

HELLEBORUS E PL. TOTA 2- helleborus e pl. tota 2 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 e pl. tota 2

Directions: FOR ORAL USE.

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

Active Ingredient: Helleborus (Christmas rose) 2X

Inactive Ingredients: Water, Salt

Prepared using rhythmical processes.

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 shopuriel.com Lot:

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**Helleborus
e pl. tota 2X**

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

Helleborus e pl. tota 2X

HELLEBORUS E PL. TOTA 2

helleborus e pl. tota 2 liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-5037
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
HELLEBORUS FOETIDUS ROOT (UNII: 332Y7EES96) (HELLEBORUS FOETIDUS ROOT - UNII:332Y7EES96)			HELLEBORUS FOETIDUS ROOT	2 [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:48951-5037-1	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(48951-5037)

Revised: 12/2024

Uriel Pharmacy Inc.