AT-A-GLANCE
PRODUCT SUMMARY

Pulmotil® (tilmicosin)

Pulmotil® is an innovative bovine respiratory disease (BRD) treatment for groups of cattle in the early stages of an outbreak that provides 14 days of sustained in-feed therapy, a practice that reduces stress associated with cattle handling.

14-DAY BRD CONTROL — Pulmotil reduces the risk of a BRD outbreak once cattle are on feed and reduces the impact of subclinical BRD and associated economic losses.

IN-FEED GROUP THERAPY — Feeding Pulmotil is a treatment program that allows you to provide 14 days of sustained in-feed therapy to groups of cattle showing early signs of BRD.

REDUCED MORBIDITY — The four-study registration summary showed Pulmotil reduced BRD pulls by 35.6% and reduced cumulative morbidity through day 28.[1]

Effective BRD control isn't just a chute-side protocol anymore.

Pulmotil is also available in 18 and 5.68 formulations to meet your operation's needs.

PULMOTIL FEEDING GUIDELINES & LABEL DIRECTIONS

INDICATIONS: For the control of bovine respiratory disease (BRD) associated with *Mannheimia haemolytica*, *Pasteurella multocida* and *Histophilus somni* in groups of beef and non-lactating dairy cattle, where active BRD has been diagnosed in at least 10% of the animals in the group.

FEEDING DIRECTIONS: Feed continuously for a single, 14-day period at 568 to 757 g/ton of tilmicosin (100% DM basis) in a Type C medicated feed as the sole ration to provide 12.5 mg/kg of body weight/day.

STORAGE: Store at less than or equal to 25°C (77°F). Excursions to 40°C (104°F) are acceptable. Avoid excessive moisture.

The label contains complete use information, including cautions and warnings. Always read, understand and follow the label and use directions.

IMPORTANT SAFETY INFORMATION

- **CAUTION:** Federal law restricts medicated feed containing this veterinary feed directive (VFD) drug to use by or on the order of a licensed veterinarian.
- Cattle intended for human consumption must not be slaughtered within 28 days of the last treatment.
- To assure both food safety and responsible use, treatment must be initiated within the first 45 days of the production period. The treatment should not occur concurrent with or following administration of an injectable macrolide, or within 3 days following administration of a non-macrolide injectable BRD therapy.
- VFD expiration date must not exceed 45 days from the time of issuance. VFDs shall not be refilled.
- Use only in cattle fed in confinement for slaughter.
- Cattle with severe clinical illness should be evaluated for individual treatment with an alternative non-macrolide therapy.
- Do not use in female dairy cattle 20 months of age or older or in veal calves.
- Safety has not been established for cattle intended for breeding.
- Do not allow horses or other equines access to feeds containing tilmicosin.

For the control of Bovine Respiratory Disease (BRD) associated with *Mannheimia haemolytica*, *Pasteurella multocida* and *Histophilus somni* in groups of beef and non-lactating dairy cattle, where active BRD has been diagnosed in at least 10% of the animals in the group: Feed continuously for a single, 14-day period at 568 to 757 g/ton of tilmicosin (100% DM basis) in a Type C medicated feed as the sole ration to provide 12.5 mg/kg of body weight/day.

[1]Pulmotil 90 Freedom of Information Summary (NADA 141-064).

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