MAGNESIA SULPHURICA- magnesium sulfate heptahydrate pellet Boiron

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.

Magnesia sulphurica 30C

Magnesia sulphurica 30C

(**contains 0.443 mg of the active ingredient per pellet)

Irritability or itchy skin rash*

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

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

Keep out of reach of children

Do not use if pellet dispenser seal is broken.

Contains approx 80 pellets.

How to dispense pellets? Turn tube upside down. Twist until 5 pellets are dispensed into cap. Carefully remove the cap and use it to pour pellets under the tongue. *CLAIMS BASED ON TRADITIONAL HOMEOPATHIC PRACTICE NOT ACCEPTED MEDICAL EVIDENCE. NOT FDA EVALUATED.

*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

lactose, sucrose

Adults and children: At the onset of symptoms, dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

1-800-BOIRON-1 (1-800-264-7661), BoironUSA.com Info@boiron.com Distributed by Boiron, Inc. Newtown Square, PA 19073





Drug Facts

Active ingredient^{**}: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

Uses: See symptoms on front panel.

Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before use. Keep out of reach of children.

Directions: Adults and children: At the onset of symptoms, dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

Other information: Do not use if pellet dispenser seal is broken.

Drug Facts (continued) Inactive ingredients: lactose, sucrose



MAGNESIA SULPHU	JRICA					
magnesium sulfate heptahy	drate pellet					
Product Information						
Product Type	HUMAN OTC DRUG	Ite	m Code	e (Source)	NDC:	0220-3267
Route of Administration	ORAL					
Active Ingredient/Active	e Moiety					
Ingre	dient Name			Basis of Streng	Jth	Strength
MAGNESIUM SULFATE HEPTAHYDRATE (UNII: SK47B8698T)MAGNESIUM SULFATE(MAGNESIUM CATION - UNII:T6V3LHY838)HEPTAHYDRATE					30 [hp_C] in 30 [hp_C]	
Institut Insurationta						
Inactive Ingredients						
Ingredient Name						Strength
SUCROSE (UNII: C151H8M554)						
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)						
Product Characteristics	5					
Color	white	Score				
Shape	ROUND	Size				4mm
Flavor		Imprint	Code			

Co	ntains			
Pa	ackaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:0220-3267- 41	30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	
Μ	arketing	Information		
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	approved meopathic		03/03/1983	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment								
Name	Address	ID/FEI	Business Operations					
Boiron		282560473	manufacture(0220-3267)					

Revised: 11/2023

Boiron