RUTA GRAV- ruta graveolens flowering top tablet Hyland's Inc.

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.

RUTA GRAV. 30X
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SPRAINS, BACKACHE OR EYE STRAIN
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Made according to the Homeopathic Pharmacopoeia of the United States since 1903.

Warnings

Do not use if cap band is missing or broken.

If you are pregnant or nursing, consult a licensed health care professional before using this product.

If symptoms persist for 7 days or worsen, contact a licensed practitioner.

Keep this and all medicines out of the reach of children.

To be used according to label indications and/or standard homeopathic indications.

Directions

Adults: Dissolve 4 tablets under tongue 4 times a day.

Children: consult a healthcare professional.

Inactive Ingredients

Acacia Gum and Lactose

Questions?

800-624-9659

PRINCIPAL DISPLAY PANEL - 250 Tablet Bottle Label

SINCE 1903 **Hyland's** [®] HOMEOPATHIC NDC 54973-2925-4
RUTA GRAV.
30X
SPRAINS, BACKACHE OR
EYE STRAIN *
250 TABLETS
*Claims are based on traditional homeopathic practice, not accepted medical evidence.

Not FDA evaluated.

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Standard Homeopathic Company Los Angeles, CA 90061 Questions? 800-624-9659



NDC 54973 - 2925 - 4 RUTA GRAV.

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RUTA GRAV

ruta graveolens flowering top tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54973-2925
Route of Administration	ORAL		

Active Ingredient/Active Moiety Ingredient Name RUTA GRAVEOLENS FLOWERING TOP (UNII: N94C2U587S) (RUTA GRAVEOLENS FLOWERING TOP - UNII:N94C2U587S) RUTA GRAVEOLENS FLOWERING TOP 30 [hp_X]

Inactive Ingredients		
Ingredient Name	Strength	
ACACIA (UNII: 5C5403N26O)		
LACTOSE (UNII: J2B2A4N98G)		

Product Characteristics			
Color	brown	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	
Contains			

ı	Packaging				
	# Item Cod	e Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:54973-2925-4	250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/1955		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/1955	

Labeler - Hyland's Inc. (008316655)

Establishment			
Name	Address	ID/FEI	Business Operations
Hyland's Inc.		008316655	manufacture(54973-2925) , pack(54973-2925) , label(54973-2925)

Revised: 12/2022 Hyland's Inc.