

MILK OF MAGNESIA ORIGINAL- magnesium hydroxide liquid
Preferred Pharmaceuticals Inc.

gc mom

Active ingredient (in each 15 mL tablespoonful)

Magnesium hydroxide 1200 mg

Purpose

Saline laxative

Uses

- relieves occasional constipation (irregularity)
- usually produces bowel movement in 1/2 to 6 hours

Warnings

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet
- stomach pain, nausea, or vomiting
- a sudden change in bowel habits that lasts over 14 days

Ask a doctor or pharmacist before use if you are

taking a prescription drug.

This product may interact with certain prescription drugs.

Stop use and ask a doctor if

- you have rectal bleeding or no bowel movement after using this product. These could be signs of a serious condition.
- you need to use a laxative for more than 1 week

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- shake well before use

- do not exceed the maximum recommended daily dose in a 24 hour period
- dose may be taken once a day preferably at bedtime, in divided doses, or as directed by a doctor
- drink a full glass (8 oz) of liquid with each dose

adults and children 12 years and older	2 to 4 tablespoonfuls
children 6 to 11 years	1 to 2 tablespoonfuls
children under 6 years	ask a doctor

Other information

- **each 15 mL tablespoonful contains:** magnesium 500 mg
- store at room temperature and avoid freezing
- close cap tightly after use

Inactive ingredients

purified water, sodium hypochlorite

Relabeled By: Preferred Pharmaceuticals Inc.

package Label

<p>Milk of Magnesia Brand Name Item</p> <p>Each 15ml (tablespoonful) contains: Magnesium hydroxide 1200mg...saline laxative</p> <p>Pkg Size: Exp Date: Lot#: Batch#: Ins: Mfg: Geri-Care Pharmaceutical Corp Prod#: Warning Ask a doctor before use if you have kidney disease, a magnesium-restricted diet, stomach pain, nausea, or vomiting; a sudden change in bowel habits that lasts over 14 days; or taking a prescription drug. This product may interact with certain prescription drugs. Keep this and all medication out of the reach of children. See bottle for additional drug facts. Store at room temperature and avoid freezing. Original flavor. Effective overnight relief.</p>	 <p>CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed</p>	<p>Milk of Magnesia Qty: Ins: Lot#: Bat#: Prod# (NDC):</p>	Log	
<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Directions English</p> <p style="writing-mode: vertical-rl; transform: rotate(180deg);">Use as directed by your doctor</p>		<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Instrucciones Espanol:</p> <p style="writing-mode: vertical-rl; transform: rotate(180deg);">Uso según lo dirigido por su doctor</p>	<p>Milk of Magnesia Qty: Ins: Lot#: Bat#: Prod# (NDC):</p>	Chart
<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Directions English</p> <p style="writing-mode: vertical-rl; transform: rotate(180deg);">Use as directed by your doctor</p>	<p>Milk of Magnesia Qty: Insurance NDC: Lot#: Bat#:</p>	<p>Milk of Magnesia Qty: Ins: Lot#: Bat#: Prod# (NDC):</p>	Billing	
<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Directions English</p> <p style="writing-mode: vertical-rl; transform: rotate(180deg);">Use as directed by your doctor</p>	<p>Milk of Magnesia Qty: Ins: Lot#: Bat#: Prod# (NDC):</p>	<p>Milk of Magnesia Qty: Ins: Lot#: Bat#: Prod# (NDC):</p>	Patient	

MILK OF MAGNESIA ORIGINAL			
magnesium hydroxide liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8413(NDC:57896-649)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	1200 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM HYPOCHLORITE (UNII: DY38VHM5OD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8413-3	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M007	04/06/2023	

Labeler - Preferred Pharmaceuticals Inc. (791119022)**Registrant** - Preferred Pharmaceuticals Inc. (791119022)**Establishment**

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8413)

Revised: 7/2024

Preferred Pharmaceuticals Inc.