

HELLEBORUS COMP. F- helleborus comp. f 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. F

Directions: FOR ORAL USE.

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

Active Ingredients: Bulbus olfactorius (Bovine olfactory bulb) 8X, Arnica 12X, Helleborus (Christmas rose) 12X, Jaspis (Jasper) 12X, Tunica mucosa nasi (Bovine nasal mucosa) 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

Uriel, East Troy, WI 53120

shopuriel.com Lot:

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Lot:



**Helleborus
comp. F**

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

Helleborus comp. F

HELLEBORUS COMP. F

helleborus comp. f liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-5130	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)		SILICON DIOXIDE	12 [hp_X] in 1 mL	
BOS TAURUS OLFACTORY BULB (UNII: S46V5TR3J8) (BOS TAURUS OLFACTORY BULB - UNII:S46V5TR3J8)		BOS TAURUS OLFACTORY BULB	8 [hp_X] in 1 mL	
HELLEBORUS NIGER ROOT (UNII: 608DGJ6815) (HELLEBORUS NIGER ROOT - UNII:608DGJ6815)		HELLEBORUS NIGER ROOT	12 [hp_X] in 1 mL	
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)		ARNICA MONTANA	12 [hp_X] in 1 mL	
BOS TAURUS NASAL MUCOSA (UNII: 343455G79K) (BOS TAURUS NASAL MUCOSA - UNII:343455G79K)		BOS TAURUS NASAL MUCOSA	12 [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:48951-5130-1	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(48951-5130)