

AMINO ACID - amino acid concentrate

Agri Laboratoies, 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 CONCENTRATE

ORAL SOLUTION

For Animal Use Only

Keep Out of Reach of Children

INDICATIONS

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

DOSAGE UNDILUTED

Administer orally as a drench. The usual recommended dose in adult cattle and horses is 50 to 500 mL, depending on size and condition.

Not for Human use

STORE BETWEEN 15° and 30° C (59° and 86°F)

TAKE TIME OBSERVE LABEL DIRECTIONS

CONTENTS

Each 100 mL of aqueous solution contains:

Dextrose • H₂O 5 g
Sodium Acetate • 3H₂O 250 mg
Magnesium Sulfate • 7H₂O ... 200 mg
Potassium Chloride 200 mg
Calcium Chloride • 2H₂O150 mg

Comprised of : Niacinamide, Pyridoxine Hydrochloride (B₆), Thiamine Hydrochloride (B₁), d-Panthenol, Riboflavin (B₂), Cyanocobalamin (B₁₂), L-Leucine, L-Lysine Hydrochloride, L-Glutamic Acid, L-Valine, L-Phenylalanine, L-Arginine Hydrochloride, L-Isoleucine, L-Threonine, L-Histidine Hydrochloride • H₂O, L-Methionine, L-Cysteine Hydrochloride • H₂O, with propylne glycol 2.5%, sorbitol 2.5%, lactic acid 0.16%, citric acid 0.1%, BHA 0.005%, methylparaben 0.18%, propylparaben 0.02%, and ehtylparaben 0.01% (preservatives).

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Not for Human use

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

A-0020-06

Rev. 09-12



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

LOT NO.:

EXP. DATE:

NDC 57561-102-06

AMINO ACID CONCENTRATE

Oral Solution

For Animal Use Only

Keep Out of Reach of Children

**Net Contents:
500 mL (16.9 oz)**



COMPOSITION:

Each 100 mL of aqueous solution contains:

Dextrose • H₂O 5 g
Sodium Acetate • 3H₂O 250 mg
Magnesium Sulfate • 7H₂O 200 mg
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AMINO ACID

amino acid concentrate

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:57561-102
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CHLORIDE ANHYDROUS (UNII: OFM21057LP) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CATION	150 mg in 100 mL
DEXTROSE MONOHYDRATE (UNII: LX22YL083G) (ANHYDROUS DEXTROSE - UNII:5SL0G7R0OK)	DEXTROSE MONOHYDRATE	5 g in 100 mL
MAGNESIUM SULFATE HEPTAHYDRATE (UNII: SK47B8698T) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM SULFATE HEPTAHYDRATE	200 mg in 100 mL
POTASSIUM CHLORIDE (UNII: 660YQ981I0) (CHLORIDE ION - UNII:Q32ZN48698, POTASSIUM CATION - UNII:295O53K152)	POTASSIUM CHLORIDE	200 mg in 100 mL
SODIUM ACETATE ANHYDROUS (UNII: NVG71ZZ7P0) (SODIUM CATION - UNII:LYR4M0NH37, ACETATE ION - UNII:569DQM74SC)	SODIUM ACETATE ANHYDROUS	250 mg in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57561-102-06	500 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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unapproved drug other		09/01/1991	
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Labeler - Agri Laboratoies, Ltd. (155594450)

Revised: 6/2017

Agri Laboratoies, Ltd.