## BELLIS PERENNIS- bellis perennis 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.

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## **Bellis perennis 30C**

Bellis perennis 30C

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

Post-traumatic breast 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**<sup>\*\*</sup>: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

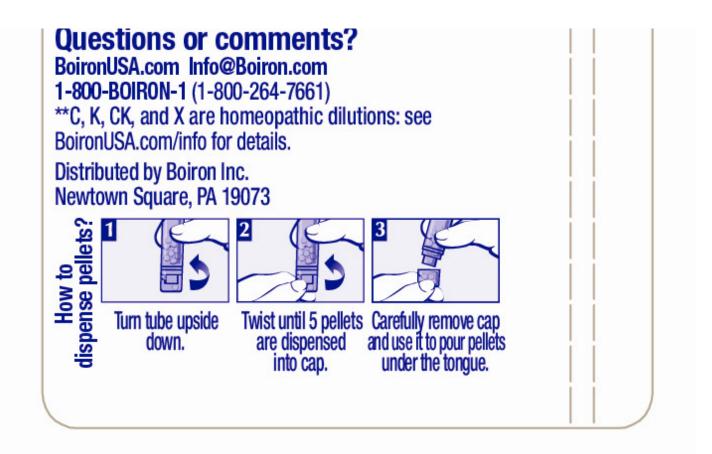
**Uses:** See symptoms on front panel.

**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



BELLIS PERENNIS				
bellis perennis pellet				
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Product Information				
Product Type	HUMAN OTC DRUG	Item Co	de (Source)	NDC:0220-0732
Route of Administration	ORAL			
<b>Active Ingredient/Active</b>	e Moiety			
Ingred	Strength			
BELLIS PERENNIS (UNII: 2HU33I03UY) (BELLIS PERENNIS - UNII:2HU33I03UY)			BELLIS PERENNIS	30 [hp_C] in 30 [hp_C]
Inactive Ingredients				
	Strength			
LACTOSE, UNSPECIFIED FORM				
SUCROSE (UNII: C151H8M554)				
<b>Product Characteristics</b>	•			
Color	white	Score		
Shape	ROUND	Size		4mm

			Imprint Code			
ntains						
Packaging						
ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date	
NDC:0220-0732- 41	30 [hp_C] in Product	1 TUBE; Type 0: Not a Con	nbination	03/03/1983		
arkating	Informa	tion				
Marketing information						
Marketing Category	Applic	ation Number or Mon Citation	ograph	Marketing Start Date	Marketing End Date	
approved meopathic				03/03/1983		
	Item Code NDC:0220-0732- 41 arketing Marketing Category approved	Item Code Para Strain Para Strain Para Strain Para Strain Product Prod	Item Code     Package Description       NDC:0220-0732- 41     30 [hp_C] in 1 TUBE; Type 0: Not a Comproduct       arketing Information       Marketing Category     Application Number or Monicitation       Approved	Item Code       Package Description         NDC:0220-0732- 41       30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product         arketing Category       Application Number or Monograph Citation         Apploation Number or Monograph Citation	Item Code       Package Description       Marketing Start Date         NDC:0220-0732- 41       30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product       03/03/1983         arketing Information         Marketing Category       Application Number or Monograph Citation       Marketing Start Date         opproved       03/03/1983	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment					
Name	Address	ID/FEI	Business Operations		
Boiron		282560473	manufacture(0220-0732)		

Revised: 11/2023

Boiron