

Feed Distributor Responsibilities, Minor Species, 4-H, and Other Questions

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Feed Distributor Responsibilities

- File one-time notice with FDA of intent to distribute VFD drugs
- ONLY fill a VFD order if VFD contains all required information
- Ensure animal feed containing the VFD drug complies with VFD order and label directions
- Retain VFD for 2 year

Feed Distributor Responsibilities

- Retain records of receipt and distribution of all medicated animal feed containing VFD drug for 2 years
- Provide VFD orders for inspection and copying by FDA upon request
- Retain records of VFD manufacturing for 1
 year and must be available for inspection and
 copying by FDA upon request

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Feed Distributor Responsibilities

- If you are originating distributor, you must obtain an acknowledgement letter from the receiving distributor before the feed is shipped
- If you are the originating distributor, you must retain a copy of each receiving distributor's acknowledgement letter for 2 years

Feed Distributor Responsibilities

- All labeling and advertising contains the following statement:
 - "Caution: Federal law restricts medicated feed containing this veterinary feed directive (VFD) drug to use by or on the order of a licensed veterinarian."

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Acknowledgement Letter

- Written communication provided to the original distributor from the receiving distributor when shipping animal feed containing a VFD drug in the absence of a valid VFD
- Different from FDA notification

Acknowledgement Letter Requirements

- Distributor will not ship the VFD feed to an animal production facility that does not have a valid VFD
- 2. Distributor will not ship the VFD feed to another distributor without receiving a similar acknowledgement letter
- 3. Distributor has complied with the distributor notification requirements

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Minor and Other Species

- Primary species
 - Cattle
 - Horses
 - Swine
 - Chickens
 - Turkeys
 - Dogs
 - Cats

- Minor Species
 - Any species other than "primary" species
 - Sheep
 - Goats
 - Buffalo/bison
 - · Honey Bees

AMDUCA

- Animal Medicinal Drug Use Clarification Act
 - Prohibits ANY extra-label use in medicated feeds
 - Includes minor species

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Minor Use and Minor Species Animal Health Act of 2004

- Minor use drugs that are intended for use in major species for diseases that occur infrequently or in limited areas and only in small numbers of animals
- Intended to make more medications legally available for treatment of minor species and uncommon diseases in the major species

Compliance Policy Guide 615.115

- "Agency ordinarily will not consider regulatory action if"...
 - Medicated feed is approved in major species, but used in minor species
 - Medicated feed is limited to farmed or confined species
 - Medicated feed is used within the context of a valid VCPR
 - Extra-label use is limited to therapeutic treatments, not production purposes
 - Veterinarian is responsible for establishing withdrawal period

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Antimicrobials for Minor Species

- Goats
 - Monensin
 - Neomycin

- Sheep
 - Neomycin
 - Oxytetracycline
 - Chlortetracycline
 - Lasalocid

Aureomycin

Sheep	
Breeding Sheep: Reduction in the incidence of (vibrionic) abortions caused by Campylobacter fetus infection susceptible to chlortetracycline.	80
Indications	In complete feed Chlortetracycline g per ton
Swine	
Increased rate of weight gain and improved feed efficiency.	10-50
Reduction in the incidence of cervical lymphadenitis (jowl abscesses) caused by Group E <i>Streptococci</i> susceptible to chlortetracycline.	50-100
Breeding Swine: Control of leptospirosis (reducing the incidence of abortion and shedding of leptospirae) caused by <i>Leptospira pomona</i> susceptible to chlortetracycline. Feed continuously for not more than 14 days.	400
Sheep	
Increased rate of weight gain and improved feed efficiency.	20-50

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CTC 4G

SHEEP (Growing Increased rate d improved feed efficiency. (20 more than 12.5 p SHEEP (Breeding) t less than 5 nor complete ration.

Reducing the incidence of (vibrionic) abortion caused by Campylobacter fetus infection susceptible to Chlortetracycline. (80 mg/head/day: Feed 0.2 lb per 10 head per day.

4-H Animals

- All animals consuming feed with a VFD drug or combination VFD drug requires a VFD form
- 4-H advisor CANNOT be the client
 - Unless they are the primary caretaker
- Group of animals located at same location
 - Each person will need their own VFD

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FDA Checklist for VFD Inspection

- Randomly select 3 VFD forms at feed distributor
 - Stop at producer and veterinarian
- Veterinarian and Client information
- Premises specified on the VFD have animals from form
- Date of VFD order
- Affirmation of intent for VFD combination drug
- Vet's signature, electronic or written

Tips for VFD Preparation

- 1. Determine which, if any, of your feeds contain feed-grade medications.
 - a) Consult veterinarian and feed distributor
- 2. Develop veterinary-client-patient relationship
- 3. Keep accurate records of all feed-grade medications
- 4. Have a designated storage area for VFD forms

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Questions

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