

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 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			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71919-595
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
RUTA GRAVEOLENS FLOWERING TOP (UNII: N94C2U587S) (RUTA GRAVEOLENS FLOWERING TOP - UNII:N94C2U587S)		RUTA GRAVEOLENS FLOWERING TOP	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:719 19-595-07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/31/2009	
2	NDC:719 19-595-08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/31/2009	
3	NDC:719 19-595-09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/31/2009	
4	NDC:719 19-595-10	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(719 19-595)

Revised: 12/2009

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