MAG-AL LIQUID- aluminum hydroxide and magnesium hydroxide suspension PAI Holdings, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Mag-AL Liquid

Drug Facts

Active ingredients (in each 5 mL = 1 teaspoonful) Purpose Aluminum hydroxide (equiv. to dried gel, USP) 200 mg Antacid Magnesium hydroxide 200 mg Antacid

Uses

for the relief of:

- · acid indigestion
- heartburn
- sour stomach
- upset stomach due to these symptoms

Warnings

Do not take more than 16 teaspoonfuls in a 24-hour period or use the maximum dosage for more than 2 weeks except under the advice and supervision of a physician.

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet

Ask a doctor or pharmacist before use if you are

• presently taking a prescription drug. Antacids may interact with certain prescription drugs.

Stop use and ask a doctor if

• symptoms last more than 2 weeks.

Keep out of reach of children.

Directions

- shake well before using
- do not take more than 16 teaspoonfuls in 24 hours or use the maximum dosage for more than 2 weeks

adults and children 12 years and older	take 2 to 4 teaspoonfuls four times a day	
	or as directed by a physician	
children under 12 years	consult a physician	

Other information

- each 5 mL contains: magnesium 83 mg, sodium 1.34 mg
- does not meet USP requirements for preservative effectiveness
- store at controlled room temperature 20° 25°C (68° 77°F)
- protect from freezing
- White colored, peppermint flavored liquid supplied in the following oral dosage form:

NDC 0121-1760-30:30 mL unit dose cup, in a tray of ten cups.

Inactive ingredients

Antifoam af emulsion, butylparaben, hydroxypropyl methylcellulose, flavoring, propylene glycol, propylparaben, purified water, sodium saccharin, and sorbitol solution.

Questions or comments?

Call 1-800-845-8210

MANUFACTURED BY

Pharmaceutical Associates, Inc. Greenville, SC 29605

R07/21

PRINCIPAL DISPLAY PANEL - 30 mL Dose Cup Tray Label

Delivers 30 ML

NDC 0121-1760-30

MAG-AL Liquid

Each 30 mL contains:

Magnesium Hydroxide 1200 mg

Aluminum Hydroxide 1200 mg

SHAKE WELL

Package Not Child-Resistant



MAG-AL LIQUID

aluminum hydroxide and magnesium hydroxide suspension

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0121-1760	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	200 mg in 5 mL		
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	200 mg in 5 mL		

Inactive Ingredients		
Ingredient Name	Strength	
BUTYLPARABEN (UNII: 3QPI1U3FV8)		
PROPYLPARABEN (UNII: Z8IX2SC10H)		
HYPROMELLOSE 2910 (4000 MPA.S) (UNII: RN31520P35)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
SORBITOL (UNII: 506T60A25R)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
PEPPERMINT OIL (UNII: AV092KU4JH)		
CYCLOMETHICONE 4 (UNII: CZ227117JE)		
WATER (UNII: 059QF0KO0R)		

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor	PEPPERMINT	Imprint Code	
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:0121- 1760-30	10 in 1 CASE	01/14/2004		
1		10 in 1 TRAY			
1		30 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	01/14/2004	

Labeler - PAI Holdings, LLC (044940096)

Establishment				
Name	Address	ID/FEI	Business Operations	
PAI Holdings, LLC dba Pharmaceutical Associates, Inc.		097630693	manufacture(0121-1760)	

Revised: 3/2023 PAI Holdings, LLC