

MILK OF MAGNESIA- magnesium hydroxide suspension
Natco Pharma USA LLC

Milk of Magnesia, USP Original Flavor DASH DOSE TM

Active Ingredient (in each 30 mL cup)

Magnesium hydroxide 2400 mg

Purpose

Saline laxative

Uses

As a Laxative

- relieves occasional constipation (irregularity)
- usually produces bowel movement in ½ to 6 hours

Warnings

Do not use if you have ever had an allergic reaction to this product or any of its ingredients

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 (1-800-222-1222).

Principal Display Panel

Delivers 30 mL

NDC 69339-153-01

Milk of Magnesia, USP

2,400mg Magnesium Hydroxide/30mL

Shake Well

For Institutional Use Only

3 69339 15301 7

Lot# 123456

Exp: MM/YYYY

DASH Pharmaceuticals LLC

Upper Saddle River, NJ 07458

LD153.01.R0721

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**DASH-
DOSE**

MILK OF MAGNESIA

magnesium hydroxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69339-153
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM HYDROXIDE	2400 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	WHITE (opaque)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69339-153-17	40 in 1 BOX, UNIT-DOSE	12/27/2021	
1	NDC:69339-153-01	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	M007	12/27/2021	

Labeler - Natco Pharma USA LLC (079590418)

Establishment

Name	Address	ID/FEI	Business Operations
GCP Laboratories		965480861	manufacture(69339-153)

Revised: 11/2023

Natco Pharma USA LLC