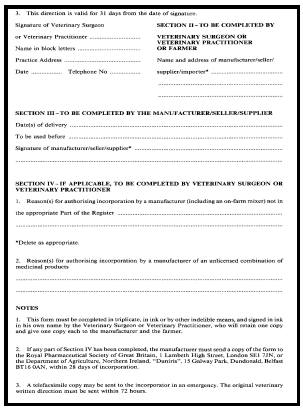
Code of practice for category A registered manufacturers of medicated animalfeedingstuffs.

Great Britain. Ministry of Agriculture, Fisheries and Food - eCFR :: 21 CFR Part 207



3. This direction is valid for 31 days from the date of signature.

Signature of Veterinary Surgeon or Veterinary Practitioner .....
Name in block letters .....
Practice Address .....
Date ..... Telephone No .....

SECTION II - TO BE COMPLETED BY VETERINARY SURGEON OR VETERINARY PRACTITIONER OR FARMER

Name and address of manufacturer/seller/supplier/importer* .....

SECTION III - TO BE COMPLETED BY THE MANUFACTURER/SELLER/SUPPLIER

Date(s) of delivery .....
To be used before .....
Signature of manufacturer/seller/supplier* .....

SECTION IV - IF APPLICABLE, TO BE COMPLETED BY VETERINARY SURGEON OR VETERINARY PRACTITIONER

1. Reason(s) for authorising incorporation by a manufacturer (including an on-farm mixer) not in the appropriate Part of the Register .....

*Delete as appropriate.

2. Reason(s) for authorising incorporation by a manufacturer of an unlicensed combination of medicinal products .....

NOTES

1. This form must be completed in triplicate, in ink or by other indelible means, and signed in ink in his own name by the Veterinary Surgeon or Veterinary Practitioner, who will retain one copy and give one copy each to the manufacturer and the farmer.

2. If any part of Section IV has been completed, the manufacturer must send a copy of the form to the Royal Pharmaceutical Society of Great Britain, 1 Lambeth High Street, London SE1 7JN, or the Department of Agriculture, Northern Ireland, "Dunrita", 15 Galway Park, Dundonald, Belfast BT16 0AN, within 28 days of incorporation.

3. A telefacsimile copy may be sent to the incorporator in an emergency. The original veterinary written direction must be sent within 72 hours.

Description: -
-
Portrait miniatures -- Private collections -- New York (State) -- New York -- Exhibitions.
Portrait miniatures, American -- Exhibitions.
Metropolitan Museum of Art (New York, N.Y.) -- Exhibitions.
Manney, Gloria -- Art collections -- Exhibitions.
Manney, Richard -- Art collections -- Exhibitions.
Mozart, Wolfgang Amadeus, -- 1756-1791.
Medicated feeds.
Veterinary drugs.Code of practice for category A registered manufacturers of medicated animalfeedingstuffs.
-Code of practice for category A registered manufacturers of medicated animalfeedingstuffs.
Notes: PB 0766.
This edition was published in 1991

Filesize: 25.61 MB

Tags: #The #Medicines #(Medicated #Animal #Feeding #Stuffs) #Regulations #1992

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Generally, there are few problems resulting from interpretation differences, but medicated feed manufacturers should be aware that differences of opinions can and do occur over the meaning of the regulations. FDA may also cancel the reservation of a proposed NDC at any time on the request of the person whose labeler code is included in the proposed NDC.

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Salvager means a person who owns or operates an establishment that engages in salvaging. Focus is on prevention of contamination by physical separation. Material change means any change in any drug listing information, as required under , , , , or except changes in format of labeling, labeling changes of an editorial nature, or inclusion of a bar code or initial inclusion of an NDC on the label.

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See comments on Section 225. The definitions and interpretations of terms in sections 201 and 510 of the Federal Food, Drug, and Cosmetic Act apply to the terms used in this part, if not otherwise defined in this section.

FDA Regulation of Medicated Feed

Operation should be on a predetermined, systematic and documented basis. No changes have been applied to the text. The second and current was published on June 3, 2015.

CFR

Equipment cleanout addresses this second goal. A drug approved for use in or on animal food as a VFD drug can be fed to animals only under the professional supervision of a licensed veterinarian.

CFR

Content of labeling means: 1 For human prescription drugs that are subject to section 505 of the Federal Food, Drug, and Cosmetic Act or section 351 of the Public Health Service Act: The content of the prescription drug labeling as specified in , , and , including all text, tables, and figures.

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All manufacturers, repackers, relabelers, or of Type B or Type C medicated feeds are exempt from listing.

Related Books

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- Teacher and school law - cases and materials in the legal foundations of education
- Fait divers.
- Women after prison
- The Non-Domestic Rating (Demand Notices) (Wales) Regulations 1993 (Statutory Instruments: 1993: 252)