

**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