RUTA GRAVEOLENS 7133- ruta graveolens liquid Professional Complementary Health Formulas

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.

DRUT

ACTIVE INGREDIENTS

Ruta graveolens 6X, 9X, 12X, 30X

QUESTIONS

Professional Formulas
PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

Use according to standard homeopathic indications for self-limiting conditions. Common Materia Medica indications: muscle soreness, bruises, minor strains and sprains.*

*Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS

In case of overdose, get medical help or contact a poison control center right away. Keep out of the reach of children.

If pregnant or breastfeeding, ask a healthcare professional before use.

DIRECTIONS

Place drops under tongue 30 minutes before/after meals. Adults and children 12 years and over: Take 10 drops up to 3 times per day. Consult a physician for use in children under 12 years of age.

OTHER INFORMATION

Tamper resistant. If seal is broken, do not use. After opening, close container tightly and store at room temperature away from heat.

INACTIVE INGREDIENTS

20% ethanol, purified water.

LABEL

Est 1985

Professional Formulas

Complementary Health

Ruta graveolens

Homeopathic Remedy

1 FL. OZ. (29.5 mL)



RUTA GRAVEOLENS 7133

ruta graveolens liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-7133	
Route of Administration	ORAL			

l	Active Ingredient/Active Moiety				
l	Ingredient Name	Basis of Strength	Strength		
	RUTA GRAVEOLENS FLOWERING TOP (UNII: N94C2U587S) (RUTA GRAVEOLENS FLOWERING TOP - UNII:N94C2U587S)	RUTA GRAVEOLENS FLOWERING TOP	6 [hp_X] in 29.5 mL		

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

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l		NDC:63083- 7133-1	29.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/15/1985	

Marketing Information				
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
	08/15/1984			
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date		

Labeler - Professional Complementary Health Formulas (167339027)

Registrant - Natural Pharmaceutical Manufacturing LLC (015624923)

Establishment				
Name	Address	ID/FEI	Business Operations	
Natural Pharmaceutical Manufacturing LLC		015624923	manufacture(63083-7133)	

Revised: 8/2019 Professional Complementary Health Formulas