

LACTATED RINGERS- lactated ringers injection, solution
VetTek

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.

LACTATED RINGERS

KEEP OUT OF REACH OF CHILDREN

FOR ANIMAL USE ONLY

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

INDICATIONS:

For the correction of electrolyte depletion, metabolic acidosis and dehydration of cattle, calves, horses, sheep and swine.

DOSAGE AND ADMINISTRATION:

May be injected intravenously, subcutaneously or intraperitoneally (except in horses) using strict aseptic technique.

Cattle and Horses: 2 to 5 mL per pound of body weight depending on size and condition of animal, repeated 1 to 3 times daily or as needed.

Swine and Sheep: 2 to 5 mL per pound of body weight depending on size and condition of animal, repeated 1 to 3 times daily or as needed.

If administered subcutaneously divide the dosage into several sites of injection and massage points of injection to aid in absorption and help prevent inflammation and/or sloughing.

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

Manufactured For
VetTek
Blue Springs, MO
64014

REV 06-19

ISS19XB04

ACTIVE INGREDIENTS

Each 100 mL contains:

Sodium Chloride.....600 mg
Sodium Lactate.....310 mg
Potassium Chloride.....30 mg
Calcium Chloride Dihydrate....20 mg
Water for Injection.....q.s.

The Calcium, Potassium, Sodium, Chloride and Lactate contents are approximately 2.7, 4.0, 130, 109 and 28 mEq/liter, respectively.

Total Osmolar Concentration: 273 mOsm per liter (calculated).

CAUTION:

Solution should be warmed to body temperature prior to administration and administered at a slow rate. This is a single dose unit. It contains no preservatives. Use entire contents when first opened.

WARNING:

Do not administer to horses by intraperitoneal injection. Do not administer to animals with inadequate renal function. Not for use in lactic acidosis.

Principal Display Panel

NDC 60270-132-20

Lactated Ringers

Dura-Ster TS

TERMINALLY STERILIZED

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

KEEP OUT OF REACH OF CHILDREN

FOR ANIMAL USE ONLY

Net Contents: 1000 mL (33.8 fl. oz)

VETTEK™

Manufactured for: VetTek, Blue Springs, MO 64014

INDICATIONS: For the correction of electrolyte depletion, metabolic acidosis and dehydration of cattle, calves, horses, sheep and swine.

DOSAGE AND ADMINISTRATION: May be injected intravenously, subcutaneously or intraperitoneally (except in horses) using strict aseptic technique.

Cattle and Horses: 2 to 5 mL per pound of body weight depending on size and condition of animal, repeated 1 to 3 times daily or as needed.

Swine and Sheep: 2 to 5 mL per pound of body weight depending on size and condition of animal, repeated 1 to 3 times daily or as needed. If administered subcutaneously divide the dosage into several sites of injection and massage points of injection to aid in absorption and help prevent inflammation and/or sloughing.

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

Manufactured for:
VetTek
Blue Springs, MO
64014
REV 06-19
ISS19XB04

Each 100 mL contains:
Sodium Chloride 600 mg
Sodium Lactate 310 mg
Potassium Chloride 30 mg
Calcium Chloride Dihydrate 20 mg
Water for Injection q.s.

The Calcium, Potassium, Sodium, Chloride and Lactate contents are approximately 2.7, 4.0, 130, 109 and 28 mEq/L, respectively. Total Osmolar Concentration: 273 mOsm per liter (calculated).

CAUTION: Solution should be warmed to body temperature prior to administration and administered at a slow rate. This is a single dose unit. It contains no preservatives. Use entire contents when first opened.

WARNING: Do not administer to horses by intraperitoneal injection. Do not administer to animals with inadequate renal function. Not for use in lactic acidosis.

TAKE TIME OBSERVE LABEL DIRECTIONS

LACTATED RINGERS			
lactated ringers injection, solution			
Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:60270-132
Route of Administration	INTRAVENOUS, SUBCUTANEOUS,		

Route of Administration INTRAPERITONEAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	600 mg in 100 mL
SODIUM LACTATE (UNII: TU7HW0W0QT) (LACTIC ACID - UNII:33X04XA5AT)	SODIUM LACTATE	310 mg in 100 mL
POTASSIUM CHLORIDE (UNII: 660YQ98I10) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM CATION	30 mg in 100 mL
CALCIUM CHLORIDE (UNII: M4I0D6VV5M) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CHLORIDE	20 mg in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60270-132-20	1000 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/08/2019	

Labeler - VetTek (056387798)

Registrant - VetTek (056387798)