ARNICA SCORODITE- arnica scorodite 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 Scorodite

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: Amethyst (Nat. silicic acid anhydrate with traces of manganese and iron) 6X, Arnica 6X, Conchae (Oyster shells) 6X, Levisticum (Lovage) 6X, Natrium carb. (Sodium carbonate monohydrate) 6X, Orchis tub. decoct. (Orchid) 6X, Apatite (Nat. calcium fluorophosphate) 7X, Scorodite (Nat. ferric arsenate) 7X, Cerebellum (Bovine cerebellum) 8X

Inactive Ingredients: Sucrose, Lactose

Use: Temporary relief of headache.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Contains sugar. Diabetics and persons intolerant of sucrose (sugar): Consult a doctor before use. Do not use if allergic to any ingredient. Contains traces of lactose. 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. Natural ingredients may cause color, scent and/or taste variation.

Questions? Call 866.642.2858
Uriel, East Troy, WI 53120
www.urielpharmacy.com

ARNICA SCORODITE

arnica scorodite pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-1156
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	6 [hp_X]
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	6 [hp_X]
OSTREA EDULIS SHELL (UNII: 49OY13BE7Z) (OSTREA EDULIS SHELL - UNII:49OY13BE7Z)	OSTREA EDULIS SHELL	6 [hp_X]
LEVISTICUM OFFICINALE ROOT (UNII: 46QZ19OEX8) (LEVISTICUM OFFICINALE ROOT - UNII:46QZ19OEX8)	LEVISTICUM OFFICINALE ROOT	6 [hp_X]
SODIUM CARBONATE (UNII: 45P3261C7T) (CARBONATE ION - UNII:7UJQ5OPE7D)	SODIUM CARBONATE	6 [hp_X]
ORCHIS MASCULA TUBER (UNII: 5H2N55J61B) (ORCHIS MASCULA TUBER - UNII:5H2N55J61B)	ORCHIS MASCULA TUBER	6 [hp_X]
FLUORAPATITE (UNII: M4CM1H238J) (FLUORAPATITE - UNII:M4CM1H238J)	FLUORAPATITE	7 [hp_X]
FERROUS ARSENATE (UNII: 129CO35H12) (FERROUS ARSENATE - UNII:129CO35H12)	FERROUS ARSENATE	7 [hp_X]
BOS TAURUS CEREBELLUM (UNII: Q09851U44F) (BOS TAURUS CEREBELLUM - UNII:Q09851U44F)	BOS TAURUS CEREBELLUM	8 [hp_X]

Inactive Ingredients	
Ingredient Name	**Strength**
SUCROSE (UNII: C151H8M554)	
LACTOSE (UNII: J2B2A4N98G)	

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:48951-1156-2	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(48951-1156)