MANNITOL- mannitol injection
Covetrus

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Covetrus
Mannitol

Recommended Use:

For use as an osmotic diuretic in dogs. Mannitol is essentially inert metabolically. When given parenterally, it is freely filtered at the glomerulus which produces osmotic diuresis as more than 90% of the mannitol injected escapes reabsorption.

Each 100 mL Contains:

Mannitol USP.....20 g

Water for Injection.....q.s.

This solution contains 1098 mOsmols/Liter

Dosage and administration:

The usual canine dosage administered intravenously is 1.5 - 2.0 g per kg body weight given over a 30 minute period. This is approximately 3.4-4.5 mL/lb of body weight.

Note:

Crystals of mannitol may form in a 20% saturated solution of mannitol. Dissolve the crystals by warming in hot water or autoclaving for 15 minutes. Cool to body temperature before administering. This is a single dose vial that contains no preservatives. Use entire contents when first opened.

Store at temperatures between 15°and 30°C (59°-86°F)

NDC:11695-1254-1

Net contents: 100 mL (3.4 fl oz)

Lot No.

Exp. Date

Reorder #002469

Questions?

(855) 724-3461

Distributed by:

Covetrus North America

400 Metro Place North

Dublin, OH 43017

covetrus.com

Made in the USA

AH-002469-02

RMS-92-1192

L919-1119

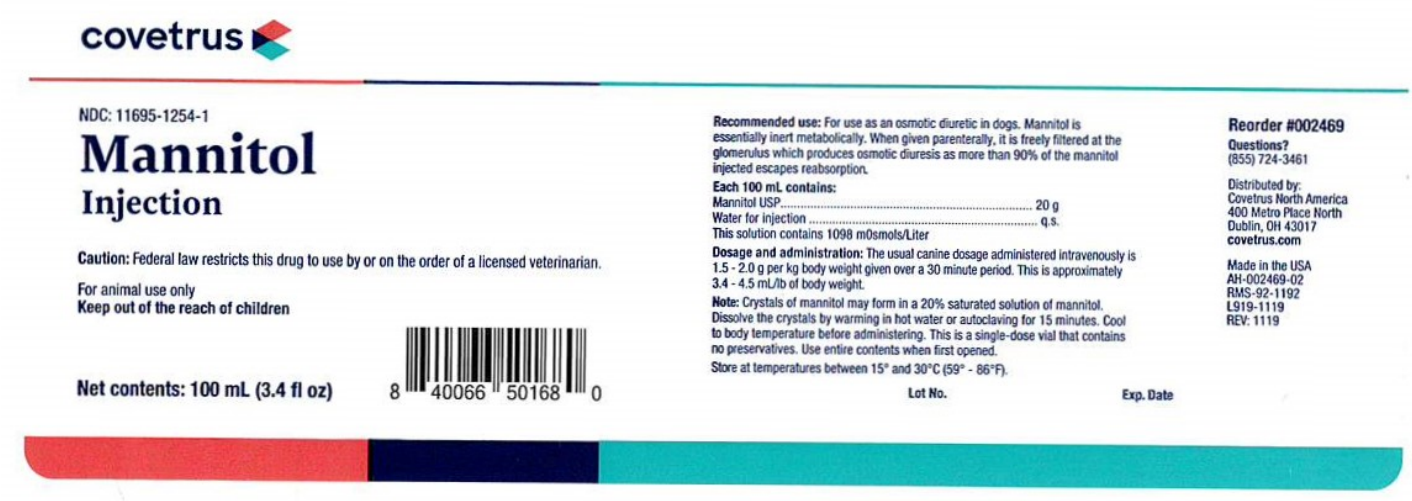
REV: 1119

Caution:

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

FOR ANIMAL USE ONLY

KEEP OUT OF THE REACH OF CHILDREN



MANNITOL

mannitol injection

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:11695-1254
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MANNITOL (UNII: 3OWL53L36A) (MANNITOL - UNII:3OWL53L36A)	MANNITOL	20 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:11695-1254-1	100 mL in 1 VIAL, SINGLE-USE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/16/2020	

Labeler - Covetrus (603750329)

Registrant - Nova-Tech, Inc (196078976)

Establishment

Name	Address	ID/FEI	Business Operations
Nova-Tech, Inc		196078976	manufacture

Revised: 3/2020 Covetrus