

**A-LYTE ORAL- amino acid solution**  
**Durvet, 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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**A-Lyte Solution**

FOR USE IN ANIMALS ONLY

KEEP OUT OF REACH OF CHILDREN

An oral source of vitamins, amino acids and electrolytes for cattle, swine, sheep and horses when dietary intake is reduced.

For use in drinking water

Supply fresh drinking water daily.

**DOSAGE UNDILUTED**

Cattle: Administer 1 oz. A-Lyte Solution per 10 pounds of body weight in the drinking water to be consumed in one day.

Horses: Administer 10 oz. A-Lyte Solution per 100 pounds of body weight in the drinking water to be consumed in one day.

Sheep and Swine: Administer 1/2 oz. A-Lyte Solution per 5 pounds body weight in the drinking water to be consumed in one day.

PROTECT FROM FREEZING

Store in controlled room temperature between 15° and 30°C (59°-86°F).

TAKE TIME OBSERVE LABEL DIRECTIONS

**COMPOSITION**

Each 100 mL of aqueous solution contains:

**ACTIVE INGREDIENTS:**

- Dextrose•H<sub>2</sub>O ..... 5 g
- Sodium Acetate•3H<sub>2</sub>O ..... 250 mg
- Magnesium Sulfate•7H<sub>2</sub>O ..... 20 mg
- Potassium Chloride ..... 20 mg
- Calcium Chloride•2H<sub>2</sub>O ..... 15 mg

**INACTIVE INGREDIENTS:** Niacinamide, Pyridoxine HCl (B<sub>6</sub>), d-Panthenol, Riboflavin (B<sub>2</sub>), Cyanocobalamin (B<sub>12</sub>), L-Argenine HCl, L-Cysteine HCl H<sub>2</sub>O, L-Glutamic Acid, L-Histidine HCl H<sub>2</sub>O, L-Isoleucie, L-Leucine, L-Lycine HCl, L-Methionine, L-Phenylalamine, L-Theonine, L-Valine with Methylparaben 0.18%, Ethylparaben 0.01%, Propylparaben 0.02%.

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Manufactured for:  
**DURVET, INC.**  
Blue Springs, Missouri 64014  
www.durvet.com



NDC 30798-001-17



**A-Lyte  
Solution**

An oral source of vitamins,  
amino acids and electrolytes



**NET CONTENTS:  
500 mL  
(16.9 FL. OZ.)**

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OF CHILDREN**

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A-0010-06      REV 07-16  
Lot No.:              EXP. DATE:



**A-LYTE ORAL**

amino acid solution

**Product Information**

<b>Product Type</b>	OTC ANIMAL DRUG	<b>Item Code (Source)</b>	NDC:30798-001
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>DEXTROSE MONOHYDRATE</b> (UNII: LX22YL083G) (ANHYDROUS DEXTROSE - UNII:5SL0G7R0OK)	DEXTROSE MONOHYDRATE	5000 mg in 100 mL
<b>CALCIUM CHLORIDE ANHYDROUS</b> (UNII: OFM21057LP) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CATION	15 mg in 100 mL
<b>MAGNESIUM SULFATE HEPTAHYDRATE</b> (UNII: SK47B8698T) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM SULFATE HEPTAHYDRATE	20 mg in 100 mL
<b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10) (POTASSIUM CATION - UNII:295053K152, CHLORIDE ION - UNII:Q32ZN48698)	POTASSIUM CHLORIDE	20 mg in 100 mL
<b>SODIUM ACETATE ANHYDROUS</b> (UNII: NVG71ZZ7P0) (ACETATE ION - UNII:569DQM745C, SODIUM CATION - UNII:LYR4M0NH37)	SODIUM ACETATE ANHYDROUS	250 mg in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>VALINE</b> (UNII: HG18B9YRS7)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:30798-001-17	500 mL in 1 BOTTLE		
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
unapproved drug other		08/11/1998		

**Labeler** - Durvet, Inc. (056387798)

Revised: 1/2017

Durvet, Inc.