

CACTUS CINIS AVENAE- cactus cinis avenae 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 Cinis Avenae

Directions: FOR ORAL USE.

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

Active Ingredients: Cactus (Queen of the night) 3X, Crataegus (Hawthorn) 3X,Magnesium phos. e cin. Avenae (Magnesium phosphate in ash of oat grains) 6X, Arnica 20X

Inactive Ingredients: Water, Salt, Lactose

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. Contains traces of lactose. 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
www.urielpharmacy.com



CACTUS CINIS AVENAE

cactus cinis avenae liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-3239
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
MAGNESIUM PHOSPHATE, TRIBASIC, PENTAHYDRATE (UNII: 453COF7817) (MAGNESIUM CATION - UNII:T6V3LHY838)		MAGNESIUM PHOSPHATE, TRIBASIC, PENTAHYDRATE	6 [hp_X] in 1 mL
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
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ) (ARNICA MONTANA FLOWER - UNII:OZ0E5Y15PZ)		ARNICA MONTANA FLOWER	20 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-3239-1	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(48951-3239)

Revised: 12/2020

Uriel Pharmacy Inc.