

**ALUMINUM HYDROXIDE, MAGNESIUM HYDROXIDE AND SIMETHICONE
SUSPENSION- aluminum hydroxide, magnesium hydroxide, and
simethicone suspension**

**ALUMINUM HYDROXIDE, MAGNESIUM HYDROXIDE AND SIMETHICONE
SUSPENSION MAXIMUM STRENGTH- aluminum hydroxide, magnesium
hydroxide, and simethicone suspension**

Major Pharmaceuticals

**Magnesium Hydroxide, Aluminum Hydroxide and Simethicone Suspension
1,200-1,200-120 mg/30 mL,
Magnesium Hydroxide, Aluminum Hydroxide and Simethicone Suspension
Maximum Strength 2,400-2,400-240 mg/30 mL**

Drug Facts

1,200-1,200-120 mg/30 mL

Active ingredients (in each 30 mL)

Purpose

Aluminum hydroxide 1,200 mg (equivalent to dried gel, USP).....Antacid
Magnesium hydroxide 1,200 mg.....Antacid
Simethicone 120 mg.....Antigas

2,400-2,400-240 mg/30 mL

Active ingredient (in each 30 mL)

Purpose

Aluminum hydroxide 2,400 mg (equivalent to dried gel, USP).....Antacid
Magnesium hydroxide 2,400 mg.....Antacid
Simethicone 240 mg.....Antigas

Usesrelieves • heartburn • sour stomach • acid indigestion • the symptoms referred to
as gas

Warnings

Ask a doctor before use if you have• kidney disease • a magnesium restricted diet

Ask a doctor or pharmacist before use if you aretaking another medication. This
product may interact with certain medications.

Stop use and ask a doctor ifsymptoms last more than 2 weeks

If pregnant or breast-feeding,ask a health professional before use.

Keep out of reach of children.

Directions • shake well before use

1,200-1,200-120 mg/30 mL

age	dose
Adults and children 12 years and older	30mL, not more than 120 mL in 24 hrs

Children under 6 years of age

Do not use unless directed by a doctor

2,400-2,400-240 mg/30 mL

age	dose
Adults and children 12 years and older	30mL, not more than 60 mL in 24 hrs
Children under 6 years of age	Do not use unless directed by a doctor

Other information

1,200-1,200-120 mg/30 mL

- **each 30 mL contains** magnesium 510 mg and sodium 18 mg
- protect from freezing
- store at 20° to 25°C (68° to 77°F).
- do not use if lid seal is open or damaged
- retain this insert for full product information

2,400-2,400-240 mg/30 mL

- **each 30 mL contains** magnesium 990 mg, sodium 30 mg
- store at 20° to 25°C (68° to 77°F).
- protect from freezing
- do not use if lid seal is open or damaged
- retain this insert for full product information

Inactive ingredients benzyl alcohol, butylparaben, carboxy-methylcellulose sodium, flavor, hypromellose, microcrystalline cellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or comments? (855) 879-4368

Distributed by:

MAJOR® PHARMACEUTICALS
Indianapolis, IN 46268

(800) 616-2471

Rev. 06/23

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

MAJOR®

NDC 0904-7325-62

Each 30 mL Contains:

Magnesium hydroxide 1,200 mg

Aluminum hydroxide 1,200 mg

Simethicone 120 mg

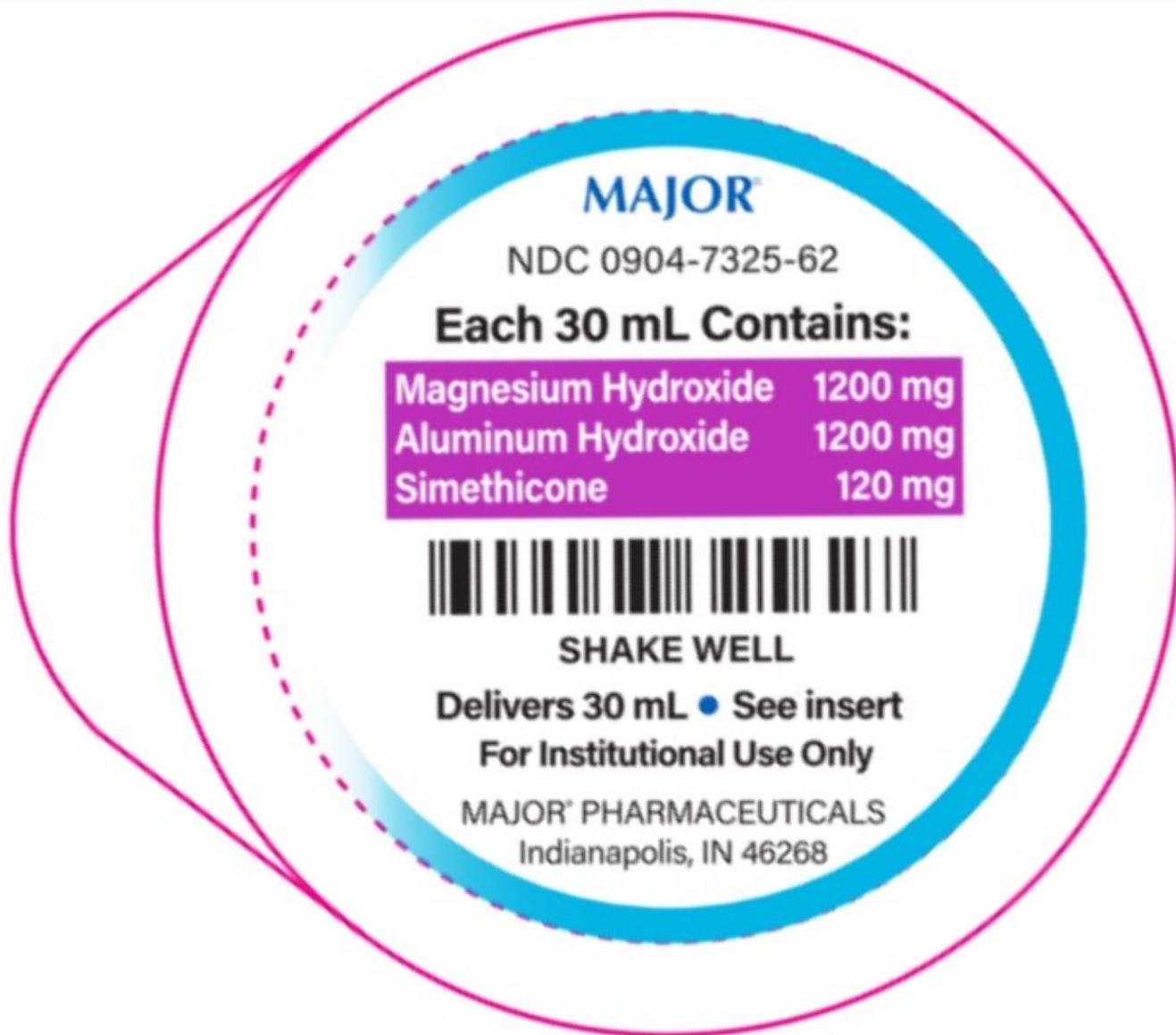
SHAKE WELL

Delivers 30 mL • See insert

For Institutional Use Only

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Indianapolis, IN 46268



PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

MAJOR®

NDC 0904-7326-62

Each 30 mL Contains:

Magnesium hydroxide 2,400 mg

Aluminum hydroxide 2,400 mg

Simethicone 240 mg

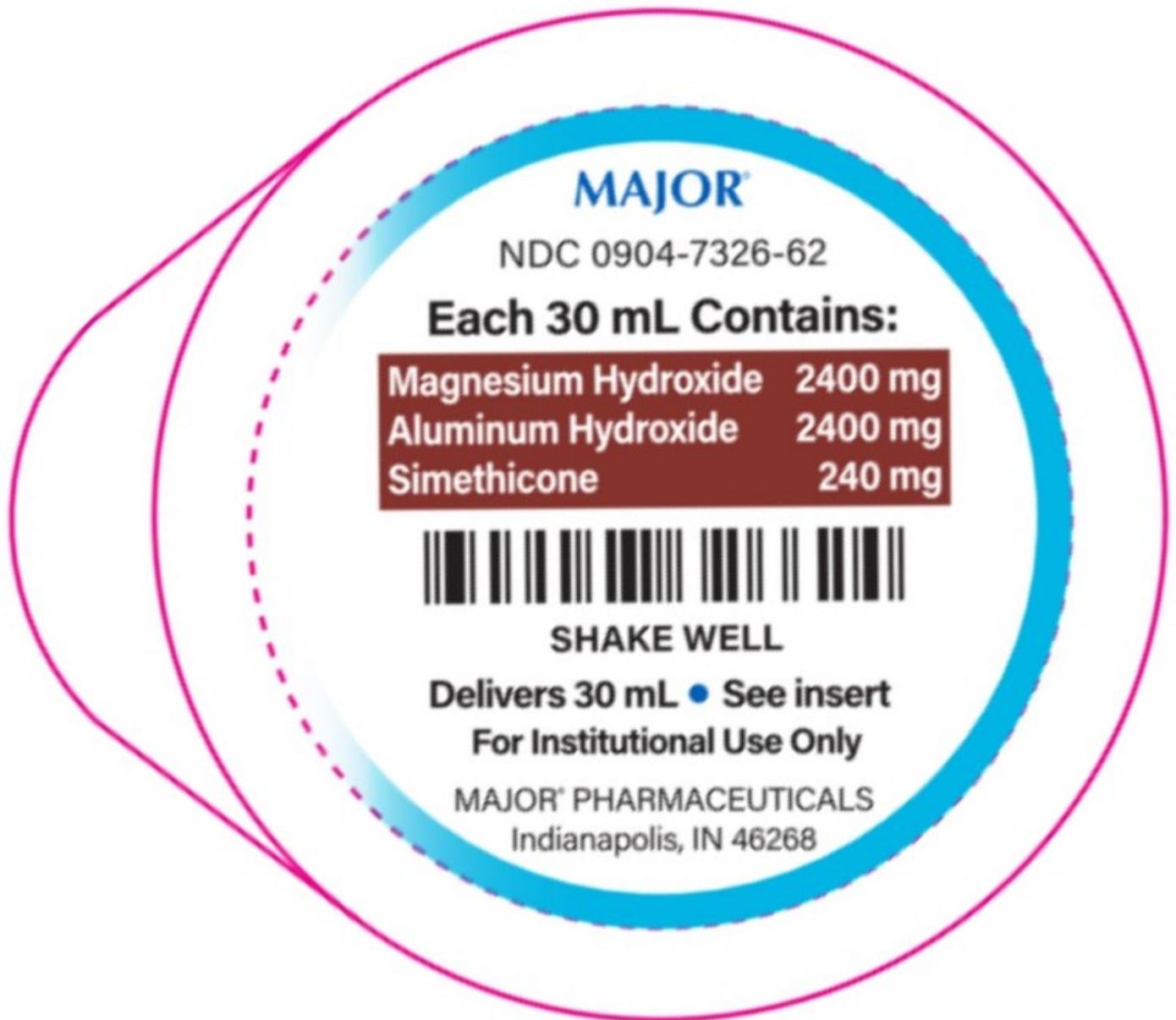
SHAKE WELL

Delivers 30 mL • See insert

For Institutional Use Only

MAJOR® PHARMACEUTICALS

Indianapolis, IN 46268



ALUMINUM HYDROXIDE, MAGNESIUM HYDROXIDE AND SIMETHICONE SUSPENSION

aluminum hydroxide, magnesium hydroxide, and simethicone suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-7325
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	1200 mg in 30 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	1200 mg in 30 mL
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y1O) (DIMETHICONE, UNSPECIFIED - UNII:92RU3N3Y1O)	DIMETHICONE, UNSPECIFIED	120 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-7325-73	10 in 1 CASE	08/01/2023	04/30/2026
1		10 in 1 TRAY		
1	NDC:0904-7325-62	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
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ALUMINUM HYDROXIDE, MAGNESIUM HYDROXIDE AND SIMETHICONE SUSPENSION MAXIMUM STRENGTH

aluminum hydroxide, magnesium hydroxide, and simethicone suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-7326
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	2400 mg in 30 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	2400 mg in 30 mL
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y10) (DIMETHICONE, UNSPECIFIED - UNII:92RU3N3Y10)	DIMETHICONE, UNSPECIFIED	240 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
PROPYLPARABEN (UNII: Z8IX25C1OH)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-7326-73	10 in 1 CASE	08/01/2023	11/30/2024
1		10 in 1 TRAY		

1	NDC:0904-7326-62	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 Drug	M001	08/01/2023	11/30/2024

Labeler - Major Pharmaceuticals (191427277)

Revised: 9/2025

Major Pharmaceuticals