ARNICA MONTANA- arnica montana 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.

Arnica montana 3X

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

Muscle Pain, Stiffness, Swelling From Injuries, Bruises*

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).

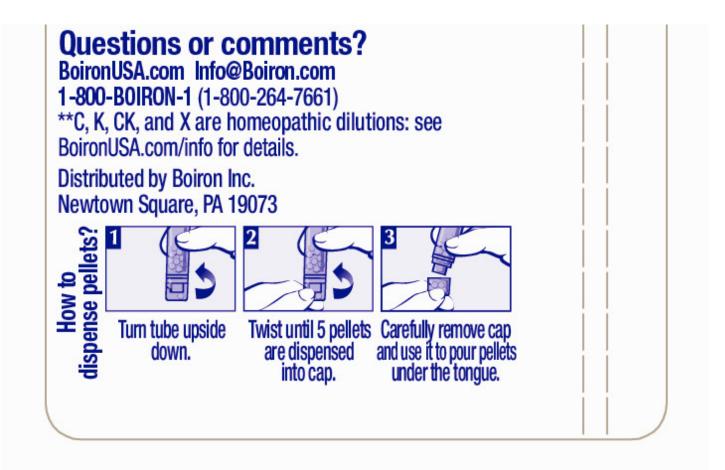
Uses: See symptoms on front panel.

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



ARNICA MONTANA						
arnica montana pellet						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code	e (Source)	NDC:0220-0515		
Route of Administration	ORAL					
Active Ingredient/Active	e Moiety					
Ingre	Strength					
ARNICA MONTANA (UNII: 080TY) UNII:080TY208ZW)	3 [hp_X] in 3 [hp_X]					
Inactive Ingredients						
	Strength					
LACTOSE, UNSPECIFIED FORM						
SUCROSE (UNII: C151H8M554)						
Product Characteristics						
Color	white	Score				
Shape	ROUND	Size		4mm		

Flavor		Imprint Code					
ontains							
Packaging							
ltem Code	P	Package Description		Marketing Start Date	Marketing End Date		
NDC:0220-0515- 41	3 [hp_X] in 1 Product)3/03/1983			
larkoting	Informa	tion					
Marketing Category	Applic	ation Number or Mo Citation	nograph	Marketing Start Date	Marketing End Date		
approved				03/03/1983			
1	ackaging Item Code NDC:0220-0515- 41 Iarketing Category	ackaging item Code NDC:0220-0515- 41 Product Product Product Product Product Product Product Product	ackaging Item Code Package Description NDC:0220-0515- 3 [hp_X] in 1 TUBE; Type 0: Not a Com Product Iarketing Information Marketing Application Number or Mo Citation	ackaging Item Code Package Description NDC:0220-0515- 3 [hp_X] in 1 TUBE; Type 0: Not a Combination 41 Product Iarketing Category Application Number or Monograph Citation	Item Code Package Description Marketing Start Date NDC:0220-0515- 3 [hp_X] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983 Item Keting Start Date Marketing Category Application Number or Monograph Citation Marketing Start Date		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-0515)			

Revised: 2/2023

Boiron