DEXTROLYTE - electrolyte with dextrose injection, solution 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.

DextroLyte

For animal use only. Keep out of reach of children and pets.

CAUTION:

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

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.

DOSAGE AND ADMINISTRATION:

Warm solution to body temperature and administer slowly (10 to 30 mL per minute) by intravenous or intraperitoneal injection, using strict aseptic procedures.

Adult Cattle 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.

CONTRAINDICTIONS:

Do not administer intraperitoneally to horses.

CONTAINS:

Each 500 mL of sterile aqueous solution contains:

Dextrose.H2O	12.50 g
Sorbitol	12.50 g
Sodium Lactate	3.95 g
Sodium Chloride	2.40 g
Potassium Chloride	0.37 g
Magnesium Chloride.6H2O	0.21 g
Calcium Chloride.2H2O	0.19 g
Milliequivalents per liter Cations	

Sodium 153 mEq/L

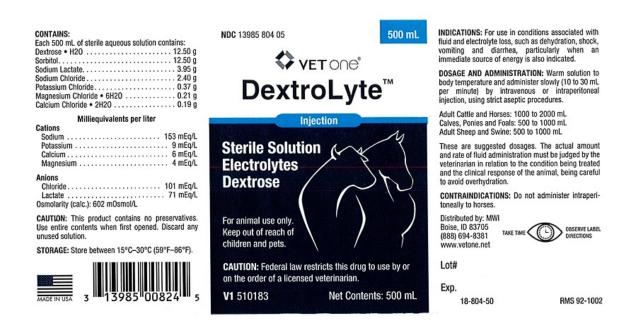
Potassium	9 mEq/L
Calcium	6 mEq/L
Magnesium	4 mEq/L
Anions	
Chloride	101 mEq/L
Lactate	71 mEq/L
Osmolarity (calc.): 602 mOsmol/L	•

CAUTION:

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

STORAGE:

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



DEXTROLYTE

electrolyte with dextrose injection, solution					
Product Information					
Product Type	PRESCRIPTION ANIMAL DRUG Item Code (Source) NDC:13		NDC:13985-804		
Route of Administration	INTRAVENOUS, INTRAPERITONEAL				
Active Ingredient/Active Moiety					
Ingredient Name		Basis of Strength	Strength		

Dextrose Monohydrate (UNII: LX22YL083G) (Dextrose - UNII:IY9 XDZ35W2)	Dextrose Monohydrate	12.50 g in 500 mL
Sorbitol (UNII: 506T60A25R) (Sorbic Acid - UNII:X045WJ989B)	Sorbitol	12.50 g in 500 mL
Sodium Lactate (UNII: TU7HW0W0QT) (Lactic Acid - UNII:33X04XA5AT)	Sodium Lactate	3.95 g in 500 mL
Sodium Chloride (UNII: 451W47IQ8X) (Sodium Cation - UNII:LYR4M0NH37)	Sodium Chloride	2.40 g in 500 mL
Potassium Chloride (UNII: 660 YQ98 I10) (Potassium Cation - UNII:295053K152)	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: M4I0 D6 VV5M) (Calcium Cation - UNII:2M8 3C4R6 ZB)	Calcium Chloride	0.19 g in 500 mL

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13985-804-05	500 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/15/2011	

Labeler - MWI (VetOne) (019926120)

Registrant - MWI (VetOne) (019926120)

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

Revised: 8/2011 MWI (VetOne)