

CINNABAR DANDELION- cinnabar dandelion 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.

Cinnabar Dandelion

Directions: FOR ORAL USE.

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

Active Ingredients: Agropyron (Couch grass) 3X, Taraxacum (Dandelion) 3X, Pyrite (Nat. Iron disulfide) 8X, Kali carb e cinere Fagi (Potassium carbonate from beech wood ash) 10X, Cinnabaris (Nat. mercuric sulfide) 20X

Inactive Ingredients: Water, Salt, Lactose

Uses: Temporary relief of cold, flu and sore throat symptoms.

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. 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.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 shopuriel.com Lot:

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Lot:

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

CINNABAR DANDELION			
cinnabar dandelion liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-3101
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ELYMUS REPENS ROOT (UNII: 3IXW0F6P8W) (ELYMUS REPENS ROOT - UNII:3IXW0F6P8W)	ELYMUS REPENS ROOT	3 [hp_X] in 1 mL
TARAXACUM PALUSTRE ROOT (UNII: GCZ4W7077C) (TARAXACUM PALUSTRE ROOT - UNII:GCZ4W7077C)	TARAXACUM PALUSTRE ROOT	3 [hp_X] in 1 mL
FERROUS DISULFIDE (UNII: 132N09W4PR) (FERROUS CATION - UNII:GW89581OWR)	FERROUS DISULFIDE	8 [hp_X] in 1 mL
CARBONATE ION (UNII: 7UJQ5OPE7D) (CARBONATE ION - UNII:7UJQ5OPE7D)	CARBONATE ION	10 [hp_X] in 1 mL
MERCURIC CATION (UNII: ED30FJ8Y42) (MERCURIC CATION - UNII:ED30FJ8Y42)	MERCURIC CATION	20 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LACTOSE (UNII: J2B2A4N98G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-3101-1	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(48951-3101)

Revised: 3/2024

Uriel Pharmacy Inc.