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 Sterile Nonpyrogenic Solution

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**

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 degrees C - 30 degrees C (59 degrees F - 86 degrees F)

EACH 100 mL CONTAINS:

Sodium Chloride......600 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 mOsmol 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

Iss. 04-09

Manufactured for:

A and G Pharmaceuticals

Clarksburg, NJ 08510

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

Manufactured by Nova-Tech, Inc. Grand Island, NE 68801





Manufactured for:

EACH 100 mL CONTAINS:

Lot No.

Sodium Chloride		
Sodium Lactate	310	mg
Potassium Chloride	. 30	mg
Calcium Chloride Dihydrate	20	mg
Water for Injection		q.s.
The Calcium, Potassium and Sodium	conte	ents

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Exp. Date



A&G Pharmaceuticals Clarksburg, NJ 08510

actated ringers injection, sol	ution			
Product Information				
Product T ype	PRESCRIPTION ANIMAL DRUG	PRESCRIPTION ANIMAL DRUG		NDC:57699- 803
Route of Administration	of Administration INTRAVENOUS, SUBCUTANEOUS, INTRAPERITONEAL			
Active Ingredient/Active	Moiety			
0	Moiety Ingredient Name	Basi	s of Strength	Strength
5	5		s of Strength m Chloride	
Sodium Chloride (UNII: 451W47	Ingredient Name	Sodiu	0	Strength 600 mg in 1000 mL 310 mg in 1000 mL
Sodium Chloride (UNII: 451W47 Sodium Lactate (UNII: TU7HW0	Ingredient Name IQ8X) (Sodium Cation - UNII:LYR4M0NH37)	So diui So diui	m Chloride	600 mg in 1000 mL

# Item	Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:57699	-803-60	1000 mL in 1 BOTTLE, PLASTIC			
Marketing Information					
Marketin	g Inforn	nation			
Marketin Marketing C	U	1ation Application Number or Monograph Cita	ation Marketing Start D	ate Marketing End Date	
	ategory		ation Marketing Start Data 0 1/25/20 13	ate Marketing End Date	

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

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

Establishment

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

Establishment						
Name	Address	ID/FEI	Business Operations			
K+S KALI GmbH		507531213	api manufacture			

Revised: 2/2015

A & G Pharmaceuticals, Inc.