ARNICA BERBERIS TEUCRIUM- arnica berberis teucrium 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.

Arnica Berberis Teucrium

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: Berberis (Barberry) 2X, Teucrium scor. (Wood sage) 2X, Arnica 4X

Inactive Ingredient: Organic sucrose

Use: Temporary relief of headache.

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

ARNICA BERBERIS TEUCRIUM

arnica berberis teucrium pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-1113
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	2 [hp_X]
TEUCRIUM SCORODONIA WHOLE (UNII: 2MJ724W91Q) (TEUCRIUM SCORODONIA WHOLE - UNII:2MJ724W91Q)	TEUCRIUM SCORODONIA WHOLE	2 [hp_X]
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	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:48951-1113-2	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(48951-1113)

Revised: 5/2018 Uriel Pharmacy Inc.