

GENTIANA ABSINTHIUM- gentiana absinthium 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.

Gentiana Absinthium

Directions: FOR ORAL USE.

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

Active Ingredients: Absinthium (Wormwood) 3X, Gentiana (Yellow gentian) 3X, Taraxacum (Dandelion) 3X, Nux vomica (Poison nut) 6X

Inactive Ingredients: Water, Salt

Use: Temporary relief of digestive upset.

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

www.urielpharmacy.com

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**Gentiana
Absinthium**

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

Gentiana Absinthium

GENTIANA ABSINTHIUM

gentiana absinthium liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-5007
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WORMWOOD (UNII: E84709D2XY) (WORMWOOD - UNII:E84709D2XY)	WORMWOOD	3 [hp_X]

WORMWOOD (UNII: F84709F2XV) (WORMWOOD - UNII:F84709F2XV)		WORMWOOD	in 1 mL	
GENTIANA LUTEA ROOT (UNII: S72O3284MS) (GENTIANA LUTEA ROOT - UNII:S72O3284MS)		GENTIANA LUTEA 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	
STRYCHNOS NUX-VOMICA SEED (UNII: 269XH13919) (STRYCHNOS NUX-VOMICA SEED - UNII:269XH13919)		STRYCHNOS NUX-VOMICA SEED	6 [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:48951-5007-1	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(48951-5007)

Revised: 8/2017

Uriel Pharmacy Inc.