

MAGNESIA PHOSPHORICA - magnesia phosphorica pellet
HOMEOLAB USA 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.

HOMEOPATHIC MEDICINE NDC 60512-9152-1

ACTIVE INGREDIENT HPUS

MAGNESIA PHOSPHORICA 1X

(Magnesium hydrogen phosphate)

CRAMPS

USE

For self-limiting condition listed above or as directed by a health professional.

WARNINGS

Enter section text here

Do not use if pellet-dispenser seal is broken.

Stop use and ask a doctor if symptoms persist more than 3 days or worsen.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

DIRECTIONS

(Adults / Children 2-18 years): Allow 3 or 4 pellets to dissolve in the mouth 3 times a day until symptoms are relieved or as directed by a health professional.

OTHER INFORMATION

Store at room temperature.

INACTIVE INGREDIENTS

Lactose, sucrose.

QUESTIONS?

1-800-404-4666

The letters HPUS indicate the ingredient is officially included in the Homeopathic Pharmacopoeia of the United States.

80 Pellets

Pellet dispenser

HOMEOLAB USA INC., 3025 De L'Assomption, Montreal, QC, H1N 2H2, CANADA

Product of Canada

LABEL

HOMEOPATHIC MEDICINE

MAGNESIA PHOSPHORICA 1x

Magnesium Hydrogen Phosphate

NDC 60512-9152-1

CRAMPS

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Break seal, turn & twist.

A-C

80 Pellets
Pellet dispenser



The Finest in Homeopathy
HOMEOLAB
USA

Product
of Canada

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MAGNESIA PHOSPHORICA

magnesia phosphorica pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60512-9152
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	1 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	
SUCROSE (UNII: C151H8M554)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60512-9152-1	80 in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		11/30/2011	

Labeler - HOMEOLAB USA INC (202032533)**Establishment**

Name	Address	ID/FEI	Business Operations
HOMEOLAB USA INC		202032533	manufacture

Revised: 11/2011

HOMEOLAB USA INC