

MAG PHOS- magnesia phosphorica liquid
Energique, Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Drug Facts:

ACTIVE INGREDIENT:

(in each drop): 100% of Magnesia Phosphorica 30C.

INDICATIONS:

May temporarily relieve nerve pain and cramping pain.**

**Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated

WARNINGS:

If pregnant or breastfeeding, 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.

Do not use if tamper evident seal is broken or missing.

Store in a cool, dry place.

KEEP OUT OF REACH OF CHILDREN:

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

DIRECTIONS:

Adults and children 5 to 10 drops orally, 1 time daily or as otherwise directed by a health care professional. If symptoms persist for more than 7 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

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INACTIVE INGREDIENTS:

Demineralized water, 20% Ethanol.

QUESTIONS:

Dist. by Energique, Inc.

201 Apple Blvd.

Woodbine, IA 51579 **800.869.8078**

PACKAGE LABEL DISPLAY:

ENERGIQUE

SINCE 1987

HOMEOPATHIC REMEDY

MAG PHOS

30C

1 fl. oz. (30 ml)

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LOT: XXXXXX

MAG PHOS			
magnesia phosphorica liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0353
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
MAGNESIUM PHOSPHATE, DIBASIC TRIHYDRATE (UNII: HF539G9L3Q) (MAGNESIUM CATION - UNII:T6V3LHY838)			MAGNESIUM PHOSPHATE, DIBASIC TRIHYDRATE	30 [hp_C] in 30 mL
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
ALCOHOL (UNII: 3K9958V90M)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44911-0353-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	04/20/2016	10/26/2026
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved homeopathic			04/20/2016	10/26/2026

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0353) , api manufacture(44911-0353) , label(44911-0353) , pack(44911-0353)

Revised: 6/2022

Energique, Inc.