# 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.

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#### 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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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		<b>s of Strength</b> m Chloride	
Sodium Chloride (UNII: 451W47	Ingredient Name	Sodiu	0	<b>Strength</b> 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	<b>U</b>	<b>1ation</b> 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.