MINTOX MAXIMUM STRENGTH- aluminum hydroxide, magnesium hydroxide, dimethicone suspension
MAJOR Pharmaceuticals Inc.

MAJOR MINTOX MAX

Active ingredients (in each 5 mL teaspoonful)

Aluminum hydroxide 400 mg (equivalent to dried gel, USP)

Magnesium hydroxide 400 mg

Simethicone 40mg

Purposes

Antacid

Antigas

Uses

relieves

- 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 are taking a prescription drug.

Antacids may interact with certain prescription drugs.

Stop use and ask a doctor if symptoms 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 using
- adults and children 12 years and older: take 2 to 4 teaspoonfuls two times a day or as directed by a physician
- do not take more than 8 teaspoonfuls in 24 hours or use the maximum dosage for

more than 2 weeks.
- children under 12 years: consult a physician

Other information
- **each 5 mL teaspoonful contains:** magnesium 165 mg, sodium 5 mg
- keep tightly closed
- store at room temperature and avoid freezing

Inactive ingredients

benzyl alcohol, butylparaben, caramel color, carboxymethylcellulose sodium, D and C yellow no.10, flavor, glycerin, hypromellose, microcrystalline cellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

package Label

NDC 0904-5725-14

MAJOR

Mintox

Maximum Strength

antacid/antigas

Alumina, Magnesia and Simethicone Oral Suspension USP

Fast Relief of:

HEARTBURN

ACID INDIGESTION

PRESSURE AND BLOATING

Lemon Creme

*compare to the active ingredients of Maalox Advanced Maximum Strength

12 FL OZ (355 mL)

NDC 0904-5725-14

MAJOR®

Mintox™

Maximum Strength

Antacid/Antigas

Alumina, Magnesia, and Simethicone Oral Suspension USP

Fast Relief of:

- Heartburn
- Acid Indigestion
- Pressure and Bloating

Lemon Creme

Compare to the active ingredients of Maalox® Advanced Maximum Strength*

12 FL OZ (355 mL)

226-11522-4

Drug Facts

TAMPER-EVIDENT: Do not use if the breakaway band on bottle cap is missing or broken.

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Aluminum hydroxide 400 mg (equivalent to dried gel USP)	Antacid
Magnesium hydroxide 400 mg	Antacid
Simethicone 40 mg	Antigas

Uses relieves:
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- the symptoms referred to as gas

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926-11522-6

*Major® Pharmaceuticals is not affiliated with the owner of the registered trademark MAALOX®.

Distributed by
MAJOR® PHARMACEUTICALS
17177 N Laurel Park Drive, Suite 233
Livonia, MI 48152

Questions or comments?
CALL: 800-616-2471

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Rev 11/16

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MINTOX MAXIMUM STRENGTH

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0904-5725
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE -	ALUMINUM	400 mg

UNII:5QB0T2IUN0)	HYDROXIDE	in 5 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	400 mg in 5 mL
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	40 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	LEMON (lemon)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-5725-14	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/01/2011	

Labeler - MAJOR Pharmaceuticals Inc. (191427277)

Registrant - GCP Laboratories (965480861)

Establishment

Name	Address	ID/FEI	Business Operations
GCP Laboratories		965480861	manufacture(0904-5725)