CORALLIUM RUBRUM- corallium rubrum exoskeleton 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.

Corallium rubrum 5C

Corallium rubrum 5C

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

Fitful cough worsened by cold*

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).

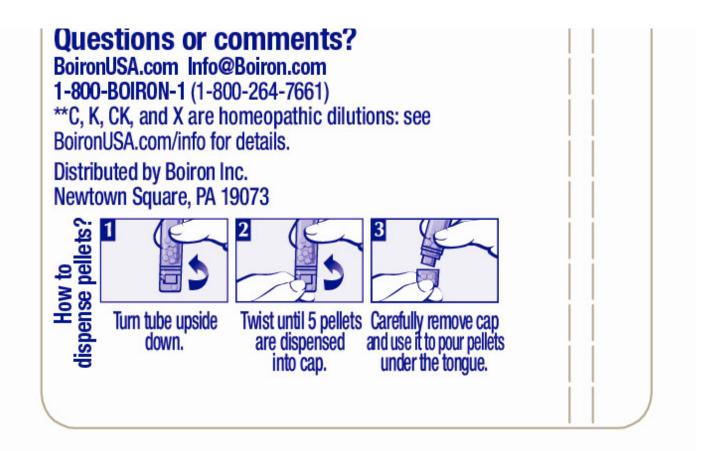
Uses: See symptoms on front panel.

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



CORALLIUM RUBRU	M								
corallium rubrum exoskeleton pellet									
Product Information									
Product Type	HUMAN OTC DRUG		Item Code (So	ource)	NDC:02	20-1541			
Route of Administration	ORAL								
Active Ingredient/Active	e Moiety								
Ingr	ength	Strength							
CORALLIUM RUBRUM EXOSKELETON (UNII: 2CA71K0DLE) (CORALLIUMCORALLIUM RUBRRUBRUM EXOSKELETON - UNII:2CA71K0DLE)EXOSKELETON					RUM	5 [hp_C] in 5 [hp_C]			
Inactive Ingredients									
Ingredient Name						Strength			
SUCROSE (UNII: C151H8M554)						5			
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)									
Product Characteristics									
Color	white	Score	2						
Shape	ROUND	Size			4	mm			
Flavor		Impri	nt Code						

	ntains			
Pa	ickaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:0220-1541- 41	5 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	
Μ	arketing	Information		
Μ	arketing Marketing Category	I nformation Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
una	Marketing	Application Number or Monograph		Marketing End Date

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment								
Name	Address	ID/FEI	Business Operations					
Boiron		282560473	manufacture(0220-1541)					

Revised: 11/2023

Boiron