

Pennox (oxytetracycline) Veterinary Feed Directive for use in Turkeys

Client: _____ Veterinarian: _____
Business or Home Address: _____
Address: _____
Phone #: _____ Phone #: _____

Approximate number of turkeys to be treated: _____

Location of animals: _____

Special Instructions and/or other animal identifications:

Indication, Drug Level in Medicated Feed, and Duration of Use (select one and specify the additional required information):

- A)** Control of hexamitiasis caused by *Hexamita meleagrides* susceptible to oxytetracycline.
Drug level: 100 g/ton
Duration of use: _____ days (7 to 14 days)
- B)** Control of infectious synovitis caused by *Mycoplasma synoviae* susceptible to oxytetracycline.
Drug level: 200 g/ton
Duration of use: _____ days (7 to 14 days)
- C)** Control of complicating bacterial organisms associated with bluecomb (transmissible enteritis, coronaviral enteritis) susceptible to oxytetracycline.
Drug level: _____ g/ton in order to provide 25 mg/lb body weight / day
Duration of use: _____ days (7 to 14 days)

Caution: Use of feed containing this Veterinary Feed Directive (VFD) drug in a manner other than as directed on the labeling (extra-label use) is not permitted.

For use in Dry Feeds Only. Not for Use in Liquid feed Supplements.

**Residue Warnings: Do not feed to Turkeys Producing Eggs for Human Consumption.
Zero-day withdrawal period.**

Combination Use:

This VFD only authorizes the use of the VFD drug(s) cited in this order and is not intended to authorize the use of such drug(s) in combination with any other animal drugs.

VFD Issuance Date: _____

VFD Expiration Date: _____

Month/Day/Year
(Not to exceed 6 months from issuance date)

Veterinarian's signature: _____

Copy – Supplier

Copy – Client

original – Veterinarian