Phibro
ANIMAL HEALTH CORPORATION ®

**AUREO S 700® Granular (chlortetracycline and sulfamethazine)** Veterinary Feed Directive for use in Beef Cattle

Veterinarian: _____ Client: _____

Address: _____ Business or Home Address: _____

_____ _____

Phone #: _____ Phone #: _____

Drug Levels: _____g/ton each for chlortetracycline and sulfamethazine (specify level to provide 350 mg/head/day chlortetracycline and 350 mg/head/day sulfamethazine).

Duration of Use: Feed for 28 days

**Indications for Use:**
Beef Cattle: As an aid in the maintenance of weight gains in the presence of respiratory disease such as shipping fever.

**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 ***Beef Cattle*** to be treated: _____

Premises or Location of animals: _____

Special Instructions and/or other animal identifications:

| |
|---|
| |

A**ffirmation of Intent (**for combination VFD drugs**):**

☐ 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 concentrations(s) [all listed doses are 90% DM basis] | Specifications* |
|---|---|
| ☐ 10 to 30 g/ton lasalocid to provide 100-300 mg/hd/day (BOVATEC®; NADA 141-535) | Beef steers and heifers fed in confinement for slaughter |
| ☐ 25-30 g/ton lasalocid to provide 250-300 mg/hd/day (BOVATEC®; NADA 141-535) | Beef steers and heifers fed in confinement for slaughter |
| ☐ 30-181.8 g/ton to provide 1 mg per 2.2 lbs bodyweight per day (maximum 360 mg lasalocid daily) (BOVATEC®; NADA 141-535) | Beef cattle up to 800 lb |
| ☐ 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.

**Withdrawal Periods and Residue Warnings:**
Withdraw 7 days prior to slaughter.
A withdrawal period has not been established for this product in pre-ruminating calves.
Do not use in calves to be processed for veal.

VFD Expiration Date: _____ month/day/year (Not to exceed 6 months from issuance date)

Veterinarian's Signature: _____ Date of Issuance (Month/Day/Year): _____

Approved by FDA under NADA # 035-805 ARS104724USA Rev. March 04, 2025

Phibro Animal Health Corporation
300 Frank W. Burr Blvd
Teaneck, NJ 07666

Copy – Supplier Copy – Client Copy – Feed Mill