LACTATED RINGERS- lactated ringers injection, solution Nova-Tech, Inc.

Reference Label Set Id: e4d8dbc5-6c0e-4a3c-8efe-ed6dc365fc71

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

Composition

Each 100 mL of sterile aquious solution contains:

The Calcium, Potassium and Sodium contents are approximately 2.7, 4.0, and 130 mEq/liter,

respectively. Total Osomolar concentration: 269 mOsm per liter (calculated).

Each 100 mL contains Sodium Chloride 600 mg; Sodium Lactate 310 mg; Potassium Chloride 30 mg; Calcium Chloride, Dihydrate 20 mg in Water for Injection.

May contain HCL or NaOH for pH adjustment.

mEq/Liter: Sodium 130; Chloride 109; Lactate 28; Potassium 4.0; Calcium 2.7.

Osomolarity: 269 mOsmol/liter (calc.).

pH: 6.6 (6.0 - 7.5).

INDICATIONS

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

DOSAGE & 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.

As directed by a veterinarian. Dosage is dependent upon the age, weight and clinical condition of the patient as well as laboratory determinations.

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.

Solution should be warmed to body temperature prior to administration and administered at a slow rate. Sterile nonpyrogenic solution. Use only if solution is clear and container is undamaged. This is a single dose unit. It contains no preservatives. Use promptly upon initial entry. If entire contents are not used, discard unused portion. Not for use in the treatment of lactic acidosis. Squeeze and inspect inner bag which maintains product sterility. Discard if leaks are found or if the solution contains visible solid particles.

WARNING:

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

Store between 15°C-30°C (59°F-86°F)

TAKE TIME OBSERVE LABEL DIRECTIONS
KEEP OUT OF REACH OF CHILDREN

RX only.

STERILE SOLUTION

FOR ANIMAL USE ONLY

CAUTION:

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

Manufactured by:

Nova-Tech, Inc. Grand Island, NE 68801 USA

NDC# 65207-803-60

Nova-Tech®

Animal Health

Net Contents:

1000 mL (33.81 fl oz)

Lot No.

Exp. Date

Assembled in USA

NDC # 65207-803-61

18-803

RMS# 92-2111

365207803613

NDC # 65207-803-80

18-803

RMS 92-2112

365207803803

Indications

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

Dosage & 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 to help prevent inflammation and/or sloughing.



Manufactured by:

Nova-Tech, Inc. Grand Island, NE 68801 USA

18-803 RMS 92-360



Composition

Each 100 mL of sterile aqueo	
Sodium Chloride	600 mg
Sodium Lactate	
Potassium Chloride	
Calcium Chloride Dihydrate	20 mg
Water for Injection	q.s

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

Total Osmolar Concentration: 269 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.

Store between 15°C-30°C (59°F-86°F).



Lot No.

Exp. Date

LACTATED RINGER'S Injection

Approx.

200

FOR ANIMAL USE ONLY

1000 mL (33.81 fl. oz.)

ACTIVE INGREDIENTS: EACH 100 mL CONTAINS SODIUM

600 MG; LACTATE POTASSIUM CHLORIDE 30 mg: CALCIUM CHLORIDE. DIHYDRATE 20 mg IN WATER FOR INJECTION.

MAY CONTAIN HCI OR NaOH FOR PH ADJUSTMENT.

mEg/LITER: SODIUM 130; CHLORIDE 109;

LACTATE 28; POTASSIUM 4.0; CALCIUM 2.7.

OSMOLARITY: 269 mOsmol/LITER (CALC.).

pH: 6.6 (6.0 - 7.5).

DOSAGE AND ADMINISTRATION: AS DIRECTED BY A VETERINARIAN. DOSAGE IS DEPENDENT UPON TAGE, WEIGHT AND CLINICAL CONDITION OF THE PATIENT AS WELL AS LABORATORY DETERMINATIONS.

TO BE CAUTION: SOLUTION SHOULD WARMED BODY TEMPERATURE PRIOR **ADMINISTRATION** ONLY ADMINISTERED AT A SLOW RATE. NONPYROGENIC SOLUTION. USE CLEAR AND CONTAINER IS UNDAMAGED.
INGLE DOSE UNIT. IT CONTAINS NO
S. USE PROMPTLY UPON INITIAL SOLUTION IS THIS IS A SINGLE S. USE PROMP ENTIRE CONTENTS PRESERVATIVES. UNUSED PORTION.
ATMENT OF LACTIC A
INNER BAG WHICH USED, ENTRY. ARE NOT DISCARD USE FOR TOM THE TREATMENT ACIDOSIS. SQUEEZE INSPECT INNER BA MAINTAINS PRODUCT IF LEAKS FOUND ARE OR THE SOLUTION CONTAINS VISIBLE SOLID PARTICLES.

WARNING: DO TOM ADMINISTER TO HORSES BY INTRAPERITONEAL INJECTION.

STORAGE: 30°C. STORE BETWEEN 15°C KEEP

FROM FREEZING.

KEEP OUT OF REACH OF CHILDREN RX ONLY.

CAUTION: FEDERAL LAW RESTRICTS THIS USE BY OR ON THE ORDER OF LICENSED 800 A

VETERINARIAN. ASSEMBLED IN U.S.A.

NDC # 65207-803-61 RMS# 92-2111



400

600

LACTATED RINGERS

Approx. 1000

INJECTION

FOR ANIMAL USE ONLY 5000 mL (169.07 fl. Oz.)

ACTIVE INGREDIENTS: EACH 100 mL CONTAINS SODIUM CHLORIDE 600 mg; SODIUM LACTATE 310 mg; POTASSIUM CHLORIDE 30 mg; CALCIUM CHLORIDE, DIHYDRATE 20 mg IN WATER FOR INJECTION. MAY CONTAIN HCI OR NaOH FOR 2000 pH ADJUSTMENT.

mEq/LITER: SODIUM 130; CHLORIDE 109; LACTATE 28; POTASSIUM 4.0; CALCIUM 2.7.

OSMOLARITY: 269 mOsmol/LITER (CALC.).

pH: 6.6 (6.0—7.5).

DOSAGE AND ADMINISTRATION: AS DIRECTED BY A VETERINARIAN. DOSAGE IS DEPENDENT UPON THE AGE. WEIGHT AND CLINICAL CONDITION OF PATIENT AS WELL AS LABORATORY DETERMINATIONS. THE

CAUTION: SOLUTION SHOULD BE WARMED TO BODY TEMPERATURE PRIOR TO ADMINISTRATION AND ADMINISTERED AT A SLOW RATE. NONPYROGENIC SOLUTION. USE ONLY IF SOLUTION IS CLEAR AND CONTAINER IS UNDAMAGED. THIS IS A SINGLE DOSE UNIT. IT CONTAINS NO PRESERVATIVES. USE PROMPTLY UPON INITIAL ENTRY. IF ENTIRE CONTENTS ARE NOT USED, DISCARD UNUSED PORTION. NOT FOR USE IN THE TREATMENT OF LACTIC SQUEEZE AND INSPECT INNER BAG WHICH **MAINTAINS** PRODUCT STERILITY. DISCARD IF LEAKS ARE FOUND OR IF THE SOLUTION **CONTAINS VISIBLE SOLID PARTICLES.**

WARNING: DO NOT ADMINISTER TO HORSES BY INTRAPERITONEAL INJECTION.

STORAGE: STORE BETWEEN 15°C - 30°C. KEEP FROM FREEZING.

KEEP OUT OF REACH OF CHILDREN





RX ONLY

CAUTION: FEDERAL LAW RESTRICTS THIS DRUG TO USE BY OR ON THE ORDER OF A LICENSED VETERINARIAN. 18-803

4000

RMS# 92-2112

ASSEMBLED IN U.S.A. NDC# 65207-803-80





LACTATED RINGERS

lactated ringers injection, solution

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:65207- 803
Route of Administration	INTRAVENOUS, SUBCUTANEOUS, 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:295053K152)	POTASSIUM CATION	30 mg in 100 mL	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M) (CALCIUM CATION - UNII: 2M83C4R6ZB)	CALCIUM CHLORIDE	20 mg in 100 mL	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:65207-803-60	1000 mL in 1 BOTTLE, PLASTIC			
2	NDC:65207-803-61	1000 mL in 1 BAG			
3	NDC:65207-803-80	5000 mL in 1 BAG			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		03/18/2020		

Labeler - Nova-Tech, Inc. (196078976)

Revised: 1/2024 Nova-Tech, Inc.