SECALE CORNUTUM- claviceps purpurea sclerotium liquid Washington Homeopathic Products

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.

DRUG FACTS

ACTIVE INGREDIENTS

SECALE

USES

To relieve the symptoms of restlessness.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

SECALE Restlessness

STOP USE AND ASK DOCTOR

If symptoms persist/worsen or if pregnant/nursing, consult your practitioner.

DIRECTIONS

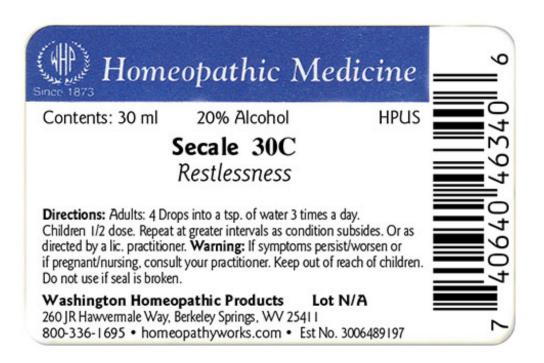
Adults: 4 drops into a tsp. of water 3 times a day. Children: 1/2 dose. Repeat at greater intervals as condition subsides. Or as directed by a lic. practitioner.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL

The OTC potency range of SECALE is 3x-30x, 2c-30c, 200c, 1m, 10m, 50m, and CM. Availability is subject to change.



ll WHP single remedies are made to order; thus, the labels are printed on the same label stock, as the orders are filled.

'Bottle Size,' 'Potency,' and 'Alcohol Percentage' vary on the label depending on customer choice. Standard bottle sizes for dilution-form remedies are 15ml, 30ml, 50ml, and 100ml.

SECALE CORNUTUM

claviceps purpurea sclerotium liquid

Draduct	Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71919-618
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLAVICEPS PURPUREA SCLEROTIUM (UNII: 0 1G9 XEA93N) (CLAVICEPS PURPUREA SCLEROTIUM - UNII: 0 1G9 XEA93N)	CLAVICEPS PURPUREA SCLEROTIUM	30 [hp_C] in 1 mL

Inactive 1	Ingredients
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Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

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Color	white (white)	Score		
Shape		Size		
Flavor		Imprint Code		

Contains

ŀ	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:71919-618- 07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/10/2009		
2	NDC:71919-618- 08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/10/2009		
3	NDC:71919-618- 09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/10/2009		
4	NDC:71919-618- 10	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/10/2009		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/10/2009	

Labeler - Washington Homeopathic Products (084929389)

Establishment				
Name	Address	ID/FEI	Business Operations	
Washington Homeopathic Products		084929389	manufacture(71919-618)	

Revised: 12/2009 Washington Homeopathic Products