

HELLEBORUS COMP. B- helleborus comp. b 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.

Helleborus comp. B

Directions: FOR ORAL USE.

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

Active Ingredients: Colchicum (Meadow saffron) 4X, Helleborus (Christmas rose) 4X, Lien (Bovine spleen) 8X, Argentum (Silver) 12X, Stibium (Antimony) 17X, Equisetum (Common horsetail) 20X, Formica (Red wood ant) 20X, Phosphorus (Yellow phosphorus) 30X, Plumbum (Lead) 30X

Inactive Ingredients: Water, Salt

Prepared using rhythmical processes.

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

Uriel, East Troy, WI 53120

shopuriel.com Lot:

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Inactive Ingredients: Water, Salt

Prepared using rhythmical processes.

Use: Temporary relief of headache.

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Uriel

Helleborus comp. B

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

Helleborus comp. B

HELLEBORUS COMP. B

helleborus comp. b liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-5115	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
COLCHICUM AUTUMNALE WHOLE (UNII: W79255C628) (COLCHICUM AUTUMNALE WHOLE - UNII:W79255C628)		COLCHICUM AUTUMNALE WHOLE	4 [hp_X] in 1 mL	
FORMICA RUFA (UNII: 55H0W83JO5) (FORMICA RUFA - UNII:55H0W83JO5)		FORMICA RUFA	20 [hp_X] in 1 mL	
BOS TAURUS SPLEEN (UNII: 190Q9MB3U3) (BOS TAURUS SPLEEN - UNII:190Q9MB3U3)		BOS TAURUS SPLEEN	8 [hp_X] in 1 mL	
ANTIMONY (UNII: 9IT35J3UV3) (ANTIMONY - UNII:9IT35J3UV3)		ANTIMONY	17 [hp_X] in 1 mL	
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)		SILVER	12 [hp_X] in 1 mL	
HELLEBORUS FOETIDUS ROOT (UNII: 332Y7EES96) (HELLEBORUS FOETIDUS ROOT - UNII:332Y7EES96)		HELLEBORUS FOETIDUS ROOT	4 [hp_X] in 1 mL	
EQUISETUM ARVENSE TOP (UNII: 1DP6Y6B65Z) (EQUISETUM ARVENSE TOP - UNII:1DP6Y6B65Z)		EQUISETUM ARVENSE TOP	20 [hp_X] in 1 mL	
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)		PHOSPHORUS	30 [hp_X] in 1 mL	
LEAD (UNII: 2P299V784P) (LEAD - UNII:2P299V784P)		LEAD	30 [hp_X] in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-5115-1	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(48951-5115)

Revised: 12/2024

Uriel Pharmacy Inc.