CROTALUS CASCAVELLA- crotalus durissus terrificus venom 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

CROTALUS CASC

USES

To relieve the symptoms of sharp pains.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

CROTALUS CASC Sharp pains

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.

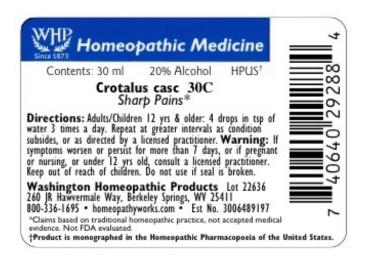
INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL

The OTC potency range of CROTALUS CASC is 8x–30x, 4c–30c, 200c, 1m, 10m, 50m, and CM.

Availability is subject to change.



All 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.

CROTALUS CASCAVELLA

crotalus durissus terrificus venom liquid

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71919-234

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CROTALUS DURISSUS TERRIFICUS VENOM (UNII: 2XF6 I0 446 G) (CROTALUS DURISSUS TERRIFICUS VENOM - UNII: 2XF6 I0 446 G)	CROTALUS DURISSUS TERRIFICUS VENOM	30 [hp_C] in 1 mL

Inactive Ingredients

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

Product Characteristics

Color	white (white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71919-234- 07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	06/05/2013	
2	NDC:71919-234- 08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	06/05/2013	
3	NDC:71919-234- 09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	06/05/2013	
4	NDC:71919-234- 10	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	06/05/2013	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		06/05/2013		

Labeler - Washington Homeopathic Products (084929389)

Establishment				
Name	Address	ID/FEI	Business Operations	
Washington Homeopathic Products		084929389	manufacture(71919-234)	

Revised: 12/2018 Washington Homeopathic Products