

# Veterinary Feed Directive for Use in Beef Cattle

## Tylovet 100<sup>®</sup>

### (tylosin phosphate)

Sequential VFD ID Number, if appropriate

 Veterinarian: \_\_\_\_\_  
 Address: \_\_\_\_\_  
 Phone #: \_\_\_\_\_  
 FAX or email: (optional) \_\_\_\_\_

 Client: \_\_\_\_\_  
 Business/Home Address: \_\_\_\_\_  
 Phone #: \_\_\_\_\_  
 FAX or email: (optional) \_\_\_\_\_

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

 For reduction of incidence of liver abscesses associated with *Fusobacterium necrophorum* and *Arcanobacterium pyogenes*.  
**Drug Level:** \_\_\_\_\_ g/ton (8 to 10 g/ton)  
**Duration of Feeding:** \_\_\_\_\_ days

**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.**

 Approximate number of **Cattle** to be treated: \_\_\_\_\_  
 Premise or Location of cattle: \_\_\_\_\_

 Special Instructions and/or other animal identifications:  

**Affirmation of Intent (for combination VFD drugs): check the appropriate box:**

- 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.
- This VFD authorizes the use of the VFD drug(s) cited in this order in the following FDA-approved, conditionally approved, or indexed combination(s) in medicated feed that contains the VFD drug(s) as a component.

|                          | Drug(s) and Dose Range(s) [all listed doses are 90% DM basis]  | Specifications*  |
|--------------------------|--|--|
| <input type="checkbox"/> | monensin at 10 to 40 g/ton to provide 0.14 to 0.42 mg/lb bodyweight/day<br>ractopamine at 9.8 to 24.6 g/ton to provide 90 to 430 mg/head/day<br>(supplied by Rumensin <sup>™</sup> and Actogain <sup>®</sup> , ANADA 200-585)  | For use in cattle fed in confinement for slaughter for the last 28 to 42 days on feed. |
| <input type="checkbox"/> | monensin at 10 to 40 g/ton to provide 0.14 to 0.42 mg /lb bodyweight/day<br>ractopamine at 8.2 to 24.6 g/ton to provide 70 to 430 mg/head/day<br>(supplied by Rumensin <sup>™</sup> and Actogain <sup>®</sup> , ANADA 200-585)   | For use in cattle fed in confinement for slaughter for the last 28 to 42 days on feed. |
| <input type="checkbox"/> | monensin at 10 to 40 g/ton to provide 0.14 to 0.42 mg/lb bodyweight/day<br>ractopamine up to 800 g/ton as a top dress to provide 70 to 400 mg/head/day<br>(supplied by Rumensin <sup>™</sup> and Actogain <sup>®</sup> , ANADA 200-585)  | For use in cattle fed in confinement for slaughter for the last 28 to 42 days on feed. |
| <input type="checkbox"/> | monensin at 10 to 40 g/ton to provide 0.14 to 0.42 mg/lb bodyweight/day<br>ractopamine at 9.8 to 24.6 g/ton to provide 90 to 430 mg/head/day<br>Melengestrol acetate, melengestrol at the rate of 0.125 to 1.0 mg/lb (top dress or mixed at bunk with other Type C feed) to provide 0.25-0.5 mg/head/day. (supplied by Rumensin <sup>™</sup> and Actogain <sup>®</sup> and melengestrol acetate, MGA <sup>®</sup> , ANADA 200-583) | For use in cattle fed in confinement for slaughter for the last 28 to 42 days on feed. |
| <input type="checkbox"/> | Other FDA-approved, conditionally approved, or indexed combination:  |  |

\*for complete information see the approved Type C medicated feed label.

- This VFD authorizes the use of the VFD drug(s) cited in this order in any FDA-approved, conditionally approved, or indexed combination(s) in medicated feed that contains the VFD drug(s) as a component.

**Warning:** No withdrawal period is required.

Date of VFD Issuance: \_\_\_\_\_ (dd/mm/yyyy)

 Date of VFD Expiration: \_\_\_\_\_ (dd/mm/yyyy)  
 (Cannot exceed 6 months after issuance)

Veterinarian's signature: \_\_\_\_\_

**Color Z Original – Veterinarian**

**Color X Copy – Supplier**

**Color Y Copy – Client**

All parties must retain a copy of this VFD for 2 years after issuance