

MANNITOL 20% - mannitol injection

MWI (VetOne)

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](#).

VetOne

MANNITOL 20%

INDICATIONS:

Mannitol Injection 20% is indicated for use as an osmotic diuretic in canine species. 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.

Storage:

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

NDC 13985-052-15

100 mL

Lot #

Exp. Date

Manufactured by: Nova-Tech, Inc. Grand Island, NE 68801 for Neogen Corporation

□ Distributed by:

MWI

Boise, ID 83705

www.VetOne.net

Made in USA

V1 501050

Net Contentes: 100 mL

Rev. 05/18

RMS-92-530

ⓘ **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**

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VET one

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TAKE CARE OBSERVE LABEL DIRECTIONS MADE IN USA

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MANNITOL 20%

mannitol injection

Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:13985-052
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
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1	NDC:13985-052-15	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		06/20/2011		

Labeler - MWI(VetOne) (019926120)

Registrant - Nova-Tech, Inc (196078976)

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

Revised: 11/2019

MWI (VetOne)