another animal species, or for another condition in the same species; or
- (b) if there is no such product that is suitable, either—
 - (i) a human medicinal product authorised in the United Kingdom; or
 - (ii) a veterinary medicinal product not authorised in the United Kingdom but authorised in another member State for use with any animal species (in the case of a food-producing animal, it must be a food-producing species); or
- (c) if there is no such product that is suitable, a veterinary medicinal product prepared extemporaneously by a pharmacist, a veterinary surgeon or a person holding a manufacturing authorisation authorising the manufacture of that type of product.

(3) In the case of a veterinary medicinal product imported from another member State, if the veterinary surgeon has not obtained a certificate from the Secretary of State under regulation 25(5) permitting importation, the veterinary surgeon must obtain a certificate from the Secretary of State before administration.

(4) Any pharmacologically active substances included in a medicinal product administered to a food-producing animal under the cascade must be listed in Table 1 in the Annex to Commission Regulation (EU) No 37/2010.

Withdrawal periods

2.—(1) A veterinary surgeon prescribing or administering a veterinary medicinal product to a food-producing animal under the cascade must specify an appropriate withdrawal period.

(2) The withdrawal period must ensure that, if there is a maximum residue limit specified for the active substance in Table 1 in the Annex to Commission Regulation (EU) No 37/2010, the level of residue of the active substance does not exceed that limit.

(3) In any event, unless the Secretary of State has specified in writing a different withdrawal period for a particular veterinary medicinal product, the withdrawal period (irrespective of whether or not a maximum residue limit is specified in Table 1 in the Annex to Commission Regulation (EU) No 37/2010) must not be less than—

- (a) 7 days for eggs;
- (b) 7 days for milk;
- (c) 28 days for meat from poultry and mammals including fat and offal;
- (d) 500 degree days⁽⁸⁾ for fish meat.

Administration to food-producing horses

3.—(1) If there is no authorised veterinary medicinal product for a food-producing horse (as shown on its horse passport) and treatment under the cascade is unsuitable, substances may be administered in accordance with [Commission Regulation \(EC\) No 122/2013](#) (establishing, in accordance with [Directive 2001/82/EC](#) of the European Parliament and of the Council on the Community code relating to veterinary medicinal products, a list of substances essential for the treatment of equidae ⁽⁹⁾).

(2) The person administering the substance must comply with Article 3(2) of [Commission Regulation \(EC\) No 122/2013](#) (recording the details of the treatment in the animal's passport).

Immunological products for serious epizootic disease

4. In the event of serious epizootic diseases, the Secretary of State may permit in writing the administration of immunological veterinary medicinal products without a marketing authorisation, in the absence of a suitable medicinal product and after informing the Commission of the detailed conditions of use and may publicise any permit as the Secretary of State sees fit.

Immunological products for an imported or exported animal

5. If an animal is imported from, or exported to, a third country, the Secretary of State may permit the administration to that animal of an immunological veterinary medicinal product that is not covered by a marketing authorisation in the United Kingdom but is authorised under the legislation of the third country.

Administration by veterinary surgeons from other member States

6.—(1) Veterinary surgeons practising in another member State may bring into the United Kingdom and administer to animals small quantities of veterinary medicinal products that are not authorised for use in the United Kingdom if—

- (a) the quantity does not exceed the requirements for the treatment of specific animals;

⁽⁸⁾ The number of days of the withdrawal period is calculated by dividing 500 by the mean temperature of the water in degrees Celsius.

⁽⁹⁾ OJ No L42, 13.2.2013, p. 1.

Status: This is the original version (as it was originally made).

- (b) the product is authorised in the member State in which the veterinary surgeon is established;
 - (c) the product is transported by the veterinary surgeon in the original manufacturer's packaging;
 - (d) in the case of administration to food-producing animals, there is a veterinary medicinal product authorised in the United Kingdom that has the same qualitative and quantitative composition in terms of active substances;
 - (e) the veterinary surgeon is acquainted with the Code of Professional Conduct for veterinary surgeons issued by the Royal College of Veterinary Surgeons⁽¹⁰⁾.
- (2) The veterinary surgeon must only supply to the owner or keeper enough veterinary medicinal product to complete the treatment of animals concerned.
- (3) The veterinary surgeon must—
- (a) ensure that the withdrawal period specified on the label of the product is complied with, or the United Kingdom withdrawal period for the equivalent product authorised in the United Kingdom if this is longer than the one on the label; and
 - (b) keep detailed records of the animals treated, the diagnosis or clinical assessment, the products administered, the dosage administered, the duration of treatment and the withdrawal period applied, and must keep them in the United Kingdom for at least three years.
- (4) The overall range and quantity of veterinary medicinal products carried by the veterinary surgeon must not exceed that generally required for the daily needs of good veterinary practice.
- (5) This paragraph does not apply in relation to immunological veterinary medicinal products.

Treatment in exceptional circumstances

7.—(1) Where the health situation so requires, and where there is no suitable veterinary medicinal product available either as an authorised product or under the cascade, a veterinary surgeon may treat an animal with a medicinal product authorised in a third country; but a veterinary surgeon who has not obtained a certificate from the Secretary of State under regulation 25(5) permitting importation must obtain a certificate from the Secretary of State before treating the animal.

- (2) The certificate may be granted subject to any condition the Secretary of State thinks fit.

Administration of a homeopathic remedy

8.—(1) A registered homeopathic remedy or a homeopathic remedy prepared and supplied by a pharmacist under paragraph 10 of Schedule 3 may be administered to an animal by anyone, subject to any restrictions specified in its registration.

(2) A homeopathic remedy that was on the market before 1st January 1994 may be administered by anyone.

(3) A veterinary surgeon may administer, either personally or under the veterinary surgeon's responsibility—

- (a) a homeopathic remedy authorised for human use, or
- (b) a homeopathic remedy prepared extemporaneously by a veterinary surgeon, a pharmacist or a person holding a manufacturing authorisation authorising the manufacture of that type of product.

⁽¹⁰⁾ Published at <http://www.rcvs.org.uk/advice-and-guidance/code-of-professional-conduct-for-veterinary-surgeons/>.

Administration under an animal test certificate

9.—(1) A medicinal product may be administered in accordance with an animal test certificate granted for research purposes by the Secretary of State.

(2) An application for an animal test certificate may be refused if this is necessary for the protection of animal or public health or the environment, and the animal test certificate may be varied, suspended or revoked in the same way as a marketing authorisation.

(3) The holder of an animal test certificate may not supply a product for administration that is not within the terms of the animal test certificate.

(4) The holder of an animal test certificate test who becomes aware of any serious adverse reaction following the administration of a product under an animal test certificate must report the reaction to the Secretary of State within 15 days of learning of it.

Offences

10. It is an offence to fail to comply with—

- (a) paragraph 3(2);
- (b) paragraph 6; or
- (c) paragraph 9(3) or (4).

SCHEDULE 5

Regulation 14

Medicated feedingstuffs and specified feed additives

CONTENTS

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3. Enforcement of Regulation [\(EC\) No 1831/2003](#)
4. Enforcement of Regulation [\(EC\) No 882/2004](#)
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7. Approval of manufacturers and distributors of feedingstuffs containing veterinary medicinal products
8. Incorporation of a veterinary medicinal product into a premixture
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10. Incorporation of a veterinary medicinal product into feedingstuffs
11. Additional record keeping requirements relating to veterinary medicinal products
12. Labelling a premixture containing a veterinary medicinal product
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19. Prescriptions for feedingstuffs containing a veterinary medicinal product
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 27. Imports from third countries
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 29. Import for incorporation into premixture or feedingstuffs for export
 30. Animals on domestic premises
 31. Offences
- Signature
Explanatory Note

Scope and interpretation

1.—(1) This Schedule applies in relation to the following (referred to in this Schedule as “specified feed additives”) when used as feed additives—

- (a) coccidiostats;
- (b) histomonostats; and
- (c) all other zootechnical additives except—
 - (i) digestibility enhancers;
 - (ii) gut flora stabilisers; and
 - (iii) substances incorporated with the intention of favourably affecting the environment.

(2) It also applies in relation to the manufacture and placing on the market of feedingstuffs containing a veterinary medicinal product.

(3) In this Schedule—

“premixture” means a mixture of a veterinary medicinal product or a specified feed additive with feedingstuffs materials, intended for further mixing with feedingstuffs before being fed to animals;

“zootechnical additive” means any additive used to maintain animals in good health or favourably affect their performance.

Enforcement of Regulation (EC) No 178/2002

2.—(1) For the purposes of Regulation (EC) No 178/2002 (of the European Parliament and of the Council laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety⁽¹¹⁾) the competent authority is the Secretary of State.

(2) No person may fail to comply with any of the following provisions of that Regulation—

- (a) Article 11 (requirements relating to imports);
- (b) Article 12 (requirements relating to exports);
- (c) Article 15(1) (prohibition on the placing on the market or feeding unsafe feedingstuffs);

⁽¹¹⁾ OJ No L 31, 1.2.2002, p. 1.

- (d) Article 16 so far as it prohibits misleading labelling, advertising or presentation of feedingstuffs;
- (e) Article 18(2) and (3) (requirements of traceability) in so far as it relates to feed business operators; and
- (f) Article 20 (responsibilities of feed business operators).

Enforcement of Regulation (EC) No 1831/2003

3.—(1) For the purposes of Regulation (EC) No 1831/2003 (of the European Parliament and of the Council on additives for use in animal nutrition⁽¹²⁾) the competent authority is the Secretary of State.

(2) An authorisation under Article 3(2) of that Regulation must be in writing.

(3) No person may possess a specified feed additive, or a premixture or feedingstuffs containing a specified feed additive, unless the specified feed additive has been authorised under Regulation (EC) No 1831/2003 or is for export to a third country.

(4) No person may fail to comply with any of the following provisions of that Regulation—

- (a) Article 3(1) or Article 3(3) (the authorisation, conditions of use and labelling of specified feed additives);
- (b) Article 12(1) or (2) (conditions relating to specified feed additives);
- (c) Article 16(1) (labelling);
- (d) Article 16(3) (additional labelling requirement);
- (e) Article 16(4) (premixtures containing specified feed additives);
- (f) Article 16(5) (packaging).

Enforcement of Regulation (EC) No 882/2004

4. For the purposes of Regulation (EC) No. 882/2004 (of the European Parliament and the Council on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules⁽¹³⁾) the competent authority is the Secretary of State.

Enforcement of Regulation (EC) No 183/2005

5.—(1) For the purposes of Regulation (EC) No 183/2005 (of the European Parliament and of the Council laying down requirements for feed hygiene⁽¹⁴⁾) the competent authority is the Secretary of State.

(2) No person may fail to comply with any of the following provisions of that Regulation—

- (a) Article 5(2), (5) or (6) (specific obligations);
- (b) Article 6(1) as read with (2) and (3) (HACCP system);
- (c) Article 7(1) (documents concerning the HACCP system);
- (d) Article 9(2) (official controls, notification and registration);
- (e) Article 10(1) (approval of feed business establishments);
- (f) Article 11 (prohibition on operating without approval or registration);
- (g) Article 17(2) (exemption from on-site visits);

⁽¹²⁾ OJ No L268, 18.10.2003, p. 29.

⁽¹³⁾ Corrected version at OJ No L191, 28.5.2004, p. 1.

⁽¹⁴⁾ OJ No L35, 8.2.2005, p. 1

Status: This is the original version (as it was originally made).

- (h) Article 18(3) (declaration of compliance);
- (i) Article 23(1) (conditions relating to imports from third countries);
- (j) Article 25 (feedingstuffs produced for export to third countries).

(3) A manufacturer must ensure that, so far as is reasonably practicable, the active ingredient is evenly incorporated throughout the feedingstuffs.

(4) In the case of the refusal, suspension or revocation of an approval under the Regulation the appeals procedure relating to a manufacturing authorisation in regulation 30 applies.

Enforcement of Regulation (EC) No 767/2009

6. No person may contravene Article 8 of Regulation (EC) No 767/2009 of the European Parliament and of the Council in relation to feedingstuffs containing specified feed additives.

Approval of manufacturers and distributors of feedingstuffs containing veterinary medicinal products

7.—(1) For the purposes of Directive 90/167/EEC laying down the conditions governing the preparation, placing on the market and use of medicated feedingstuffs in the Community⁽¹⁵⁾ the competent authority is the Secretary of State.

(2) No person may incorporate a veterinary medicinal product into a premixture or feedingstuff, or act as a distributor of premixtures or feedingstuffs containing a veterinary medicinal product, without being approved to do so by the Secretary of State.

(3) The conditions which govern approval of feed business establishments under Regulation (EC) No 183/2005 laying down requirements for feed hygiene⁽¹⁶⁾ also govern approval of manufacturers and distributors under sub-paragraph (2).

(4) The Secretary of State shall conduct inspections of manufacturers and distributors approved under sub-paragraph (2) basing the frequency of inspection on the risks associated with each premises' history and the nature of the products handled at the premises.

(5) A manufacturer must ensure that, so far as is reasonably practical, the veterinary medicinal product is evenly incorporated throughout the feedingstuffs.

(6) The provisions of this paragraph do not apply in relation to any person breeding or selling ornamental fish not intended for human consumption provided that the person does not use more than a total of 1kg of veterinary medicinal product annually for that purpose.

(7) In the case of the refusal, suspension or revocation of an approval under this paragraph the appeals procedure relating to a manufacturing authorisation in regulation 30 applies.

Incorporation of a veterinary medicinal product into a premixture

8. Any person who incorporates a veterinary medicinal product into a premixture—
- (a) must do so in accordance with the summary of product characteristics, and must take account of any interactions listed there; and
 - (b) must ensure that the veterinary medicinal product does not contain the same active substance as any other additive.

⁽¹⁵⁾ OJ No L 92, 7.4.1990, p. 42.

⁽¹⁶⁾ OJ No L 35, 8.2.2005, p. 1.

Top dressing

9. No person may promote or label any veterinary medicinal product, or anything containing a veterinary medicinal product, as being suitable for top dressing (that is, sprinkling it on to feedingstuffs without thoroughly incorporating it) unless the summary of product characteristics specifically permits this use.

Incorporation of a veterinary medicinal product into feedingstuffs

10. Any person who incorporates a veterinary medicinal product (or a premixture containing a veterinary medicinal product) into feedingstuffs—

- (a) must do so in accordance with the summary of product characteristics, and must take account of any interactions listed there;
- (b) must ensure that the veterinary medicinal product does not contain the same active substance as any other additive;
- (c) must ensure that the veterinary medicinal product is incorporated in accordance with its marketing authorisation (unless it has been prescribed under the cascade) and the prescription;
- (d) must ensure that the daily dose of the veterinary medicinal product is contained in a quantity of medicated feedingstuffs corresponding to at least half the daily feedingstuffs ration of the animals treated or, in the case of ruminants, corresponding to at least half the daily requirements of non-mineral complementary feedingstuffs.

Additional record keeping requirements relating to veterinary medicinal products

11.—(1) Any person who—

- (a) incorporates a veterinary medicinal product into a premixture;
- (b) incorporates a premixture containing a veterinary medicinal product into feedingstuffs; or
- (c) incorporates a veterinary medicinal product into feedingstuffs,

must make a daily record of—

- (d) the types and quantities of all veterinary medicinal products (and specified feed additives, if any) and premixture used in the manufacturing process; and
- (e) the quantity of feedingstuffs and premixture containing veterinary medicinal product manufactured that day.

(2) An approved distributor must make a daily record of—

- (a) the types and quantities of all premixtures and feedingstuffs containing veterinary medicinal products bought and sold that day; and
- (b) the quantity held.

(3) A manufacturer and distributor must also record, as soon as reasonably practicable, for each consignment supplied—

- (a) the date of delivery;
- (b) the name and address of each consignee (or, in the case of a manufacturer supplying to a distributor, the name and address of the distributor);
- (c) the type of feedingstuffs or premixture supplied;
- (d) the quantity;
- (e) the type of veterinary medicinal product incorporated into the feedingstuffs; and
- (f) the expiry date.

- (4) Records must be kept for five years.

Labelling a premixture containing a veterinary medicinal product

12.—(1) A premixture containing a veterinary medicinal product must be clearly and legibly labelled with the following—

- (a) the words “MEDICATED PREMIXTURE” (or, if it is to be labelled as “complementary feedingstuffs” under legislation implementing Council [Directive 79/373/EEC](#) on the marketing of compound feedingstuffs⁽¹⁷⁾, “MEDICATED COMPLEMENTARY FEEDINGSTUFFS”) in upper case letters;
- (b) the proprietary name of the veterinary medicinal product and the authorisation number;
- (c) the name and amount of the active substance (mg/kg) in the premixture;
- (d) the range of acceptable inclusion rates of the premixture into the final feedingstuffs, the range of acceptable levels of the active ingredients in the final feedingstuffs and the words “refer to the prescription for the exact inclusion rate” or equivalent wording;
- (e) warnings and contra-indications;
- (f) the withdrawal period, and a statement that, if the prescription requires a longer withdrawal period, that is the one that applies;
- (g) the expiry date;
- (h) any special storage instructions;
- (i) where a prescription is required, a statement to this effect.

(2) If there is more than one veterinary medicinal product used, the longest withdrawal period must be shown on the label.

(3) If the premixture also contains a specified feed additive to which this Schedule applies it must also contain the information required under Article 16 of Regulation [\(EC\) No 1831/2003](#)⁽¹⁸⁾.

(4) No person may supply such a premixture unless it is labelled in accordance with this paragraph.

Labelling of feedingstuffs containing a specified feed additive

13. No person may contravene the labelling requirements of Article 15 and Article 17 of Regulation [\(EC\) No 767/2009](#) of the European Parliament and of the Council.

Labelling of feedingstuffs containing a veterinary medicinal product

14.—(1) Feedingstuffs containing a veterinary medicinal product must be clearly and legibly labelled with the following—

- (a) the words “MEDICATED COMPLETE FEED” in upper case letters, or where feedingstuffs are to be labelled as a complementary feedingstuff and intended to be fed to animals without further mixing with feed materials, the words “MEDICATED COMPLEMENTARY FEEDINGSTUFF”;
- (b) the proprietary name, authorisation number and inclusion rate (kg/tonne or mg/kg) of the veterinary medicinal product incorporated into the feedingstuffs;
- (c) the name and amount of the active substance (mg/kg) in the feedingstuffs;
- (d) the species of animal for which the feedingstuffs are intended;

⁽¹⁷⁾ OJ No L86, 6.4.1979, p. 30.

⁽¹⁸⁾ OJ No L268, 18.10.2003, p. 29. Regulation (EC) no 1831/2003 was last amended by Article 29 of Regulation [\(EC\) No 767/2009](#) (OJ No L229, 1.9.2009, p. 1.)

- (e) warnings and contra-indications;
 - (f) the withdrawal period, and a statement that, if the prescription requires a longer withdrawal period, that is the one that applies;
 - (g) the expiry date;
 - (h) any special storage instructions required by the marketing authorisation;
 - (i) a statement to the effect that the feedingstuffs must only be fed in accordance with its prescription;
 - (j) the name and approval number of the manufacturer or the distributor.
- (2) If there is more than one veterinary medicinal product used, the longest withdrawal period must be shown on the label.
- (3) If the feedingstuff also contains a specified feed additive to which this Schedule applies it must also contain the information required by Articles 15 and 17 of Regulation (EC) No 767/2009 of the European Parliament and of the Council.
- (4) No person may supply feedingstuffs unless they are labelled in accordance with this paragraph.

Supply of specified feed additives

- 15.**—(1) No person other than the person who manufactured a specified feed additive or an approved distributor may supply a specified feed additive.
- (2) The person who manufactured the specified feed additive may only supply it to—
- (a) an approved distributor;
 - (b) an approved premixture manufacturer or an approved complementary feedingstuffs manufacturer; or
 - (c) a feedingstuff manufacturer approved to mix a specified feed additive directly into feedingstuff.
- (3) An approved distributor may only supply it to—
- (a) another approved distributor;
 - (b) an approved premixture manufacturer or an approved complementary feedingstuffs manufacturer; or
 - (c) a feedingstuff manufacturer approved to mix a specified feed additive directly into feedingstuff.

Supply of premixture

- 16.**—(1) No person other than the person who manufactured a premixture or an approved distributor may supply a premixture.
- (2) The person who manufactured the premixture may only supply it to—
- (a) an approved distributor; or
 - (b) a feedingstuff manufacturer approved to incorporate that premixture.
- (3) An approved distributor may only supply it to—
- (a) another approved distributor; or
 - (b) a feedingstuff manufacturer approved to incorporate that premixture.

Supply of a complementary feedingstuff

17.—(1) No person other than—

- (a) the person who manufactured a complementary feedingstuff containing a specified feed additive; or
- (b) an approved distributor

may supply a complementary feedingstuff containing a specified feed additive.

(2) The person who manufactured such complementary feedingstuff may only supply it to—

- (a) an approved distributor; or
- (b) a feedingstuff manufacturer registered to incorporate that complementary feedingstuff or approved to incorporate a premixture.

(3) An approved distributor may only supply it to—

- (a) another approved distributor, or
- (b) a feedingstuff manufacturer registered to incorporate that complementary feedingstuff or approved to incorporate a premixture.

(4) In this paragraph “complementary feedingstuff” has the meaning given in Article 3 of Regulation EC No 767/2009.

Supply of feedingstuffs containing a veterinary medicinal product

18.—(1) No person other than the person who manufactured the feedingstuffs or an approved distributor may supply feedingstuffs containing a veterinary medicinal product.

(2) The person who manufactured the feedingstuff may only supply it to—

- (a) an approved distributor; or
- (b) a person who keeps animals for feeding to those animals.

(3) A distributor may only supply it to—

- (a) another approved distributor; or
- (b) a person who keeps animals for feeding to those animals.

(4) Supply to a person who keeps animals must be in accordance with a written prescription as specified in the following paragraph.

(5) If a prescription is for a period of longer than one month, the supplier may not provide more than one month’s supply at any one time.

(6) No manufacturer or distributor may supply a feedingstuff to anyone not specified in this paragraph, or otherwise than in accordance with this paragraph.

(7) The person supplying the feedingstuff must keep the prescription for five years.

Prescriptions for feedingstuffs containing a veterinary medicinal product

19.—(1) A prescription for feedingstuffs containing a veterinary medicinal product must contain the following—

- (a) the name and address of the person prescribing the product;
- (b) the qualifications enabling the person to prescribe the product;
- (c) the name and address of the keeper of the animals to be treated;
- (d) the species of animal, identification and number of the animals;
- (e) the premises at which the animals are kept if this is different from the address of the keeper;

- (f) the date of the prescription;
 - (g) the signature or other authentication of the person prescribing the product;
 - (h) the name and amount of the product prescribed;
 - (i) the dosage and administration instructions;
 - (j) any necessary warnings;
 - (k) the withdrawal period;
 - (l) the manufacturer or the distributor of the feedingstuffs (who must be approved for the purpose);
 - (m) if the validity exceeds one month, a statement that not more than 31 days' supply may be provided at any time;
 - (n) the name, type and quantity of feedingstuffs to be used;
 - (o) the inclusion rate of the veterinary medicinal product and the resulting inclusion rate of the active substance;
 - (p) any special instructions;
 - (q) the percentage of the prescribed feedingstuffs to be added to the daily ration; and
 - (r) if it is prescribed under the cascade, a statement to that effect.
- (2) It is valid for three months or such shorter period as may be specified in the prescription.
- (3) It must be sufficient for only one course of treatment.

Writing the prescription

- 20.**—(1) The person who writes the prescription must—
- (a) give a copy to the person incorporating the veterinary medicinal product into the feedingstuffs or to the distributor of the feedingstuffs;
 - (b) give one copy to the keeper of the animals to be treated;
 - (c) keep a copy.
- (2) The person must be satisfied that—
- (a) there is no undesirable interaction between the veterinary medicinal product and any feed additive used in the feedingstuffs; and
 - (b) the active substance of the veterinary medicinal product is not the same as an active substance in any feed additive used in the feedingstuffs.
- (3) If there is no suitable veterinary medicinal product a veterinary surgeon may—
- (a) prescribe a veterinary medicinal product authorised for another species and condition; or
 - (b) include more than one veterinary medicinal product for incorporation into the feedingstuff,
- provided that all veterinary medicinal products prescribed are authorised for inclusion in feedingstuffs.

Possession

- 21.**—(1) No person other than a person holding the appropriate approval under this Schedule may be in possession of any—
- (a) specified feed additive or veterinary medicinal product to which this Schedule applies;
 - (b) premixtures containing such an additive or a veterinary medicinal product; or

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(c) feedingstuffs or complementary feedingstuffs containing such an additive or a veterinary medicinal product unless supplied under these Regulations.

(2) No person other than a manufacturer or distributor may be in possession of feedingstuffs incorporating a veterinary medicinal product unless it has been supplied under a prescription.

Sampling and analysis

22.—(1) If any enforcement action is taken under this Schedule based on a sample, that sample must have been taken and analysed in accordance with Regulation (EC) No 152/2009 laying down methods of sampling and analysis for the official control of feedingstuffs⁽¹⁹⁾.

(2) Unless otherwise specified in the marketing authorisation, it is a defence if the active ingredient in the medicated feedingstuff sample is within the following tolerances—

Tolerance table for medicated feedingstuff

<i>Level of active ingredient specified on the label</i>	<i>Tolerance</i>
≤50 mg/kg	± 50%
>50 mg/kg ≤ 500 mg/kg	± 40%
>500 mg/kg ≤ 5g/kg	± 30%
>5g/kg ≤50g/kg	± 20%
>50g/kg	± 10%

(3) Unless otherwise specified in the Commission Regulation authorising the specified feed additive in question, it is a defence if the active ingredient of a specified feed additive in a feedingstuff sample is within the tolerances set out in Articles 11(5) and paragraph 2(e) of Annex IV to Regulation (EC) No 767/2009 of the European Parliament and of the Council.

Storage

23. No person may store a veterinary medicinal product intended for incorporation into feedingstuffs, or a premixture or feedingstuffs containing a veterinary medicinal product, except in—

- (a) a suitable storage area that is locked when not in use; or
- (b) a hermetic container designed to store those products.

Packages and other containers

24. No person may place on the market feedingstuffs containing a veterinary medicinal product except in packages or containers that are sealed in such a way that, when the package or container is opened, the seal is damaged.

Transport

25.—(1) No person may transport feedingstuffs by road tankers or in bulk unless the labelling requirements are set out in a document accompanying the feedingstuffs, and the transporter must hand over details when delivering the feedingstuffs unless these have already been provided to the purchaser.

⁽¹⁹⁾ OJ No L 54, 26.2.2009, p. 1.

(2) Any person transporting feedingstuffs containing veterinary medicinal products or specified feed additives in road tankers or similar containers must ensure that the vehicle or container is cleaned before any re-use if this is necessary to prevent undesirable interaction or contamination.

(3) In the case of feedingstuffs containing a veterinary medicinal product the transporter must ensure that the vehicle is accompanied by documentation stating this.

(4) Any person operating an undertaking transporting feedingstuffs containing veterinary medicinal products or specified feed additives must give written instructions to drivers on how to load and unload vehicles so as to avoid cross-contamination, and take reasonable steps to ensure that the driver complies with those instructions.

Possession, placing on the market and use of feedingstuffs

26.—(1) No person may possess, place on the market or feed to animals any feedingstuffs incorporating veterinary medicinal products or specified feed additives unless they have been incorporated in accordance with this Schedule.

(2) No person may feed to any animal, or buy, possess or supply for the purpose of feeding to any animal, any feedingstuff containing a veterinary medicinal product or specified feed additive unless—

- (a) that veterinary medicinal product or specified feed additive is authorised for that species of animal and for the purpose for which it is used; or
- (b) in the case of a veterinary medicinal product, it was prescribed for that animal.

(3) This paragraph does not apply in relation to feedingstuffs if the veterinary medicinal product has been incorporated in accordance with an animal test certificate or the feedingstuff has been imported in accordance with this Schedule.

Imports from third countries

27. No person may import a feedingstuff containing a veterinary medicinal product from a third country.

Trade between member States

28. No person may bring in from another member State a feedingstuff containing a veterinary medicinal product unless—

- (a) the feedingstuff has been manufactured in accordance with the provisions of Council [Directive 90/167/EEC](#) (laying down the conditions governing the preparation, placing on the market and use of medicated feedingstuffs in the Community⁽²⁰⁾) and Regulation [\(EC\) No 183/2005](#) of the European Parliament and of the Council laying down requirements for food hygiene; and
- (b) it only contains a veterinary medicinal product that has the same quantitative and qualitative composition as a veterinary medicinal product authorised in the United Kingdom.

Import for incorporation into premixture or feedingstuffs for export

29.—(1) A manufacturer of premixture or feedingstuffs who imports a veterinary medicinal product authorised in another member State or third country for the purpose of incorporating it into premixture or feedingstuffs for export does not commit an offence under regulation 43(i)

(20) OJ No L 92, 7.4.90, p. 42.

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(importation of an unauthorised veterinary medicinal product) or regulation 43(j) (possession of an unauthorised veterinary medicinal product).

(2) No person may place that premixture or feedingstuff on the market in the United Kingdom once the veterinary medicinal product has been incorporated into it.

Animals on domestic premises

30.—(1) The requirements of paragraph 7 (approval of manufacturers and distributors of feedingstuffs containing veterinary medicinal product) do not apply in relation to a person who incorporates a veterinary medicinal product into feedingstuffs in domestic premises for feeding, on those premises—

- (a) non-food-producing animals; or
- (b) food-producing animals provided that the animals or products from those animals are not sold or supplied commercially.

(2) Notwithstanding paragraphs 16 and 18 of this Schedule, a veterinary surgeon, a pharmacist or a suitably qualified person who is registered in accordance with paragraph 14 of Schedule 3 may be supplied with and may supply a premixture containing a veterinary medicinal product, or feedingstuffs containing a veterinary medicinal product, to such a producer.

(3) The requirement for a written prescription does not apply in relation to such supply, but the provisions of Schedule 3 relating to supply of a veterinary medicinal product apply in relation to the supply of premixture and feedingstuffs in the same way as they apply to a veterinary medicinal product.

Offences

31. It is an offence to fail to comply with—

- (a) paragraph 2(2);
- (b) paragraph 3(3) or (4);
- (c) paragraph 5(2) or (3);
- (d) paragraph 6;
- (e) paragraph 7(2) or (5);
- (f) paragraph 8;
- (g) paragraph 9;
- (h) paragraph 10;
- (i) paragraph 11;
- (j) paragraph 12(4);
- (k) paragraph 13;
- (l) paragraph 14(4);
- (m) paragraph 15;
- (n) paragraph 16;
- (o) paragraph 17;
- (p) paragraph 18;
- (q) paragraph 20(1) or (2);
- (r) paragraph 21;
- (s) paragraph 23;

- (t) paragraph 24;
- (u) paragraph 25;
- (v) paragraph 26(1) or (2);
- (w) paragraph 27;
- (x) paragraph 28; or
- (y) paragraph 29(2).

SCHEDULE 6

Regulation 15(4)

Exemptions for small pet animals

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 3. Manufacture
 4. Approval of the active substance
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 6. Labelling
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 8. Pack size
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- Signature
Explanatory Note

Animals to which this Schedule applies

1. This Schedule applies in relation to veterinary medicinal products intended solely for the following animals kept exclusively as a pet—

- (a) aquarium animals;
- (b) cage birds;
- (c) ferrets;
- (d) homing pigeons;
- (e) rabbits;
- (f) small rodents; and
- (g) terrarium animals.

Placing on the market, importing and administering the product

2. A veterinary medicinal product intended solely for an animal to which this Schedule applies may be placed on the market, imported or administered without a marketing authorisation if it complies with this Schedule.

Manufacture

3. The product must have been manufactured by—
- (a) the holder of a manufacturing authorisation if manufactured in the United Kingdom;
 - (b) the holder of a manufacturing authorisation issued under [Directive 2001/82/EC](#) if manufactured in another member State;
 - (c) in the case of Australia, Canada, New Zealand, or Switzerland, the holder of an authorisation from the competent authority permitting the manufacture of medicinal products;
 - (d) in the case of any other country, a manufacturer whose premises have been inspected and approved by an officer of the Secretary of State.

Approval of the active substance

4.—(1) The Secretary of State may approve an active substance for use in a veterinary medicinal product manufactured under this Schedule.

(2) The Secretary of State may not grant an approval if the active substance requires veterinary control.

(3) The approval must specify the species of animals for which it is approved, and may specify how the active substance or a product containing it is to be administered.

(4) The Secretary of State may suspend or revoke the approval (or limit it to a smaller number of species) if—

- (a) it is demonstrated that the substance requires veterinary control;
- (b) serious adverse reactions are reported making suspension or revocation necessary; or
- (c) it is demonstrated that the substance—
 - (i) is carcinogenic;
 - (ii) is genotoxic; or
 - (iii) shows developmental toxicity (including teratogenicity).

(5) The procedure for the refusal, suspension or revocation of an approval under this paragraph is the same as the procedure for a marketing authorisation.

The product

5.—(1) The active substance in the veterinary medicinal product must be approved under paragraph 4.

(2) The veterinary medicinal product must not be an antibiotic.

(3) It must not contain any narcotic or psychotropic substance.

(4) It must not be intended for treatments or pathological processes that require a precise prior diagnosis or the use of which may cause effects that impede or interfere with subsequent diagnostic or therapeutic measures.

Labelling

6.—(1) The product must be clearly labelled as being exempt from the requirements of these Regulations in relation to a marketing authorisation.

(2) The labelling must show the following—

- (a) the name of the veterinary product, including, if it is part of the name, its strength and pharmaceutical form;
- (b) the authorisation number of the manufacturer;
- (c) the name and strength of each active substance;
- (d) the route of administration;
- (e) the batch number;
- (f) the expiry date;
- (g) the words “For animal treatment only”;
- (h) the contents by weight, volume or number of dose units;
- (i) the name and address of the manufacturer or distributor;
- (j) the target species;
- (k) the words “Keep out of reach of children”;
- (l) storage instructions;
- (m) the shelf-life after the immediate packaging has been opened for the first time;
- (n) disposal advice;
- (o) full indications, including—
 - (i) therapeutic indications;
 - (ii) contra-indications;
 - (iii) interaction with other medicines and other forms of interaction; and
- (p) dosage instructions.

(3) If there is insufficient room on the label, the information may instead be in a package leaflet, but the leaflet must contain all the information in the preceding sub-paragraph other than the batch number and the expiry date, but the label on the product must contain at least the following—

- (a) the name of the veterinary medicinal product;
- (b) its active substance and its strength;
- (c) the route of administration;
- (d) the batch number;
- (e) the expiry date; and
- (f) the words “For animal treatment only”.

Administration

7. The method of administration must be oral or topical or (in the case of a product for fish) by addition to the water.

Pack size

8. The pack size must only be sufficient for a single course of treatment or, in the case of a veterinary medicinal product for aquarium fish, sufficient for a single course of treatment of no more than 7 administrations to an aquarium of 25,000 litres.

Adverse reactions

9.—(1) The manufacturer, importer or retailer of a veterinary medicinal product must—

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- (a) notify the Secretary of State within 15 days of learning of any serious adverse reactions (as defined in paragraph 57 of Schedule 1); and
 - (b) make a record of each adverse reaction and serious adverse reaction on becoming aware of it and keep it for three years.
- (2) It is an offence to fail to comply with this paragraph.

SCHEDULE 7

Regulation 16

Fees

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PART 1

Introduction

Interpretation

1. In this Schedule—

“national application” means an application for a marketing authorisation that does not involve another member State;

“pharmaceutical product” means any veterinary medicinal product other than an immunological product;

“simultaneous application” is an application in which, at the time an authorisation for a product is applied for, one or more additional applications are submitted for products that are identical to the first product except that—

- (a) in the case of an immunological product, they have a lesser number of antigens than the first product, but only contain antigens contained in the first product; and
- (b) in the case of a pharmaceutical product, they have different strengths of the active substance,

and, in the case of an application involving more than one member State, the additional applications do not include a member State that was not included in the first application.

Payment of fees

2. All fees under this Schedule are payable to the Secretary of State.

Time of payment

3. All fees are payable on invoice unless otherwise specified.

Multiple inspections

4. If a site, premises or establishment is inspected for more than one type of authorisation, approval or registration at the same time, the fee is the sum of —

- (a) the highest fee payable; and
- (b) 50% of each of the other fees.

Expenses for inspections outside the United Kingdom

5. Whenever premises outside the United Kingdom are inspected, the travel and subsistence costs of the inspectors and interpreters’ fees are payable in addition to the inspection fee specified.

Translation

- 6. All translation costs are charged additionally.

PART 2

Fees relating to marketing authorisations

Specified pharmaceutical applications

7. The following table sets out the fees relating to a pharmaceutical veterinary medicinal product for—

- (a) a national application for a marketing authorisation that is—
 - (i) a full application under Part 1 of Schedule 1;
 - (ii) a bibliographic application; or
 - (iii) an application based on pharmacological equivalence;
- (b) an application for a marketing authorisation using the decentralised procedure where the United Kingdom is a concerned member State;
- (c) an application for the mutual recognition of a product authorised in another member State.

<i>Application</i>	<i>Full national application under Part 1 of Schedule 1 (£)</i>	<i>Bibliographic national application (£)</i>	<i>Pharmacologically equivalent national application</i>		<i>Decentralised application where the UK is a concerned member State or recognition of a product authorised in another member State (£)</i>
			<i>Reference product authorised in UK (£)</i>	<i>Reference product not authorised in UK (£)</i>	
Base Fee:	13,530	12,115	7,195	9,220	6,515
Additional fee if any of the target species is a food-producing animal:	3,905	3,585	2,155	2,760	1,415
Additional fee for each active ingredient not previously included in a veterinary					

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<i>Application</i>	<i>Full national application under Part 1 of Schedule 1 (£)</i>	<i>Bibliographic national application (£)</i>	<i>Pharmacologically equivalent national application</i>		<i>Decentralised application where the UK is a concerned member State or recognition of a product authorised in another member State (£)</i>
			<i>Reference product authorised in UK (£)</i>	<i>Reference product not authorised in UK (£)</i>	
medicinal product authorised in the United Kingdom—					
food-producing animal:	7,465	6,595	5,885	7,495	2,630
non-food-producing animal:	6,525	5,855	5,590	7,155	2,295
Additional fee for each additional pack type:	740	740	605	775	330
Additional fee for each additional active ingredient (food-producing animal):	6,465	6,125	4,040	5,165	2,085
Additional fee for each additional active ingredient (non-food-producing animal):	4,310	4,105	3,235	4,135	1,475
Additional fee if there is more than one target species, for each additional species (food-producing animal):	3,970	3,565	2,425	3,100	1,280 Applies for a maximum of 2 additional species
Additional fee if there is more than one target species, for each additional species (non-food-producing animal):	2,495	2,090	1,550	1,980	805 Applies for a maximum of 2 additional species
Additional fee for each additional recommended route of administration	2,695	2,490	1,620	2,070	940

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<i>Application</i>	<i>Full national application under Part 1 of Schedule 1 (£)</i>	<i>Bibliographic national application (£)</i>	<i>Pharmacologically equivalent national application</i>		<i>Decentralised application where the UK is a concerned member State or recognition of a product authorised in another member State (£)</i>
			<i>Reference product authorised in UK (£)</i>	<i>Reference product not authorised in UK (£)</i>	
(food-producing animal):					
Additional fee for each additional recommended route of administration (non-food-producing animal):	1,215	1,010	740	945	405
Simultaneous applications: fee for each additional product in the application:	2,895	2,895	2,895	3,705	1,685

Decentralised pharmaceutical application where the United Kingdom is the reference member State

8. The fee for a decentralised application for a pharmaceutical product where the United Kingdom is the reference member State is the same as for a national application as set out in the table in paragraph 7, with the addition of the fees in the following table.

Fees for decentralised pharmaceutical application where the United Kingdom is the reference member State

<i>Application</i>	<i>Additional fee for a pharmacologically equivalent product (£)</i>	<i>Additional fee otherwise (£)</i>
Food-producing animal: one member State:	5,230	3,705
Non-food-producing animal: one member State:	3,985	3,220
Each additional member State:	530	530
Simultaneous application: fee for each additional product in the application:		
one member State:	6,670	6,670
each additional member State:	120	120

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Application for a marketing authorisation for an immunological or biosimilar product

9.—(1) The fee for a national application for a marketing authorisation relating to an immunological or biosimilar product, a decentralised application where the United Kingdom is the concerned member State or the mutual recognition of a product authorised in another member State is in accordance with the following table.

(2) In this paragraph a biosimilar application means an application made in accordance with Article 13(4) of [Directive 2001/82/EC](#) and a biosimilar product means a product which is the subject of such an application.

Fees for specified immunological and biosimilar applications

<i>Application</i>	<i>National application for a marketing authorisation (£)</i>	<i>Decentralised application where the UK is a concerned member State or recognition of a product authorised in another member State (£)</i>
1. Immunological or biosimilar product other than in paragraph 2 below: Base fee:	11,775	5,785
The following fees are in addition to the base fee—		
Additional fee for each active ingredient not previously included in a veterinary medicinal product authorised in the United Kingdom, and for each new combination of active ingredients:	7,405	2,490
Additional fee for each adjuvant or preservative not previously included in a veterinary medicinal product authorised in the United Kingdom and for each new combination of adjuvants or preservatives:	1,345	675
More than one antigenic component – fee for each additional component:	1,350	405
More than one species – fee for each additional species:	5,380	1,615
		Applies for a maximum of 2 additional species
More than one route of administration – fee for each additional route of administration:	5,380	1,615
Simultaneous application - fee for each additional product in the application:	2,895	1,685
2. Immunological or product that is identical to a product already authorised in the United Kingdom but with a lesser number of antigens and that only contains antigens contained in that product:	10,430	5,380

Decentralised immunological application where the United Kingdom is the reference member State

10. The fee for a decentralised application for a marketing authorisation for an immunological product where the United Kingdom is the reference member State is the same as for a national application set out in the previous table, with the addition of the fees in the following table—

Fees for decentralised immunological application where the United Kingdom is the reference member State

<i>Application</i>	<i>Additional fee (£)</i>
One member State:	3,470
Each additional member State:	530
Simultaneous applications: fee for each additional product in the application:	
one member State:	6,670
each additional member State:	120

Applications for a marketing authorisation using data already assessed

11. The fees for applications for marketing authorisations using identical data submitted simultaneously or on the basis of information provided under Article 13(c) of [Directive 2001/82/EC](#) are in accordance with the following table.

Fees for a marketing authorisation using data already assessed

<i>Application</i>	<i>Fee (£) per authorisation</i>
Decentralised application where the United Kingdom is the reference member State—	
one member State:	4,165
each additional member State:	530
Any other application:	945

Application for an exceptional marketing authorisation (pharmaceutical)

12. The fee for an application for an exceptional marketing authorisation for a pharmaceutical product is in accordance with the following table.

Fees for an exceptional marketing authorisation for a pharmaceutical product

<i>Application</i>	<i>Provisional (£)</i>	<i>Limited (£)</i>
Base Fee:	12,015	6,765
The following fees are in addition to the base fee—		
Additional fee if any of the target species is a food-producing animal:	3,905	1,952

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<i>Application</i>	<i>Provisional (£)</i>	<i>Limited (£)</i>
Additional fee for each active ingredient not previously included in a veterinary medicinal product authorised in the United Kingdom—		
food-producing animal:	5,850	3,732
non-food-producing animal:	4,910	3,262
Additional fee for each additional pack type:	710	370
Additional fee for each additional active ingredient (food-producing animal):	5,955	3,232
Additional fee for each additional active ingredient (non-food-producing animal):	3,800	2,155
Additional fee if there is more than one target species, for each additional species (food-producing animal):	2,965	1,985
Additional fee if there is more than one target species, for each additional species (non-food-producing animal):	1,485	1,247
Additional fee for each additional recommended route of administration (food-producing animal):	2,185	1,347
Additional fee for each additional recommended route of administration (non-food-producing animal):	710	608
Simultaneous applications— fee for each additional product in the application:	2,895	1,447

Fees for an application for an exceptional marketing authorisation (immunological)

13. The fee for an application for an exceptional marketing authorisation for an immunological product is in accordance with the following table.

Fees for an exceptional marketing authorisation for an immunological product

<i>Application</i>	<i>Provisional (£)</i>	<i>Limited (£)</i>
Base fee:	10,810	5,887

The following fees are in addition to the base fee—

Additional fee for each active ingredient not previously included in a veterinary medicinal product authorised in the United Kingdom, and for each new combination of active ingredients:	5,650	3,702
Additional fee for each adjuvant or preservative not previously included in a veterinary medicinal product authorised in the United Kingdom and for each new combination of adjuvants or preservatives:	1,350	672

<i>Application</i>	<i>Provisional (£)</i>	<i>Limited (£)</i>
More than one antigenic component – fee for each additional component:	1,190	675
More than one species – fee for each additional species:	4,060	2,690
More than one route of administration – fee for each additional route of administration:	4,060	2,690
Simultaneous application - fee for each additional product in the application:	2,895	1,447

Fee for the conversion from an exceptional to a full marketing authorisation

14. The fee for the conversion of an exceptional marketing authorisation to a full marketing authorisation is £3,000.

Application for a marketing authorisation relating to a parallel import

15. The fee for a marketing authorisation for a parallel import is in accordance with the following table.

Parallel imports

<i>Application</i>	<i>Fee (£)</i>
Application where the imported product has been authorised in accordance with the mutual recognition procedure or decentralised procedure, and the United Kingdom is included in these procedures— import from one or more member States:	1,755
Application to add an additional member State after the marketing authorisation has been granted – fee for each member State:	455
Application where the imported product has not been authorised in accordance with the mutual recognition procedure or the decentralised procedure but where the imported product originates from the same manufacturing site as the product authorised in the United Kingdom to which the imported product is considered to be essentially similar:	2,130
Any other application – fee for each member State from which the product is imported:	4,710

Application to change the distribution category of a product authorised through the centralised procedure

16. The fee to change the distribution category of a product authorised through the centralised procedure is £3,135.

Status: This is the original version (as it was originally made).

Application for a variation to a marketing authorisation dealt with under national or mutual recognition variation procedures.

17.—(1) This paragraph applies in relation to an application for a variation to one or more marketing authorisations except where paragraph 18, 19 or 21 applies.

(2) The fees for the variations to which this paragraph applies are set out in the following table.

(3) Where applications are made at the same time seeking an identical change to the terms of more than one marketing authorisation, and those applications are based on identical data, fees are payable as for a grouped variation.

(4) References in this paragraph to a grouped variation being “led” by a particular type of variation indicate that the principal variation in that group is a variation of that type.

<i>Type of variation</i>	<i>National</i>	<i>UK is the reference member State</i>	<i>UK is a concerned member State</i>
Single variations; one change for each product			
Extension:			
Change of strength or potency or the addition of a new strength or potency:	6,670		1,998
Change of pharmaceutical form or the addition of a new pharmaceutical form:	8,415		2,301
Change of route of administration, or the addition of a new one, of—			
(i) an immunological product, or a pharmaceutical product for a food-producing animal:	5,390		1,737
(ii) a pharmaceutical product for a food-producing animal:	7,135		2,058
Change or addition of a food producing target species:	9,620		2,547
Change of active substance, including:	8,415		2,301
use of a different salt, ester, complex or derivative of the same therapeutic moiety:			
use of a different biologically active substance with a slightly different molecular structure:			
modification of the vector used to produce the antigen or the source material, including a new master cell bank from a different source:			

<i>Type of variation</i>	<i>National</i>	<i>UK is the reference member State</i>	<i>UK is a concerned member State</i>
use of a new ligand or coupling mechanism for a radiopharmaceutical:			
change of the extraction solvent or change of the ratio of herbal drug to herbal drug preparation:			
Change of bioavailability:	8,415		2,301
Change of pharmacokinetics:	8,415		2,301
Simultaneous application: fee for each additional product in the application:	2,895		1,011
Type II:	2,895	6,030	1,872
Type IB:	885	1,325	531
Type IA:	455	685	273

Grouped variations**Extension-led:**

The fee for an application for an extension-led grouped variation is the fee for that extension as specified above plus —

- (a) if there is one variation in addition to the extension, the fee for that variation as specified above; or
- (b) if there is more than one variation in addition to the extension, the fee that would be payable for a grouped variation of that type as specified below.

Type II led:

For the first nine changes:	6,280	12,060	3,768
For each subsequent group of up to ten changes:	4,500	4,500	2,700

Type IB led:

For the first nine changes:	1,770	2,650	1,062
For each subsequent group of up to ten changes:	4,500	4,500	2,700

Type IA led:

For the first nine changes:	885	1,325	531
For each subsequent group of up to ten changes:	4,500	4,500	2,700

Application for a variation to a marketing authorisation dealt with under worksharing procedures

18.—(1) This paragraph applies in relation to an application for a variation to a marketing authorisation dealt with in accordance with worksharing procedures as set out in Article 20 of Commission Regulation (EC) No 1234/.

Status: This is the original version (as it was originally made).

(2) The fee for a worksharing application, involving marketing authorisations obtained by a national procedure in the United Kingdom only, is the fee specified in the following table in the column headed “UK Only”.

(3) The fee for a worksharing application, involving marketing authorisations obtained through a national procedure in the United Kingdom and any other member State, is specified in the following table by reference to the United Kingdom’s role in the procedure, as “UK Reference Authority”, “UK Co-Reference Authority” or “Other”.

(4) The fee for a worksharing application, involving at least one marketing authorisation obtained through the mutual recognition or decentralised procedure, is specified in the following table by reference to the United Kingdom’s role in the procedure, as “UK Reference Authority”, “UK Co-Reference Authority” or “UK Concerned member State”.

(5) The fee for any kind of variation where the Agency co-ordinates worksharing is £455 for each marketing authorisation.

<i>Type of application</i>	<i>UK Only</i>	<i>Where the application involves nationally authorised products in more than one member State</i>			<i>Application involves mutually recognised products</i>		
	UK Only	UK Reference Authority	UK Co-Reference Authority	Other	UK Reference Authority	UK Co-Reference Authority	UK Concerned member State
<i>Worksharing applications</i>							
The following fees apply for each change to each product:							
Type II							
For the first nine changes:	6,240	12,060	7,485	12,060	13,265	6,745	3,372
For each subsequent group of up to ten changes:	4,500	4,500	4,500	4,500	4,500	4,500	2,700
Type IB							
For the first nine changes:	1,770	2,650	2,120	2,650	2,915	1,905	954
For each subsequent group of up to ten changes:	4,500	4,500	4,500	4,500	4,500	4,500	2,700

Application for an extension dealt with under the decentralised procedure where the United Kingdom is the reference member State

19. The fee for a decentralised application for an extension where the United Kingdom is the reference member State is the same as for a national application as set out in the table in paragraph 17, with the addition of the supplementary fees in the following table (save that, where the application is for the addition of more than one species, only one supplementary fee applies).

**Decentralised application for an extension where
the United Kingdom is the reference member State**

<i>Application</i>	<i>Supplementary fee (£)</i>
Pharmaceutical product for a food-producing animal – one member State:	3,705
Pharmaceutical product for a non-food-producing animal – one member State:	3,220
Immunological product – one member State:	3,460
Each additional member State:	530
Simultaneous application: fee for each additional product in the application:	
one member State:	6,670
each additional member State:	120

Provision of information relating to the recognition of a United Kingdom marketing authorisation or an extension

20.—(1) Where an application is made for the Secretary of State to provide information to other member States to enable them to recognise a marketing authorisation already granted by the United Kingdom the following fees are payable.

(2) Those fees also apply where a marketing authorisation has been granted in more than one member State, the holder applies for an extension for that marketing authorisation and the United Kingdom acts as reference member State.

(3) Where a valid application to provide information to another member State is received within six months of the original grant of the marketing authorisation, or where the Secretary of State has already provided the information to a member State, and a further valid application is made to provide the information to an additional member State within six months of the date the last information was provided, the fees are—

<i>Type of application</i>	<i>Fee for^a pharmacologically equivalent product^(a)</i>	<i>Fee (other products) (£)</i>
Pharmaceutical product for a food-producing animal – one member State:	3,940	2,440
Pharmaceutical product for a non-food-producing animal one member State:	2,645	1,895
Immunological product – one member State:	2,130	2,130
Each additional member State:	535	535

(a) This fee is payable if the application for the marketing authorisation was on the basis that the product was pharmacologically equivalent to another veterinary medicinal product.

(4) Where the information to be provided relates to a product granted a marketing authorisation using identical data submitted simultaneously or on the basis of information provided under Article 13(c) of [Directive 2001/82/EC](#) the fees are—

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<i>Application</i>	<i>Fee (£)</i>
Provision of information to—	
one member State:	4,165
each additional member State:	530

(5) In any other case the fees are—

<i>Type of application</i>	<i>Fee for a pharmacologically equivalent product (£)^(a)</i>	<i>Fee (other products) (£)</i>
Pharmaceutical product for a food-producing animal – one member State:	12,015	10,515
Pharmaceutical product for a non-food-producing animal – one member State:	8,115	7,365
Immunological product – one member State:	8,940	8,940
Each additional member State:	535	535

(a) This fee is payable if the application for the marketing authorisation was on the basis that the product was pharmacologically equivalent to another veterinary medicinal product.

(6) In the case of simultaneous applications, the above fees are payable for each additional product in the application for one member State, with a fee of £115 for each additional product for each additional member State.

Exception for a variation relating to animal testing

21. If the only purpose of a variation is to remove animal testing or to reduce the numbers of animals used in testing, no fee is payable for the variation in the case of a national authorisation, and the United Kingdom element of the fee for the variation is not payable for an authorisation obtained through the mutual recognition procedure or the decentralised procedure.

Application for the renewal of a national marketing authorisation

22.—(1) The fee for an application for the renewal of a marketing authorisation is £1,360.

(2) The fee for the first reassessment of an exceptional marketing authorisation is £305, and the fee for each subsequent reassessment is £1,360.

Application for the renewal of a marketing authorisation obtained through mutual recognition or the decentralised procedure

23. The fee for an application for the renewal of a marketing authorisation obtained through mutual recognition or the decentralised procedure is —

- (a) £1,835 if the United Kingdom is the reference member State; and
- (b) £1,225 if the United Kingdom is a concerned member State.

Registration of a homeopathic remedy

24. The fee for an application for the registration of a homeopathic remedy is in accordance with the following table.

Fee for the registration of a homeopathic remedy

<i>Type of application</i>	<i>Fees (£)</i>
If all stocks and the formulation have already been assessed by the Secretary of State—	
not more than five stocks:	160
more than five stocks:	375
If either all the stocks have already been assessed by the Secretary of State but there is a new formulation, or if the formulation has already been assessed by the Secretary of State but one or more of the stocks have not been already assessed—	
not more than five stocks:	455
more than five stocks:	665
If the formulation and at least one of the stocks has not already been assessed by the Secretary of State—	
not more than five stocks:	760
more than five stocks:	985
If the product is already authorised for human use in the United Kingdom, or for human or veterinary use in the United Kingdom or in another member State—	
not more than five stocks:	160
more than five stocks:	375

Renewal of a homeopathic remedy

25. The fee for the renewal of a homeopathic remedy is £320.

Annual fees for marketing authorisations

26.—(1) Within 30 days of receiving a written demand from the Secretary of State, a holder of a marketing authorisation must provide the Secretary of State with a statement of turnover for the previous calendar year.

(2) The annual fee, rounded to the next £1, is—

$$£ \frac{0.67T}{100} + £230n$$

where—

(a) *T* is the annual turnover in the previous calendar year;

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(b) and n is the number of active marketing authorisations held at any time during the previous calendar year.

(3) In the case of an authorisation holder with a turnover relating to all marketing authorisations held of less than £230,000, the annual fee, rounded to the next £1 is—

$$£ \frac{0.67T}{100} + £200n$$

where—

(a) T is the annual turnover in the previous calendar year;

(b) and n is the number of active marketing authorisations held at any time during the previous calendar year.

(4) In this paragraph—

“turnover” means the sales value at manufacturers’ prices of all authorised veterinary medicinal products sold or supplied in the United Kingdom;

“manufacturers’ prices” means the prices charged (excluding value added tax) for authorised products by manufacturers to wholesalers, except to the extent that—

(a) the products are supplied by manufacturers direct to retailers, in which case it means the prices charged for the products by the manufacturers to the retailers reduced by such sum as, in the opinion of the Secretary of State, represents the difference between the prices paid by the retailers and those which could be expected to be charged by the manufacturers to wholesalers according to the practice prevailing during the period in question with regard to such products;

(b) a marketing authorisation holder sells or supplies products that the marketing authorisation holder has neither manufactured nor obtained from the manufacturer, in which case it means the prices paid by the marketing authorisation holder for those products.

Auditor’s certificate

27.—(1) The Secretary of State may at any time require an audit certificate in support of a statement of turnover.

(2) If the holder of the marketing authorisation does not provide an audit certificate before the date stipulated in the demand, an additional fee is payable for that year of £11,300 plus an additional £2,245 in respect of each marketing authorisation held.

(3) If the Secretary of State is not satisfied that the audit certificate provides sufficient assurance that the figures fairly present the financial records of the company, the Secretary of State may require the marketing authorisation holder to produce a further certificate and specify what further assurances are needed; and if these are not provided by the required date, the additional fee specified in sub-paragraph (2) is payable.

(4) Nothing in this paragraph limits the powers of an inspector to examine financial records.

PART 3

Fees payable by manufacturers

Application for a manufacturing authorisation

28. The fee for an application for a manufacturing authorisation for a veterinary medicinal product is—

- (a) £3,040; or
- (b) £530 if the authorisation only covers veterinary medicinal products manufactured under Schedule 6 (exemptions for small pet animals).

Application for a variation of a manufacturing authorisation

29. The fee for an application for the variation of a manufacturing authorisation is—

- (a) £636 if the variation requires scientific or pharmaceutical assessment;
- (b) £443 if the variation only involves a change of ownership;
- (c) £210 if the authorisation only covers veterinary medicinal products manufactured under Schedule 6 (exemptions for small pet animals); and
- (d) otherwise £350.

Application for an authorisation to manufacture an autogenous vaccine or a product for administration under the cascade

30.—(1) The fee for an application for a standard authorisation to manufacture an autogenous vaccine or a veterinary medicinal product for administration under the cascade is—

- (a) £3,435 for a site in the United Kingdom;
- (b) £3,270 for a site outside the United Kingdom.

(2) The fee for each inspection after a standard authorisation has been granted is (in each case) the same as the fee specified in paragraph (1).

(3) In the case of an application for an individual authorisation to manufacture a single batch of autogenous vaccine, or a single batch of veterinary medicinal product for administration under the cascade the fee is £1,635.

(4) The fee to vary an authorisation is £305 if no further inspection is required, and otherwise is the full application fee.

Annual fees

31.—(1) An annual fee of £550 is payable in respect of each manufacturing authorisation held (other than as specified in this paragraph).

(2) The annual fee for a manufacturing authorisation for an autogenous vaccine or a veterinary medicinal product for administration under the cascade is 0.67% of the turnover in the previous calendar year rounded to the next £1, with a minimum fee of £10.

(3) There is no annual fee for a manufacturing authorisation for a veterinary medicinal product manufactured in accordance with Schedule 6 for small pet animals.

(4) In this paragraph “turnover” means the sales value at manufacturers’ prices net of value added tax of all authorised veterinary medicinal products sold or supplied in the United Kingdom.

Status: This is the original version (as it was originally made).

Site inspections – type of site

32. For the purposes of deciding the fee for a site inspection—

“super site” is a site at which 250 or more relevant persons are employed;

“major site” is a site at which 60 or more, but fewer than 250, relevant persons are employed;

“standard site” is a site at which 10 or more, but fewer than 60 relevant persons are employed;

“minor site” is a site at which fewer than 10 relevant persons are employed;

“relevant person” means a person employed on the premises and systems inspected.

Inspection of a site where immunological veterinary medicinal products are manufactured

33. The fees for the inspection of a site where immunological veterinary medicinal products are manufactured are in accordance with the following table.

Sites where immunological veterinary medicinal products are manufactured

<i>Type of site</i>	<i>Fee (£)</i>	
	<i>United Kingdom site</i>	<i>Site outside United Kingdom</i>
Super site	24,071	22,867
Major site	16,785	15,946
Standard site	6,661	6,327
Minor site	4,757	4,519

Inspection of a site where sterile veterinary medicinal products are manufactured

34. The following fees are payable for the inspection of a site where no immunological veterinary medicinal products are manufactured, but where sterile products are manufactured.

Sites where sterile veterinary medicinal products are manufactured

<i>Type of site</i>	<i>Fee (£)</i>	
	<i>United Kingdom site</i>	<i>Site outside the United Kingdom</i>
Super site	23,324	22,157
Major site	13,010	12,359
Standard site	8,244	7,832
Minor site	5,022	4,770

Inspection of a site where no immunological or sterile veterinary medicinal products are manufactured

35. The following fees are payable for the inspection of a site where only non-immunological and non-sterile veterinary medicinal products are manufactured—

Site where no immunological or sterile veterinary medicinal products are manufactured

<i>Type of site</i>	<i>Fee (£)</i>	
	<i>United Kingdom site</i>	<i>Site outside the United Kingdom</i>
Super site	14,180	13,471
Major site	8,325	7,909
Standard site	6,854	6,511
Minor site	3,789	3,600
If the site is only involved in the manufacture of veterinary medicinal products authorised under Schedule 6 (exemptions for small pet animals—		
Standard site	5,055	4,802
Minor site	2,728	2,592

Inspection of a site where veterinary medicinal products are assembled

36. The following fees are payable for the inspection of a site where the only manufacturing process in relation to veterinary medicinal products is their assembly after the product has been put into its immediate container.

Site where medicinal products are assembled

<i>Type of site</i>	<i>Fee (£)</i>	
	<i>United Kingdom site</i>	<i>Site outside the United Kingdom</i>
Super site	11,025	10,474
Major site	5,949	5,652
Standard site	4,917	4,671
Minor site	2,035	1,933

Test sites

37. The fee for the inspection of a test site is £3,344, or £3,177 for a site outside the United Kingdom.

Animal blood bank or equine stem cell centre authorisations

38.—(1) The fee for an authorisation to operate a blood bank is—

- (a) on a first inspection £3,113; and
- (b) on each subsequent inspection—
 - (i) £3,113 for a site in the United Kingdom; and
 - (ii) £2,966 for a site outside the United Kingdom.

(2) The fee for an authorisation to operate an equine stem cell centre is £3,427, and £3,092 for each subsequent inspection.

(3) The fee for a variation to an authorisation to operate a blood-bank or equine stem cell centre is £320.

PART 4

Fees relating to a wholesale dealer's authorisation

Application for a wholesale dealer's authorisation

39.—(1) The fee for an application for a wholesale dealer's authorisation is—

- (a) £1,745;
- (b) £785 if the application is accompanied by an estimate that the first year's turnover will be less than £35,000; or
- (c) £785 if the authorisation only relates to products classified as AVM-GSL, homeopathic remedies, or products authorised under Schedule 6 (exemptions for small pet animals).

(2) An applicant who has paid a fee of £785 on the grounds of turnover must send a declaration of turnover for the first year of trading on the anniversary of the grant of the authorisation, and if the figure is more than £35,000 must pay the balance of £960 within 30 days.

(3) If the applicant paid £1,745 but the turnover for the first year of trading was lower than £35,000, if the applicant sends a declaration certifying the turnover, the Secretary of State must refund the excess.

(4) Nothing in this paragraph limits the powers of an inspector to examine financial records.

(5) In this paragraph "turnover" means the sales value net of value added tax of all veterinary medicinal products (whether or not authorised for use in the United Kingdom) sold by way of wholesale dealing by the holder in the United Kingdom.

Variation of a wholesale dealer's authorisation

40. The fee for an application to vary a wholesale dealer's authorisation is—

- (a) £515 if the variation requires scientific or pharmaceutical assessment;
- (b) £430 if the variation only involves a change of ownership; and
- (c) otherwise £300.

Annual fee for a wholesale dealer's authorisation

41.—(1) The annual fee for a wholesale dealer's authorisation is—

- (a) £483; or
- (b) £315, if—
 - (i) the holder certifies when making the payment that the turnover during the previous year was less than £35,000; or
 - (ii) the authorisation only relates to products classified as AVM-GSL or homeopathic remedies;
- (c) £215 if the authorisation only relates to products authorised under Schedule 6 (exemptions for small pet animals).

(2) In this paragraph "turnover" means the sales value net of value added tax of all veterinary medicinal products (whether or not authorised for use in the United Kingdom) sold by way of wholesale dealing by the holder in the United Kingdom.

Inspection of a wholesale dealer's premises

- 42.** The fee for the inspection of a wholesale dealer's premises is—
- (a) £3,058; or
 - (b) £1,442 if—
 - (i) the authorisation only relates to products classified as AVM-GSL or homeopathic remedies; or
 - (ii) the turnover relating to all veterinary medicinal products in the calendar year preceding the inspection was less than £35,000;
 - (c) £830 if the authorisation only relates to products authorised under Schedule 6 (exemptions for small pet animals).

PART 5**Fees relating to feedingstuffs****Fees for approvals and annual fees relating to feedingstuffs in Great Britain**

43.—(1) Subject to sub-paragraph (3) the fee for the application for approval of establishments manufacturing feedingstuffs and approval of distributors of feedingstuffs in Great Britain is £70.

(2) An annual fee of £70 is payable in respect of any such approval.

(3) No fee is payable under sub-paragraph (1) in respect of an establishment where specified feed additives are manufactured if a veterinary medicinal product intended to be incorporated into feedingstuffs is manufactured at that establishment in accordance with a manufacturing authorisation.

(4) Fees relating to feedingstuffs are payable with the application or on invoice for the subsequent annual fee.

(5) Where more than one manufacturing activity is carried out at one establishment only one fee (the highest) is payable.

Inspection fees relating to feedingstuffs in Great Britain

44. Fees for the inspection of establishments manufacturing or distributing feedingstuffs in Great Britain are in accordance with the following table.

Inspection fees

	<i>Type of establishment inspected</i>	<i>Fee payable (£)</i>
1	Establishment manufacturing a specified feed additive: ⁽¹⁾	1,810
2	Establishment manufacturing a premixture:	1,090
3	Establishment manufacturing feedingstuffs using specified feed additives and veterinary medicinal products directly at any concentration, or using premixtures	1,090

- (1) No fee is payable for premises that already have a manufacturing authority relating to veterinary medicinal products for incorporating into feedingstuffs.

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	<i>Type of establishment inspected</i>	<i>Fee payable (£)</i>
	or specified feed additive complementary feedingstuffs:	
4	Establishment manufacturing feedingstuffs for placing on the market using a veterinary medicinal product or premixture where the concentration of veterinary medicinal product in the feedingstuffs is 2 kg per tonne or more:	961
5	Establishment manufacturing feedingstuffs using premixtures or specified feed additive complementary feedingstuffs containing specified feed additives when the feedingstuffs are to be placed on the market:	405
6	Establishment manufacturing feedingstuffs for the manufacturers own use using a veterinary medicinal product or premixture where the concentration of veterinary medicinal product in the feedingstuffs is 2 kg per tonne or more:	320
7	Establishment manufacturing feedingstuffs using premixtures containing specified feed additives when the feedingstuffs are to be used by the person manufacturing the feedingstuffs:	240
8	Establishment distributing specified feed additives, premixtures or feedingstuffs containing specified feed additives, or premixtures or complementary feedingstuffs containing veterinary medicinal products:	227

(1) No fee is payable for premises that already have a manufacturing authority relating to veterinary medicinal products for incorporating into feedingstuffs.

Fees payable in relation to feedingstuffs in Northern Ireland

45.—(1) The annual fees payable for the approval of establishments manufacturing and distributing feedingstuffs in Northern Ireland are in accordance with the following table.

(2) Fees are payable with the application or, for the subsequent annual fee, on invoice.

(3) Where more than one manufacturing activity is carried out at one establishment only the highest fee is payable.

Approval fees

	<i>Type of establishment</i>	<i>Fee payable (£)</i>
1	Establishment manufacturing a specified feed additive ^(a) :	545
2	Establishment manufacturing a premixture:	435

(a) No fee is payable for establishments that already have a manufacturing authority relating to veterinary medicinal products for incorporating into feedingstuffs.

	<i>Type of establishment</i>	<i>Fee payable (£)</i>
3	Establishment manufacturing feedingstuffs using specified feed additives and veterinary medicinal products directly at any concentration, or using premixtures or specified feed additive complementary feedingstuffs:	435
4	Establishment manufacturing feedingstuffs for placing on the market using a veterinary medicinal product or premixture where the concentration of veterinary medicinal product in the feedingstuffs is 2 kg per tonne or more:	320
5	Establishment manufacturing feedingstuffs using premixtures or specified feed additive complementary feedingstuffs containing specified feed additives when the feedingstuffs are to be placed on the market:	170
6	Establishment manufacturing feedingstuffs for the manufacturers own use using a veterinary medicinal product or premixture where the concentration of veterinary medicinal product in the feedingstuffs is 2 kg per tonne or more:	131
7	Establishment manufacturing feedingstuffs using premixtures containing specified feed additives when the feedingstuffs are to be used by the person manufacturing the feedingstuffs:	110
8	Establishment distributing specified feed additives, premixtures or feedingstuffs containing specified feed additives, or premixtures or feedingstuffs containing veterinary medicinal products:	70
(a)	No fee is payable for establishments that already have a manufacturing authority relating to veterinary medicinal products for incorporating into feedingstuffs.	

Fees relating to premises for supply by suitably qualified persons

46.—(1) The fee to approve of premises for the retail supply of veterinary medicinal products by suitably qualified persons is—

- (a) £265; or
- (b) if the premises are only authorised to supply veterinary medicinal products for the treatment of—
 - (i) horses (or horses and companion animals) £145; or
 - (ii) companion animals £110.

(2) The subsequent annual fee is—

- (a) £185; or
- (b) if the premises are only authorised to supply veterinary medicinal products for the treatment of—
 - (i) horses (or horses and companion animals) £95; or
 - (ii) companion animals £70.

PART 6

General

Testing samples

47. The fee for testing a sample required to be submitted by the Secretary of State is the full economic cost of the test.

Animal test certificates

48.—(1) The fee for an animal test certificate is £345 in the case of—

- (a) an immunological veterinary medicinal product that has been authorised in another member State for the species on which the proposed test will be conducted;
- (b) a pharmaceutical veterinary medicinal product that has been authorised in another member State for use with a food-producing species on which the proposed test will be conducted where the same or similar dosage regime and method of administration is to be used in the medicinal test as is authorised; or
- (c) a pharmaceutical veterinary medicinal product authorised in another member State for human or animal use where the test is to be conducted on companion animals only.

(2) The fee for an animal test certificate to administer medicinal products in a small scale trial to test them for clinical safety or efficacy is £30.

(3) In any other case the fee is £815.

(4) The fee for an application for a variation of the certificate is £265 for each change.

(5) The fee for an application to renew a certificate is £130.

(6) The Secretary of State may waive the fee if satisfied that the application is in relation to developing a veterinary medicinal product for a limited market (for example, for a minor species, a minor use, or for a disease with restricted regional distribution).

Importation of a veterinary medicinal product for treatment under the cascade

49.—(1) The fee for a certificate to import (if necessary) and be in possession of and administer a veterinary medicinal product under the cascade is—

- (a) £15 if the veterinary medicinal product is authorised in another member State;
- (b) £30 if the veterinary medicinal product is authorised in a third country.

(2) The fee is payable in respect of each animal treated, but in the case of administration to and treatment of a discrete group of animals, the Secretary of State may notify the applicant in writing that a fee for only one animal is payable.

(3) There is no fee if the application is made using the website of the Veterinary Medicines Directorate.

Wholesale dealer's import certificate

50.—(1) The fee payable by the holder of a wholesale dealer's authorisation for a certificate to import and store a veterinary medicinal product not authorised in the United Kingdom to enable it to be supplied for administration under Schedule 4 is £1,320.

(2) The fee is only payable if, in the twelve month period immediately before the application, the applicant has supplied the veterinary medicinal product to which the certificate relates in accordance with at least 100 certificates.

Specific batch control

51. The fee for an authorisation to release a veterinary medicinal product, under specific batch control is—

- (a) £560; or
- (b) £100 for each batch if a number of specific batch control applications are made at the same time and all the batches are affected by the same issue.

Submission of control tests of an immunological product

52. The fee for the submission of the results of tests carried out on a batch of immunological products other than autogenous vaccines prior to release is £80.

Export certificates

53. The fee for an application for an export certificate is £30, and £15 for each certified copy.

Provision of advice

54. The fee for an application for written advice from the Secretary of State as to whether or not a product requires a marketing authorisation is £885.

Appeals to the Veterinary Products Committee

55. The fee for an appeal to the Veterinary Products Committee is £1,500.

Fee relating to an appointed person

56. The appellant is liable for the full economic cost of a referral to an appointed person subject to a maximum of £5,000.

Fees relating to a veterinary surgeon's practice premises

57.—(1) The fee for the inspection of a veterinary surgeon's practice premises is £350.

(2) The initial registration and annual fee for the registration of veterinary practice premises with the Royal College of Veterinary Surgeons to supply veterinary medicinal products is £34.

(3) Notwithstanding paragraph 2 of this Schedule, this is payable to the Royal College of Veterinary Surgeons.

Refund of fees relating to the Veterinary Products Committee or appointed persons

58. The Secretary of State must refund the fee payable in relation to an appeal to the Veterinary Products Committee or to an appointed person if, as a result of the appeal, the Secretary of State changes the decision that was the subject of the appeal.

Fees relating to an improvement notice

59. If an improvement notice is served under these Regulations, the fee for any subsequent inspection necessary as a result of the notice is the full economic cost of the inspection, payable by the person on whom the notice was served.

Non-payment of fees

60. Where any fee (other than any fee relating to a manufacturing authorisation or wholesale dealer's authorisation) is not paid, the Secretary of State may, after giving one month's written warning, suspend the processing of any application from the person who has not paid the fee.

Waiver or reduction of fees

61.—(1) If the Secretary of State is satisfied that for reasons of human or animal health or the protection of the environment it is desirable that a product should be authorised for veterinary use or that an authorised product should remain on the market the Secretary of State may waive or reduce any fees payable under these Regulations.

(2) An applicant or the holder of a marketing authorisation must provide full written justification for any waiver or reduction.

Reduction of fees when an application is withdrawn

62.—(1) Where an application for a marketing authorisation, or any variation referred to in paragraph 17 or 18 as a Type II variation, an extension, an extension-led grouped variation or a Type II led grouped variation is withdrawn before determination, the fee is reduced in accordance with this paragraph.

(2) If no assessment (veterinary, scientific or pharmaceutical) has begun, the reduction is 90%.

(3) If assessment has begun but the Secretary of State has not yet requested further data, the reduction is 50%.

(4) If the Secretary of State has requested further information but it has not yet been provided, the reduction is 25%.

(5) If the further information requested has been supplied but has not yet been fully assessed or the application has not been referred to the Veterinary Products Committee, the reduction is 10%

(6) Once the further information has been fully assessed, or the application has been referred to the Veterinary Products Committee, there is no reduction.