

MINTOX MAXIMUM STRENGTH- aluminum hydroxide, magnesium hydroxide, dimethicone suspension
Proficient Rx LP

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 71205-515-35

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)



Scan Here



NDC 71205-515-35

Relabeled By: Proficient Rx LP
Thousand Oaks, CA 91320

Mintox Antacid
12 FL OZ (355 mL) Oral Suspension
Lot #:00000 SN# MASTER
NDC 71205-515-35 Exp:00/00/00

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GTIN: 00371205515355
SN# MASTER
Exp. 00/00/00
Lot #:00000



5

Mintox Antacid

12 FL OZ (355 mL) Oral Suspension

Each 5 mL teaspoonful contains: Aluminum hydroxide 400 mg (equivalent to dried gel USP) Antacid / Magnesium hydroxide 400 mg Antacid / Simethicone 40 mg Antigas

See bottle. Lemon Creme flavored

Product ID: SM051535

Dist. By: MAJOR® PHARMACEUTICALS 17177 N Laurel Park Drive, Suite 233 Livonia, MI 48152

Store at room temperature and avoid freezing

Keep medication out of the reach of children

MINTOX MAXIMUM STRENGTH

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|------------------------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:71205-515(NDC:0904-5725) |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---------------------|----------------|
| ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0) | ALUMINUM HYDROXIDE | 400 mg 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 |
|--|----------|
| 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: 059QF0K00R) | |
| SACCHARIN SODIUM (UNII: SB8ZUX40TY) | |
| SORBITOL (UNII: 506T60A25R) | |

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:71205-515-35 | 355 mL in 1 BOTTLE; Type 0: Not a Combination Product | 12/16/2020 | |

Marketing Information

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

Labeler - Proficient Rx LP (079196022)

Establishment

| Name | Address | ID/FEI | Business Operations |
|------------------|---------|-----------|---------------------|
| Proficient Rx LP | | 079196022 | RELABEL(71205-515) |

Revised: 11/2023

Proficient Rx LP