SYMPHYTUM OFFICINALE- comfrey root 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.

Symphytum officinale 30X

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

Symptoms of bone trauma*

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).

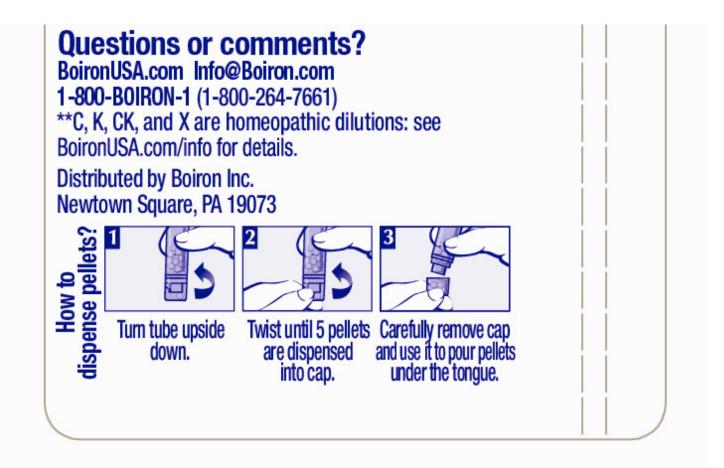
Uses: See symptoms on front panel.

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



SYMPHYTUM OFFIC	CINALE					
comfrey root pellet						
Product Information						
Product Type	HUMAN OTC DRUG		Item Cod	le (Source)	NDC:0220-4897	
Route of Administration	ORAL					
	NA - ¹ - 1					
Active Ingredient/Active	emolety			Basis of		
Ingred	Strength					
COMFREY ROOT (UNII: M9VVZ08EKQ) (COMFREY ROOT - UNII: M9VVZ08EKQ)			COMFREY ROOT	30 [hp_X] in 30 [hp_X]		
Inactive Ingredients						
	Strength					
SUCROSE (UNII: C151H8M554)						
LACTOSE, UNSPECIFIED FORM	I (UNII: J2B2A4N98G)					
Product Characteristics						
Color	white	Score	e			
Shape	ROUND	Size			4mm	

ntains				e			
	Contains						
- I							
Packaging							
ltem Code	P	Package Description		Marketing Start Date	Marketing End Date		
NDC:0220-4897- 41	30 [hp_X] in 2 Product			03/03/1983			
arkotina	Informa	tion					
Marketing Category	Applic	ation Number or Mon Citation	nograph	Marketing Start Date	Marketing End Date		
ipproved neopathic				03/03/1983			
	Item Code IDC:0220-4897- 11 Arketing Category pproved	Item Code ADC:0220-4897- ATREETING Informa Marketing Category pproved	Item Code Package Description ADC:0220-4897- 11 30 [hp_X] in 1 TUBE; Type 0: Not a Composition Arketing Information Marketing Category Application Number or Mon Citation	Item Code Package Description ADC:0220-4897- 11 30 [hp_X] in 1 TUBE; Type 0: Not a Combination Product Arketing Category Application Number or Monograph Citation	Item Code Package Description Marketing Start Date ADC:0220-4897- 11 30 [hp_X] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983 arketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date pproved 03/03/1983 03/03/1983 03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-4897)			

Revised: 7/2023

Boiron