DEXTROSE- dextrose injection, solution Vedco

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 50% SOLUTION KEEP OUT OF REACH OF CHILDREN

ACTIVE INGREDIENTS:

Each 100 mL contains:

Dextrose . H₂O 50 gms.

DOSAGE AND ADMINISTRATION:

For intraven-

ous administration only.

Cattle: 100 to 500 mL depending on size and

condition.

Treatment may be repeated in several hours or

on successive days as needed.

Store at 15 degrees C to 30 degrees C (59 degrees F-86 degrees F).

INDICATIONS:

For use as an aid in the treatment of acetonemia (Ketosis) in cattle.

CAUTION:

Intravenous administration must be done slowly and made under strict asepsis. Solution should be warmed to body temperature prior to administration.

This is a single dose container. This product contains no preservative. After a quantity has been withdrawn for injection, the remainder should be discarded. Do not administer intraperitoneally.

FOR ANIMAL USE ONLY

TAKE TIME OBSERVE LABEL DIRECTIONS

VINV-DEXT-ROUN Rev. 10-15

Distributed By

VEDCO, INC. RMS 92-365 St. Joseph, MO 64507 18-801

NET CONTENTS: 500 mL (16.9 fl. oz.)

□Lot No. Exp. Date

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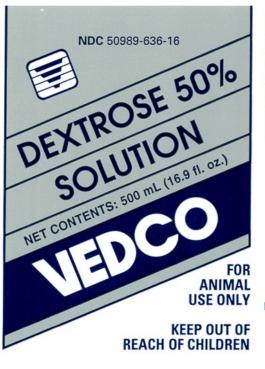


VINV-DEXT-ROUN

Rev. 10-15

Distributed By VEDCO, INC. St. Joseph, MO 64507

RMS 92-365 18-801



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EACH 100 mL CONTAINS:

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Lot No.

Exp. Date

DEXTROSE

dextrose injection, solution

Product Information			
Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:50989-636
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
DEXTROSE MONOHYDRATE (UNII: LX22YL083G) (ANHYDROUS DEXTROSE - UNII:5SL0G7R0OK)	DEXTROSE MONOHYDRATE	50 g in 100 mL	

ı	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:50989-636-16	500 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/0 1/20 15	

Registrant - Vedco (021634266)

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

Revised: 12/2015 Vedco