MAGNESIA PHOSPHORICA- magnesia phosphorica spray Ratis, LLC

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) Magnesia Phosphorica 12X 100%.

PURPOSE:

Magnesia Phosphorica - Antispasmodic pain*

USES:

May temporarily relieve pain, muscle cramps, menstrual cramps, headache, earache, toothache, and nerve injuries.*

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

WARNINGS:

For oral use only.

If pregnant or breast-feeding, ask a health care 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

KEEP OUT OF REACH OF CHILDREN:

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

DIRECTIONS:

- Adults: 2 sprays 3 times a day in mouth or half a cup of water
- Children under 12: 2-12: 1 spray as above
- Consult a physician for use in children under 12 years of age.

Store in a cool, dry place.

INACTIVE INGREDIENTS:

Demineralized water, 20% organic ethanol

QUESTIONS:

Questions?

www.HomeopathyStore.com

(888) 405-7551

Dist. by Ratis, LLC

1201 N Orange St,

Ste 7594

Wilmington, DE 19801

PACKAGE LABEL DISPLAY:

SCHUESSLER CELL SALTS

LACTOSE FREE

ANNA

KARE

Magnesia Phosphorica

HOMEOPATHIC ORAL SPRAY

1 FL. OZ (30ML)



leadener. No text

Drug Facts (continued)

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

Directions

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- Children under 12: 1 spray as above
- Consult a physician for use in children under 12 years of age.

Other information

Store in a cool, dry place.

Drug Facts (continued)

Inactive ingredients

Demineralized water, 20% organic ethanol

Questions?

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MAGNESIA PHOSPHORICA

magnesia phosphorica spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71753-8008

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength
MAGNESIUM PHOSPHATE, DIBASIC TRIHYDRATE (UNII: HF539G9L3Q) MAGNESIUM PHOSPHATE,

12 [hp_X]

(MAGNESIUM CATION - UNII:T6V3LHY838)

MAGNESIUM CATION - UNII:T6V3LHY838)

MAGNESIUM PHOSPHATE, DIBASIC TRIHYDRATE 12 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71753- 8008-1	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/30/2020	

	Marketing Information		
Marketing End Date	Marketing Start Date	Application Number or Monograph Citation	Marketing Category
	09/30/2020		unapproved homeopathic
	09/30/2020		

Labeler - Ratis, LLC (964594324)

Revised: 5/2023 Ratis, LLC