

BELLIS PERENNIS- bellis perennis pellet
OHM PHARMA INC.

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.

Bellis Perennis

DRUG FACTS: ACTIVE INGREDIENTS: BELLIS PERENNIS

CONTENT: Approx. 100 Pellets

USE: To be used according to standard homeopathic indications.

WARNINGS: Keep out of reach of children. If pregnant or breast-feeding, ask a health care professional before use. **Stop use and ask a health care professional** if symptoms persist for more than 3 days or worsen.

Keep out of reach of children.

DIRECTIONS: Dissolve 3-5 pellets under the tongue 3 times a day or as directed by a professional.

OTHER INFORMATION: Store at room temperature. **Do not use** if pellet dispenser seal is broken.

INACTIVE INGREDIENTS:Organic sucrose, lactose free.

Manufactured according to the Homeopathic Pharmacopoeia of the United States (HPUS). **PRODUCT OF USA.**

Mfg. By: OHM PHARMA, INC. Mineral Wells, TX 76067

www.ohmpharma.com FDA Est # 3003231743

BELLIS PERENNIS

The OTC Potency range from Bellis Perennis 6X-30X, 3C-30C, 200C, 1M.

Standard bottle sizes for dilution-form can range from 30mL to 60mL.



To be used according to standard homeopathic indications.

BELLIS PERENNIS

bellis perennis pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66096-798
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BELLIS PERENNIS (UNII: 2HU33I03UY) (BELLIS PERENNIS - UNII:2HU33I03UY)	BELLIS PERENNIS	6 [hp_C] in 6 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	white	Score	
Shape	ROUND	Size	4mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66096-798-01	6 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	11/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		11/01/2019	

Labeler - OHM PHARMA INC. (030572478)

Revised: 12/2021

OHM PHARMA INC.