GRATIOLA OFFICINALIS- gratiola officinalis 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.

Gratiola officinalis 30C

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

Diarrhea triggered by cold drinks*

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

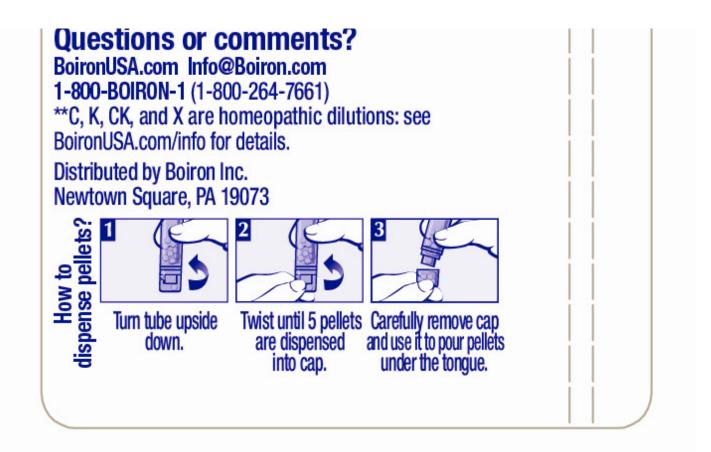




Drug Facts Active ingredient**: See product name on front panel (contains 0.443 mg of the active ingredient per pellet). Uses: See symptoms on front panel. Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before 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



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GRATIOLA OFFICIN	ALIS					
gratiola officinalis pellet						
Product Information						
Product Type	HUMAN OTC DRUG Item Code (Source)		NDC:0220-2329			
Route of Administration	ORAL					
Active Ingredient/Active	e Moiety					
Ingr	Strength					
GRATIOLA OFFICINALIS (UNII: BDD9991A36) (GRATIOLA OFFICINALIS - GRATIOLA OFFICINALIS - OFFICINALIS				30 [hp_C] in 30 [hp_C]		
Inactive Ingredients						
	Strength					
LACTOSE, UNSPECIFIED FORM						
SUCROSE (UNII: C151H8M554)						
Product Characteristics						
Color	white	Scor	e			
Shape	ROUND	Size			4mm	

tains kaging							
tem Code	Package De	Package Description		Marketing End Date			
rkoting	nformation						
Marketing Category			Marketing Start Date	Marketing End Date			
proved eopathic			03/03/1983				
	rketing Marketing Category	Product rketing Information Marketing Category Droved	Product rketing Information Marketing Category Application Number or Monograph Citation proved Citation	Product 03/03/1983 rketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date oroved 03/03/1983 03/03/1983			

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-2329)			

Revised: 11/2023

Boiron