MAGNESIA CARBONICA- magnesium carbonate 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 carbonica 6C

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

Diarrhea or facial pain*

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).

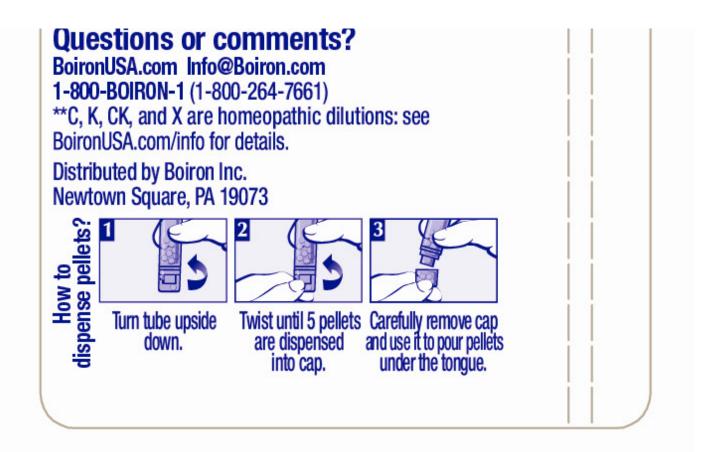
Uses: See symptoms on front panel.

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 CARBON	NICA						
magnesium carbonate pelle	t						
Product Information							
Product Type	HUMAN OTC DRUG		Item Code (Source)		ND	NDC:0220-3250	
Route of Administration	ORAL						
Active Ingredient/Active	e Moiety						
Ingredient Name Basis of						Strength	
ingre				Strength		Strength	
MAGNESIUM CARBONATE (UNII: 0E53J927NA) (CARBONATE ION -MAGNESIUMUNII:7UJQ50PE7D)CARBONATE				6 [hp_C] in 6 [hp_C]			
Inactive Ingredients							
y	Ingredient Na	me				Strength	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)						j	
SUCROSE (UNII: C151H8M554)							
Product Characteristics	5						
Color	white	Score	•				
Shape	ROUND	Size				4mm	

Flavor		Imprint C	Imprint Code			
C	ontains					
Ρ	ackaging					
#	ltem Code	P	ackage Description	Marketing Start Date	Marketing End Date	
1	NDC:0220-3250- 41	6 [hp_C] in 1 Product	TUBE; Type 0: Not a Combination	03/03/1983		
M	larketing	Informa	tion			
	Marketing Category	Applic	ation Number or Monograp Citation	h Marketing Start Date	Marketing End Date	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment								
Name	Address	ID/FEI	Business Operations					
Boiron		282560473	manufacture(0220-3250)					

Revised: 11/2023

Boiron