

**Tilmicosin and Monensin Type C Medicated Complete Feed
Bovine Respiratory Disease and Feed Efficiency in Confined Cattle**

FOR USE IN CATTLE ONLY

CAUTION: Federal law limits this drug to use under the professional supervision of a licensed veterinarian. Animal feed bearing or containing this veterinary feed directive drug shall be fed to animals only by or upon a lawful veterinary feed directive issued by a licensed veterinarian in the course of the veterinarian's professional practice.

INDICATIONS

For improved feed efficiency and control of bovine respiratory disease (BRD) associated with *Mannheimia haemolytica*, *Pasteurella multocida*, and *Histophilus somni* in groups of cattle fed in confinement for slaughter, where active BRD has been diagnosed in at least 10% of the animals in the group.

ACTIVE DRUG INGREDIENTS

Tilmicosin (as tilmicosin phosphate)568 to 757 grams per ton*^a.
Monensin, USP.....5 to 40 grams per ton*^b.

*Final printed label must bear a single drug concentration.

^a 100% dry matter basis.

^b 90% dry matter basis.

GUARANTEED ANALYSIS

Crude Protein, not less than..... %
¹
Non-Protein Nitrogen (NPN) , not more than..... %
Crude Fat, not less than..... %
Crude Fiber, not more than..... %
Calcium, not less than..... %
Calcium, not more than..... %
Phosphorus, not less than..... %
²
Salt , not less than..... %
²
Salt , not more than..... %
³
Sodium , not less than..... %
³
Sodium , not more than..... %
Potassium, not less than..... %
^{2, 4}
Vitamin A , not less than..... I.U./lb

¹ When added.

² If added.

³ Shall be guaranteed only when total sodium exceeds that furnished by the maximum salt guarantee.

⁴ Other than precursors of Vitamin A.

INGREDIENTS

Each ingredient must be named in accordance with the names and definitions adopted by the Association of American Feed Control Officials (AAFCO).

FEEDING DIRECTIONS

This Type C medicated feed is to be fed continuously as the sole ration for a single, 14 day period at 568 grams to 757 grams tilmicosin per ton on a 100% dry matter basis (511.2 to 681.3 g per ton on a 90% dry matter basis) to provide 12.5 mg tilmicosin/kg/head/day and 5 grams to 40 grams monensin per ton on a 90% dry matter basis (5.6 to 44.4 grams per ton on a 100% dry matter basis) to provide 50 to 480 mg monensin/head/day. No additional improvement in feed efficiency has been shown from feeding monensin at levels greater than 30 grams per ton on a 90% dry matter basis (360 mg monensin per head per day).

CAUTION:

Do not allow horses or other equines access to feeds containing tilmicosin or monensin. Ingestion of monensin by horses has been fatal. Monensin medicated cattle feeds are safe for use in cattle only. Consumption by unapproved species may result in toxic reactions. The safety of tilmicosin has not been established in cattle intended for breeding purposes.

Feeding undiluted or mixing errors resulting in high concentrations of monensin has been fatal to cattle. Must be thoroughly mixed in feeds before use. Do not exceed the levels of monensin recommended in the feeding directions, as reduced average daily gains may result.

To assure both food safety and responsible use in cattle, the treatment of cattle with this medicated feed is required to 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

Tilmicosin medicated feed treatment has not been evaluated in cattle with severe clinical disease. Cattle with severe clinical illness should be evaluated for individual treatment with an alternative non-macrolide therapy.

Complete Type C medicated feeds containing tilmicosin should not be pelleted. Do not use in any feeds containing bentonite, cottonseed meal, or cottonseed hulls. Bentonite, cottonseed meal, or cottonseed hulls in feeds may affect the efficacy of tilmicosin.

The expiration date for a tilmicosin Veterinary Feed Directive (VFD) for cattle must not exceed 45 days from the time of issuance. VFDs for tilmicosin phosphate shall not be refilled.

WARNINGS:

RESIDUE WARNING: Cattle: Cattle intended for human consumption must not be slaughtered within 28 days of the last treatment with tilmicosin.

Tilmicosin is not approved for use in female dairy cattle 20 months of age or older. Use in these cattle may cause drug residues in milk. Tilmicosin and monensin are not approved for use in calves intended to be processed for veal. A withdrawal period has not been established in pre-ruminating calves.

User Safety Warnings: Avoid inhalation, oral exposure, and direct contact with skin or eyes. Operators mixing and handling PULMOTIL[®] 90 and RUMENSIN[®] 90 should use protective clothing, impervious gloves, goggles, and a NIOSH-approved dusk mask. Wash thoroughly with soap and water after handling. If accidental eye contact occurs, immediately rinse thoroughly with water. If irritation persists, seek medical attention. Not for human consumption. Keep out of reach of children. The Material Safety Data Sheet contains more detailed occupational safety information. To report adverse effects in users, to obtain more information, or to obtain a Material Safety Data Sheet, call 1-800-428-4441.

Lot no. _____ Net Weight: _____ lb (_____ kg)

MANUFACTURED BY
Bluebird Feed Company
Any town, USA 12345

For emergency medical information, to report an adverse effect, or for technical service call: 1-800-428-4441.

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