

ONOPORDON COMP.- onopordon comp. pellet
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.

Onopordon comp.

Directions: FOR ORAL USE ONLY.

Dissolve pellets under the tongue 3-4 times daily. Ages 12 and older: 10 pellets. Ages 2-11: 5 pellets. Under age 2: Consult a doctor.

Active Ingredients: 100gm contains: 25gm Onopordon (Cotton thistle) 1X, 25gm Primula (Cowslip) 1X, 25gm Hyoscyamus (Henbane) 4X

Inactive Ingredient: Organic sucrose

Use: Promotes healthy circulatory support.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Contains sugar. Diabetics and persons intolerant of sucrose (sugar): Consult a doctor before use. 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. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858

Uriel, East Troy WI 53120

www.urielpharmacy.com



ONOPORDON COMP.

onopordon comp. pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-7090
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	1 [hp_X]
PRIMULA VERIS FLOWER (UNII: W5BET37294) (PRIMULA VERIS FLOWER - UNII:W5BET37294)	PRIMULA VERIS FLOWER	1 [hp_X]

HYOSCYAMUS NIGER LEAF (UNII: 32IT7G8BAW) (HYOSCYAMUS NIGER LEAF - UNII:32IT7G8BAW)			HYOSCYAMUS NIGER LEAF	4 [hp_X]
Inactive Ingredients				
Ingredient Name			Strength	
SUCROSE (UNII: C151H8M554)				
Product Characteristics				
Color	white	Score	no score	
Shape	ROUND	Size	3mm	
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-7090-2	1350 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	09/01/2009	
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(48951-7090)

Revised: 6/2018

Uriel Pharmacy Inc.