

TILMOVET[®] PREMIX

FOR USE IN SWINE FEEDS ONLY – MUST BE THOROUGHLY MIXED IN FEEDS BEFORE USE.

FOR
VETERINARY
USE ONLY

Package size: 5 & 10 kg
DIN: 02349507

Medicated Premix for Swine

ACTIVE INGREDIENT: Each bag contains: 200g tilmicosin
(as tilmicosin phosphate) per kilogram.



INDICATIONS:

Indication 1:

As an aid in reducing the severity of swine respiratory disease associated with *Actinobacillus pleuropneumoniae* and *Pasteurella multocida* when fed to pigs approximately 7 days prior to anticipated disease outbreak.

Dosage and administration:

- Feed at 200 g tilmicosin activity per tonne (1000 kg) for a 21-day period, beginning approximately 7 days before an anticipated outbreak. Feed continuously as the sole ration.
- To provide 200 g tilmicosin activity per tonne (1000 kg) in complete feed, thoroughly mix 1.0 kg TILMOVET Premix into 999 kg non-medicated swine feed.

Indication 2:

As an aid in reducing the severity of porcine polyserositis and arthritis associated with *Haemophilus parasuis* (Glasser's Disease) when fed to pigs approximately 7 days prior to anticipated outbreak.

Dosage and Administration:

- Feed at 400 g tilmicosin activity per tonne (1000 kg) for a 21-day period, beginning approximately 7 days before an anticipated outbreak. Feed continuously as the sole ration.
- To provide 400 g tilmicosin activity per tonne (1000 kg) in complete feed, thoroughly mix 2.0 kg TILMOVET Premix into 998 kg non-medicated swine feed.

CONTRA-INDICATIONS:

- 1) Do not use in animals hypersensitive to tilmicosin.
- 2) Tilmicosin is known to be toxic for horses. Do not allow horses or other equines access to feeds containing tilmicosin.

CAUTIONS:

- 1) Do not use in any feed (supplement, concentrate, or complete feed) containing bentonite. Bentonite in feeds may affect the efficacy of tilmicosin.
- 2) The safety of tilmicosin has not been established in boars used for breeding.

WARNINGS:

- 1) Treated animals must not be slaughtered for use in food for at least 14 days after the last treatment with this drug.
- 2) When mixing and handling TILMOVET premix, use protective clothing, impervious gloves, and a dust mask. Operators should wash thoroughly with soap and water after handling. If accidental eye contact occurs, immediately rinse thoroughly with water, if irritation persists, seek medical attention.
- 3) Keep out of reach of children.
- 4) To report adverse effects in users or to obtain occupational safety information, call 1-800-265-1763.

STORAGE

Store in the original container, do not store above 30°C, protect from freezing.
Shelf life after first opening of container: 3 months.

PRESENTATION

Tilmovet Premix is supplied in 5 and 10 kg three-ply paper bags.



MANUFACTURED BY: HUVEPHARMA AD, 33, James Boucher Blvd., Sofia 1407, Bulgaria
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