AMINO ACID ORAL- amino acid solution Agri Laboratories, Ltd.

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.

AMINO ACID ORAL SOLUTION

For Animal Use Only

Keep Out of Reach of Children

INDICATIONS

For use as a supplemental nutritive source of amino acids, electrolytes, B complex vitamins, and dextrose in cattle and horses.

DOSAGE AND ADMINISTRATION

Administer orally as a drench. The usual recommended dose in adult cattle and horses is 500 to 3000 mL, depending on size and condition.

Not for human use.

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

TAKE TIME OBSERVE LABEL DIRECTIONS

COMPOSITION

Each 100 mL of aqueous	solution contains:
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Dextrose•H ₂ O	5 g
Sodium Acetate•3H ₂ O	250 mg
Magnesium Sulfate•7H ₂ O	20 mg
Potassium Chloride	20 mg
Calcium Chloride•2H ₂ O	15 mg

Comprised of: Niacinamide, Pyridoxine HCl (B_6), d-Panthenol, Riboflavin (B_2), Cyanocobalamin (B_{12}), L-Argenine HCl, L-Cysteine HCl H $_2$ O, L-Glutamic Acid, L-Histidine HCl H $_2$ 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%.

COMPOSITION:

Each 100 mL of aqueous solution contains:

Comprised of: Niacinamide, Pyridoxine HCI, Thiamine HCI, d-Panthenol, Riboflavin, Cyanocobalamin (B_{12}), L-Argenine HCI, L-Cysteine HCI H $_2$ O, L-Glutamic Acid, L-Histidine HCI H $_2$ O, L-Isoleucine, L-Leucine, L-Lycine HCI, L-Methionine, L-Phenylalanine, L-Threonine, L-Valine, with Methylparaben 0.18%, Ethylparaben 0.01%, Propylparaben 0.02%.

Manufactured for: Agri Laboratories, Ltd. St. Joseph, MO 64503

LOT NO.:

EXP. DATE:

NDC 57561-101-06

Amino Acid Oral Solution

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> Net Contents: 500 mL (16.9 oz)



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OBSERVE LABEL DIRECTIONS

A-0010-06

Rev. 09-12

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AMINO ACID ORAL

amino acid solution

Product Information

Product TypeOTC ANIMAL DRUGItem Code (Source)NDC:57561-101

Route of Administration ORAL

Active Ingredient/Active Moiety Basis of Strength Ingredient Name Strength DEXTROSE MONOHYDRATE (UNII: LX22YL083G) (ANHYDROUS DEXTROSE -DEXTROSE 5000 mg MONOHYDRATE in $100\ mL$ UNII:5SL0G7R0OK) CALCIUM CHLORIDE ANHYDROUS (UNII: OFM21057LP) (CALCIUM CATION -15 mg CALCIUM CATION UNII:2M83C4R6ZB) in 100 mL MAGNESIUM SULFATE HEPTAHYDRATE (UNII: SK47B8698T) (MAGNESIUM CATION MAGNESIUM SULFATE 20 mg HEPTAHYDRATE in 100 mL - UNII:T6 V3LHY838) POTASSIUM CHLORIDE (UNII: 660 YQ98 I10) (POTASSIUM CATION -20 mg POTASSIUM CHLORIDE UNII:295O53K152, CHLORIDE ION - UNII:Q32ZN48698) in 100 mL 250 mg SODIUM ACETATE ANHYDROUS (UNII: NVG71ZZ7P0) (ACETATE ION -SODIUM ACETATE UNII:569 DQM74SC, SODIUM CATION - UNII:LYR4M0 NH37) **ANHYDROUS** in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
VALINE (UNII: HG18B9 YRS7)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date

1 NDC:57561-101-06	500 mL in 1 BOTTLE					
Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
unapproved drug other		08/11/1995				

Labeler - Agri Laboratories, Ltd. (155594450)

Revised: 1/2017 Agri Laboratories, Ltd.