

# RE-COVR™

(tripelennamine hydrochloride injection)

250 mL 20 mg per mL

FDA approved under NADA #006-417



## ■ Rapid acting antihistamine

- Speeds recovery time. **RE-COVR** blocks harmful histamine effects which can prolong recovery process.

- **Approved for use in:**
  - Cattle (Beef and Dairy)
  - Horses

- Provides effective therapy for a wide variety of histamine-associated conditions, as well as supportive therapy for disease where histamine production is suspected.

Dosages:	Species	Route	Dose
	Cattle	IV or IM	2.5 ml/100 lbs.*
	Horses	IM only	2.5 ml/100 lbs.*

\*May be repeated every 6-12 hours if necessary

- **Withdrawal:** **Milk Discard** - 24 hours  
**Slaughter** - 4 days\*  
\*After last treatment

- **RE-COVR** is the ideal product for applications involving histamine toxemias and allergic reactions.

## APPLICATIONS INVOLVING HISTAMINE TOXEMIAS:

- Pneumonia
- Mastitis
- Septic Metritis
- Retained Placenta
- Milk Fever
- Grass Tetany
- Displaced Abomasum
- Bloat
- Grain Overload
- Toxic Indigestion
- Intestinal Obstruction
- Colic
- Second Degree Burns
- Shipping Fever

## APPLICATIONS INVOLVING ALLERGIC REACTIONS:

- Asthma (Heaves)
- Urticaria
- Milk Allergy
- Insect Bites
- Anaphylactic Shock
- Drug Sensitivities
- Plant Poisonings
- Snake Bites
- Edematous Swelling

# RE-COVR™

(tripelennamine hydrochloride injection)

20 mg per mL  
Antihistamine

**Horses** - For intramuscular injection only

**Cattle** - For intravenous or intramuscular injection

**Caution:** Federal law restricts this drug to use by or on the order of a licensed veterinarian.

**DESCRIPTION:** Tripelennamine hydrochloride is a white, crystalline material which is stable, nonhygroscopic, and readily soluble in water. Tripelennamine hydrochloride is characterized by its capacity to antagonize many of the pharmacologic effects of histamine. **RE-COVR™** (tripelennamine hydrochloride injection) is supplied as a sterile solution in multiple dose vials containing 20 mg of tripelennamine hydrochloride, USP per mL, and may contain sodium hydroxide for pH adjustment.

**INDICATIONS:** For use in horses and cattle in conditions in which antihistamine therapy may be expected to lead to alleviation of some signs of disease. Not for use in beef calves less than 2 months of age, dairy calves, and veal calves.

**USER SAFETY WARNINGS:** Not for use in humans. Keep out of reach of children. To obtain a Safety Data Sheet, contact KineticVet at 1-877-786-9882 or [www.KineticVet.com](http://www.KineticVet.com).

**ANIMAL SAFETY WARNINGS:** Administration of tripelennamine hydrochloride may give rise to excitement, ataxia, and convulsions. Central nervous system stimulation in the form of hyperexcitability, nervousness, and muscle tremors lasting up to 20 minutes have been noted in horses following administration.

Depression of the central nervous system and incoordination may occur when the drug is used at therapeutic dose levels. Disturbances in gastrointestinal function may occur in some instances.

**OTHER WARNINGS:** Do not use in horses intended for human consumption.

**CONTACT INFORMATION:** Contact KineticVet at (877) 786-9882 or [www.KineticVet.com](http://www.KineticVet.com). To report suspected adverse drug experiences, contact KineticVet at (877) 786-9882. For additional information about reporting adverse drug experiences for animal drugs, contact FDA at 1-888-FDA-VETS or <http://www.fda.gov/reportanimalae>.

**HOW SUPPLIED:** 100 mL, 250 mL and 500 mL multiple dose vials.

## DOSAGE AND ADMINISTRATION:

**Horses:** Administer intramuscularly only at a dose of 0.5 mg per lb of body weight (2.5 mL for each 100 lbs of body weight). This dose may be repeated in 6-12 hours if necessary.

**Cattle:** Administer intravenously or intramuscularly at a dose of 0.5 mg per lb of body weight (2.5 mL for each 100 lbs of body weight). This dose may be repeated in 6-12 hours if necessary. The intravenous route of administration may provide a more rapid onset of action. Use aseptic technique to administer **RE-COVR™** (tripelennamine hydrochloride injection).

Warm the solution to near body temperature prior to administration. Intramuscular injection should be made into the heavy musculature of the hind leg or cervical area.

While venomous snake bites have been treated with antihistaminic drugs, other conjunctive therapy is required because of toxic reactions associated with the protein complex of venom.

Do not puncture the stopper more than 30 times, use within the 24-month product expiry.

## WARNINGS AND PRECAUTIONS:

### Withdrawal Periods and Residue

**Warnings:** Cattle: Milk taken during treatment and for 24 hours after the last treatment must not be used for human consumption. Cattle must not be slaughtered for human consumption within 4 days following the last treatment with this drug product. Not for use in beef calves less than 2 months of age, dairy calves, and veal calves. A withdrawal period has not been established in the pre-ruminating calves.

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**INDICATIONS:** For use in horses and cattle in conditions in which antihistaminic therapy may be expected to lead to alleviation of some signs of disease. Not for use in beef calves less than 2 months of age, dairy calves, and veal calves. See package insert for complete indications for use.

**STORAGE AND HANDLING:** Store at 20°C to 25°C (68°F - 77°F), excursions permitted between 15°C to 30°C (59°F and 86°F). Keep from freezing. Protect from light.

## Manufactured For:

KineticVet | PO Box 12388 | Lexington, KY 40583

Made in USA

[www.KineticVet.com](http://www.KineticVet.com)

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PRODUCT #-2004-07-00



# KINETICVET™

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