

SPIGELIA COMP- spigelia comp 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.

Spigelia comp

Directions: FOR ORAL USE.

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

Active Ingredients: Aconitum (Monkshood) 8X, Arnica 8X, Natrium nitrosum (Sodium nitrite) 8X, Spigelia (Pinkroots) 8X, Argentum nitricum (Silver nitrate) 10X, Phosphorus (Yellow phosphorus) 12X

Inactive Ingredients: Water, Salt

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. 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

www.urielpharmacy.com Lot:



SPIGELIA COMP

spigelia comp liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8368
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACONITUM NAPELLUS (UNII: U0NQ8555JD) (ACONITUM NAPELLUS - UNII:U0NQ8555JD)	ACONITUM NAPELLUS	8 [hp_X] in 1 mL
SODIUM NITRITE (UNII: M0KG633D4F) (NITRITE ION - UNII:J39976L608)	NITRITE ION	8 [hp_X] in 1 mL
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ) (ARNICA MONTANA FLOWER - UNII:OZ0E5Y15PZ)	ARNICA MONTANA FLOWER	8 [hp_X] in 1 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	12 [hp_X] in 1 mL
SPIGELIA MARILANDICA ROOT (UNII: 467D26HS0B) (SPIGELIA MARILANDICA ROOT	SPIGELIA MARILANDICA	8 [hp_X]

- UNII:467D26HS0B)	ROOT	in 1 mL
SILVER NITRATE (UNII: 95IT3W8JZE) (SILVER CATION - UNII:57N7B0K90A)	SILVER NITRATE	10 [hp_X] in 1 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8368-1	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(48951-8368)