

PRIMULA ONOPORDON- primula onopordon liquid
Uriel Pharmacy 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.

Primula Onopordon

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredient: Onopordon (Cotton thistle) 3X, Primula (Cowslip) 3X, Hyoscyamus (Henbane) 4X

Inactive Ingredients: Water, Salt

Use: Temporary relief of stress.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use.

Questions? Call 866.642.2858 Made by Uriel, East Troy, WI 53120 shopuriel.com

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Lot:



**Primula
Onopordon**

Homeopathic Ampules
net vol. 0.3 fl. oz (10 x 1 ml)

Primula Onopordon

PRIMULA ONOPORDON

primula onopordon liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8398
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ONOPORDUM ACANTHIUM FLOWER (UNII: AP97AUF88E) (ONOPORDUM ACANTHIUM FLOWER - UNII:AP97AUF88E)			ONOPORDUM ACANTHIUM FLOWER	3 [hp_X] in 1 mL
PRIMULA VERIS FLOWER (UNII: W5BET37294) (PRIMULA VERIS FLOWER - UNII:W5BET37294)			PRIMULA VERIS FLOWER	3 [hp_X] in 1 mL
HYOSCYAMUS NIGER LEAF (UNII: 32IT7G8BAW) (HYOSCYAMUS NIGER LEAF - UNII:32IT7G8BAW)			HYOSCYAMUS NIGER LEAF	4 [hp_X] in 1 mL
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8398-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved homeopathic			09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-8398)

Revised: 10/2023

Uriel Pharmacy Inc.