

ULTIGENE- electrolyte solution with dextrose injection, solution

Vedco

Reference Label Set Id: c67c10cb-141e-496e-8ba8-64054bcf9ad1

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.

ELECTROLYTE SOLUTION WITH DEXTROSE

Composition

Each 500 mL of sterile aqueous solution contains:

Dextrose.H ₂ O	12.50g
Sorbitol	12.50g
Sodium Lactate	3.95g
Sodium Chloride	2.40g
Potassium Chloride	0.37g
Magnesium Chloride.6H ₂ O	0.21g
Calcium Chloride.2H ₂ O	0.19g
Milliequivalents per liter	
Cations	
Sodium	153mEq/L
Potassium	9mEq/L
Calcium	6mEq/L
Magnesium	4mEq/L
Anions	
Chloride	101 mEq/L
Lactate	71 mEq/L
Osmolarity (calc.)	602 mOsmol/L

Indications

For use in conditions associated with fluid and electrolyte loss such as dehydration, shock, vomiting, and diarrhea, particularly when an immediate source of energy is also indicated

Contraindications

Do not administer intraperitoneally to horses

Dosage and Administration

This product contains no preservatives. Use entire contents when first opened. Discard any unused solution.

Warm solution to body temperature and administer slowly (10 to 30 mL per minute) by intravenous or

intraperitoneal injections, using strict aseptic procedures.

Adult Cattles and Horses - 1000 to 2000 mL

Calves, Ponies, and Foals - 500 to 1000 mL

Adult, Sheep and Swine - 500 to 1000 mL

These are suggested dosages. The actual amount and rate of fluid administration must be judged by the veterinarian in relation to the condition being treated and the clinical response of the animal, being careful to avoid overhydration

FOR ANIMAL USE ONLY

Warning

KEEP OUT OF REACH OF CHILDREN

CAUTION:

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

Storage

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



Distributed By:

VEDCO INC

St. Joseph, MO, 64507

Made in El Salvador



Principal Display Panel

NDC 50989-886-16

ULTIGIENE™TM TERMINALLY STERILIZED

ELECTROLYTE SOLUTION WITH DEXTROSE

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

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Rev.7/19

1.82" (46.23 mm) por 0.437" (11.1 mm)
área sin barniz

NDC 50989-886-17

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ELECTROLYTE SOLUTION WITH DEXTROSE

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Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:50989-886
Route of Administration	INTRAVENOUS, INTRAPERITONEAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Dextrose Monohydrate (UNII: LX22YL083G) (Anhydrous Dextrose - UNII:5SL0G7R0OK)	Dextrose Monohydrate	12.50 g in 500 mL
Sorbitol (UNII: 506T60A25R) (Sorbitol - UNII:506T60A25R)	Sorbitol	12.50 g in 500 mL
Sodium Lactate (UNII: TU7HW0W0QT) (Sodium Cation - UNII:L4R4M0NH37)	Sodium Lactate	3.95 g in 500 mL
Sodium Chloride (UNII: 451W47IQ8X) (Chloride Ion - UNII:Q32ZN48698)	Sodium Chloride	2.40 g in 500 mL
Potassium Chloride (UNII: 660YQ98I10) (Potassium Cation - UNII:295O53K152)	Potassium Chloride	0.37 g in 500 mL
Magnesium Chloride (UNII: 02F3473H9O) (Magnesium Cation - UNII:T6V3LHY838)	Magnesium Chloride	0.21 g in 500 mL
Calcium Chloride (UNII: M4I0D6VV5M) (Calcium Cation - UNII:2M83C4R6ZB)	Calcium Chloride	0.19 g in 500 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50989-886-16	500 mL in 1 BOTTLE, PLASTIC		
2	NDC:50989-886-17	1000 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/05/2019	

Labeler - Vedco (021634266)**Registrant** - Vedco (021634266)

Revised: 8/2019

Vedco