RUTA GRAVEOLENS- ruta graveolens flowering top pellet 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, stop use and consult your practitioner.

DIRECTIONS

Adults: Dissolve 3 to 5 under the tongue three times a day or as directed by Lic. Practitioner. Take at greater intervals as condition subsides. Children: Dissolve 3 to 5 under the tongue three times a day or as directed by Lic. Practitioner. Take at greater intervals as condition subsides.

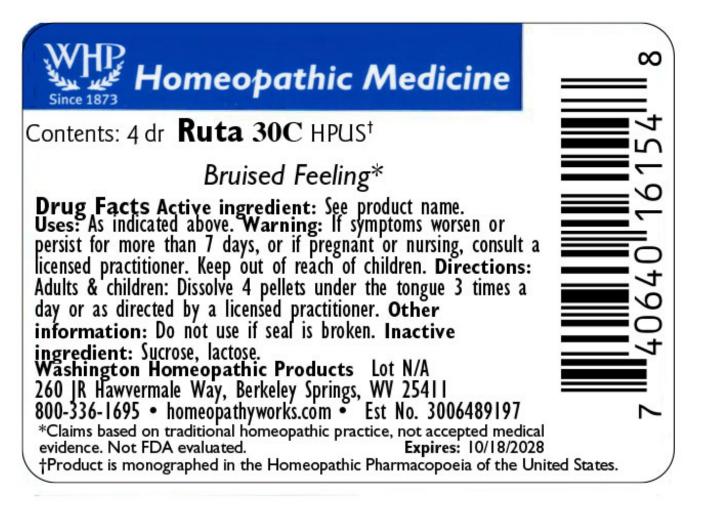
INACTIVE INGREDIENTS

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' and 'Potency' vary on the label depending on customer choice.

Standard bottle sizes for pellet-form remedies are 2 dram, 4 dram, 1 ounce, 2 ounce, and 4 ounce.

RUTA GRAVEOLENS ruta graveolens flowering top pellet						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:68428-609		
Route of Administration	ORAL					
Active Ingredient/Active Moiety						
Ingre	edient Name		Basis of S	trength	Strength	
RUTA GRAVEOLENS FLOWERING TOP (UNII: N94C2U587S) (RUTA GRAVEOLENS FLOWERING TOP - UNII:N94C2U587S)			RUTA GRAVEO FLOWERING TO		30 [hp_C]	

Inactive Ingredients					
	Ingredient Name		Strength		
SUCROSE (UNII: C151H8M554	L)				
LACTOSE (UNII: J2B2A4N98G)					
Product Characteristics					
Color	white (white)	Score			
COIOI					
		Size			
Shape Flavor		Size Imprint Code			

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:68428- 609-03	75 in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/31/2009		
2	NDC:68428- 609-05	150 in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/31/2009		
3	NDC:68428- 609-11	300 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/31/2009		
4	NDC:68428- 609-12	600 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/31/2009		
5	NDC:68428- 609-06	1200 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	
	approved meopathic		12/31/2009		

Labeler - Washington Homeopathic Products (084929389)

Establishment					
Name	Address	ID/FEI	Business Operations		
Washington Homeopathic Products		084929389	manufacture(68428-609)		

Revised: 10/2023

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