

LACTATED RINGERS - lactated ringers injection, solution

A & G Pharmaceuticals, Inc.

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 Injection

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

FOR ANIMAL USE ONLY

KEEP OUT OF REACH OF CHILDREN

NET CONTENTS: 1000 mL

Sterile Nonpyrogenic Solution

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 - 30°C (59°F - 86°F)

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 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.

Manufactured by

Nova-Tech, Inc.

Grand Island, NE 68801

TAKE TIME OBSERVE LABEL DIRECTIONS

18-803-60

RMS 92-559

Iss. 04-09

Manufactured for:

A&G Pharmaceuticals

Clarksburg, NJ 08510

Lot No.

Exp. Date

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Product Information

Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:57699-803
Route of Administration	INTRAVENOUS, SUBCUTANEOUS, INTRAPERITONEAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Sodium Chloride (UNII: 451W47IQ8X) (Sodium Cation - UNII:LYR4M0NH37)	Sodium Chloride	600 mg in 1000 mL
Sodium Lactate (UNII: TU7HW0W0QT) (Sodium Cation - UNII:LYR4M0NH37)	Sodium Lactate	310 mg in 1000 mL
Potassium Chloride (UNII: 660YQ98I10) (Potassium Cation - UNII:295O53K152)	Potassium Chloride	30 mg in 1000 mL
Calcium Chloride (UNII: M4I0D6VV5M) (Calcium Cation - UNII:2M83C4R6ZB)	Calcium Chloride	20 mg in 1000 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57699-803-60	1000 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/25/2013	

Labeler - A & G Pharmaceuticals, Inc. (182147033)

Revised: 4/2024

A & G Pharmaceuticals, Inc.