MAGNESIUM CITRATE- magnesium citrate liquid NuCare Pharmaceuticals, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredients

Magnesium citrate 1.745 g per fl oz

purpose

Saline laxative

Use

for relief of occasional constipation (irregularity). Generally produces bowel movement in 1/2 to 6 hours

Warnings

Warnings for use of product

Do not use if

you are on a low salt diet

Ask a doctor before use if you have

- kidney disease
- a magnesium or sodium-restricted diet
- abdominal pain, nausea, or vomiting
- noticed a sudden change in bowel habits that persists over a period of two weeks
- already used a laxative for a period longer than 1 week

Ask a doctor or pharmacist if

you are taking any other drug. Take this prouct 2 or more hours before or after other durgs. Laxatives may affect how other drugs work

Stop use and ask a doctor

if you have rectal bleeding or failure to have a bowel movement after use. These could be signs of a serious condition

If pregnant or breast-feeding

ask a help 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

- drink a full glass (8 ounces) of liquid with each dose
- may be taken as a single daily dose or in divided doses
- adults and children 12 years of age and over 6.5 to 10 fl oz in 24 hours
- children 6 to under 12 years of age 3 to 7 fl oz maximum 7 fl oz in 25 hours
- children 2 to 6 years of age 2 to 3 fl oz maximum 3 fl oz in 24 hours
- children under 2 years of age- ask a doctor

discard unused product within 24 hours of opening bottle

Other information

- •magnesium content 290 mg per 1 fl oz (30 mL)
- •sodium content 45 mg per 1 fl oz (30 mL)
- •store at temperaturs between 46° and 86° F (8° and 30° C)

inactive ingredients

citric acid, lemon oil, polyethylene glycol, purified water, sodium bicarbonate, sodium saccharin, sucrose

Adverse reaction

Distributed By McKesson

One Post Street, San Francisco, CA 94104

Money Back Guarantee

Principal display panel

NuCare Pharmaceuticals, Inc. Oistributed by: McKesson, San Francisco 94104 Packaged By: NuCare Pharmaceuticals, Inc. Orange, CA 92867 NDC: 68071-4060-1 by your physician. Use only as directed **Magnesium Citrate** 10oz Oral Soln. 68071408001*10*000000*000000 San Francisco, Magnesium Citrate 1.745g per fl = OZ See manufacturer's label for full list of ingredients. Product #: R0684010 Rev 01/01/19

WARNING: KEEP OUT OF REACH OF CHILDREN

Magnesium Citrate

Lot: 000000 NDC: 68071-4060-01

MFR NDC: 49346-696-49 Exp.: 00-00

Serial# 00000000001

Magnesium Citrate

Lot: 000000 NDC: 68071-4060-01 MFR NDC: 49346-696-49 Exp.: 00-00

Serial# 00000000001



GTIN 00368071406016 Serial# 00000000001

Exp. Date 00-00 LOT#: 000000

Call your doctor for medical advice about side effects. You may report side effects to FDA at

1-800-FDA-1088.

STORE AT CONTROLLED TEMPERATURE 59-86°F.

MAGNESIUM CITRATE

magnesium citrate liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68071-4060(NDC:49348-696)

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MAGNESIUM CITRATE (UNII: RHO2601T9V) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM CITRATE	1.745 g in 29.6 mL

Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
LEMON OIL (UNII: 19GRO824LL)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
WATER (UNII: 059QF0KO0R)		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)		
SACCHARIN (UNII: FST467XS7D)		
SUCROSE (UNII: C151H8M554)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:68071- 4060-1	300 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	08/18/2017		

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part334	03/20/2012				

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment					
Name	Address	ID/FEI	Business Operations		
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-4060)		

Revised: 2/2022 NuCare Pharmaceuticals,Inc.