DEXTROSE- dextrose injection, solution
Durvet

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%

For intravenous 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.

CAUTIONS

Intravenous administration must be done slowly and made under stric 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 After a quantity has been withdrawn for injection, the remainder should be discarded.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

ACTIVE INGREDIENTS:

Each 100 mL contains:

Dextrose . H_2O ..... 50 gms.

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

INDICATIONS:

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

KEEP OUT OF REACH OF CHILDREN

FOR USE IN ANIMALS ONLY

TAKE TIME. OBSERVE LABEL DIRECTIONS

REV 06-19

ISS19XB04

Manufactured for:

DURVET, INC.

Blue Springs, Missouri 64014

www.durvet.com

Principal Display Panel

NDC 30798-127-17

durvet

DEXTROSE 50% SOLUTION

Dura-Ster TS

TERMINALLY STERILIZED

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

NET CONTENTS: 500 mL

(16.9 fl. oz.)

DEXTROSE

dextrose injection, solution

Product Information			
Product Type	OTC ANIMAL DRUG	**Item Code (Source)**	NDC:30798-127
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, INTRAPERITONEAL, SUBCUTANEOUS		

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:30798-127-17	500 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
unapproved drug other		11/07/2019	

Labeler - Durvet (056387798)

Registrant - Durvet (056387798)

Revised: 11/2019 Durvet