

DEXTROSE- dextrose injection, solution
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.

Dextrose Solution 50%

Dosage And Administration:

Administer 50 mL per 100 lbs of body weight intravenously only. Dosage may be repeated in 8 to 10 hours or on successive days as needed.

Caution:

This product should be warmed to body temperature and administered slowly. This product contains no preservative. Do not use if solution is not clear. Entire contents should be used upon opening. Discard any unused portion.

FOR ANIMAL USE ONLY
KEEP OUT OF REACH OF CHILDREN

Active Ingredients:

Dextrose Monohydrate50 % w/v

Inactive Ingredients:

Water for Injection q.s.

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

TAKE TIME OBSERVE LABEL DIRECTIONS

NDC: 11695-1597-3

18-801

RMS 92-1024

Volume: 16.907 (500 mL)

Questions? (855) 724-3461

Reorder #069168

Manufactured by:

Nova-Tech, Inc.

Grand Island, NE 68801

Distributed by:

Covetrus North America

400 Metro Place North

Dublin, OH 43017

covetrus.com

Made in the USA

AH-069168-L-02

Rev: 0523

Lot No. Exp. Date

Indications:

For use in cattle as an aid in the treatment of uncomplicated primary ketosis.



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NDC: 11695-1597-3

Dextrose solution 50%

Sterile solution

**FOR ANIMAL USE ONLY
KEEP OUT OF REACH OF CHILDREN**

Net contents: 16.91 fl oz (500 mL)

Reorder #069168

Active ingredients:
Dextrose Monohydrate 50% w/v

Inactive ingredients:
Water for Injection q.s.

Store between 15° - 30°C (59° - 86°F).

Questions?
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DEXTROSE

dextrose injection, solution

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:11695-1597
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROSE MONOHYDRATE (UNII: LX22YL083G) (ANHYDROUS DEXTROSE - UNII:5SLOG7ROOK)	DEXTROSE MONOHYDRATE	50 g in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11695-1597-3	500 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/21/2017	

Labeler - Covetrus (603750329)

Registrant - Covetrus (603750329)

Establishment

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

Revised: 12/2023

Covetrus