

CACTUS CRATAEGUS SPECIAL ORDER- cactus crataegus special order 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.

Cactus Crataegus Special Order

Directions: FOR ORAL USE.

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

Active Ingredients: Cactus ex herba 3X, Crataegus e fol. et fruct. 3X, Melissa ex herba 3X

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 www.urielpharmacy.com

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Uriel, East Troy, WI 53120
www.urielpharmacy.com Lot:



**Cactus
Crataegus
Special Order**

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

Cactus Crataegus s.o.

CACTUS CRATAEGUS SPECIAL ORDER

cactus crataegus special order liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-3003
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SELENICEREUS GRANDIFLORUS STEM (UNII: 7114SV0MYK) (SELENICEREUS GRANDIFLORUS STEM - UNII:7114SV0MYK)	SELENICEREUS GRANDIFLORUS STEM	3 [hp_X] in 1 mL
HAWTHORN LEAF WITH FLOWER (UNII: 6OM09RPY36) (HAWTHORN LEAF WITH FLOWER - UNII:6OM09RPY36)	HAWTHORN LEAF WITH FLOWER	3 [hp_X] in 1 mL
MELISSA OFFICINALIS (UNII: VE70190J0N) (MELISSA OFFICINALIS		3 [hp_X]

MELISSA OFFICINALIS (UNII: YF70189L0N) (MELISSA OFFICINALIS - UNII:YF70189L0N)			MELISSA OFFICINALIS	5 Amp_AJ 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:48951-3003-1	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(48951-3003)

Revised: 5/2018

Uriel Pharmacy Inc.