RUTA GRAVEOLENS- ruta graveolens flowering top 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

RUTA

USES

To relieve the symptoms of bruised feeling.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

RUTA Bruised feeling

STOP USE AND ASK DOCTOR

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

DIRECTIONS

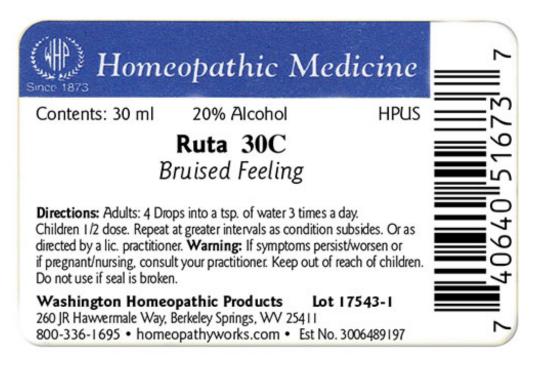
Adults: 4 drops into a tsp. of water 3 times a day. Children: 1/2 dose. Repeat at greater intervals as condition subsides. Or as directed by a lic. practitioner.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL

The OTC potency range of RUTA is 3x-30x, 2c-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.

RUTA GRAVEOLENS

ruta graveolens flowering top liquid

Dua duat	Information
Product	Intormation

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71919-595
Route of Administration	ORAL		

Active Ingredient/Active Moiety

П	S S		
	Ingredient Name	Basis of Strength St	trength
	RUTA GRAVEOLENS FLOWERING TOP (UNII: N94C2U587S) (RUTA GRAVEOLEN FLOWERING TOP - UNII:N94C2U587S)		[hp_C] 1 mL

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inditive ingredients			
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

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:71919-595- 07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/31/2009		
2	NDC:71919-595- 08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/31/2009		
3	NDC:71919-595- 09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/31/2009		
4	NDC:71919-595- 10	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/31/2009		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/31/2009	

Labeler - Washington Homeopathic Products (084929389)

Establishment				
Name	Address	ID/FEI	Business Operations	
Washington Homeopathic Products		084929389	manufacture(71919-595)	

Revised: 12/2009 Washington Homeopathic Products