SINUS RELIEF FORMULA- pulsatilla (pratensis), sinapis nigra, luffa operculata, euphorbium officinarum, guaiacum, mercurius sulphuratus ruber spray Nutritional Specialties, 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.

DRUG FACTS:

ACTIVE INGREDIENTS:

Pulsatilla 4X, Sinapis Nigra 4X, Luffa Operculata 5X, Euphorbium Officinarum 8X, Guaiacum 12X, Mercurius Sulphuratus Ruber 12X.

PURPOSE:

Aids in temporary relief of sinus congestion and irritation.†

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

WARNINGS:

Professional Use Only

If pregnant or breast-feeding, ask a health professional before use.

In case of overdose, get medical help or contact a Poison Control Center right away.

If condition worsens, seek medical attention.

KEEP OUT OF REACH OF CHILDREN

Do not use if tamper evident seal is broken or missing.

Store in a cool place after opening

KEEP OUT OF REACH OF CHILDREN:

In case of overdose, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Adults: 2 to 3 sprays orally, three times daily. Children under twelve one half adult dosage. Do not take within 15 minutes of consuming food, beverage or brushing teeth. Consult a physician for use in children under 12 years of age.

INDICATIONS:

Aids in temporary relief of sinus congestion and irritation.†

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

INACTIVE INGREDIENTS:

Alcohol USP 20%, Purified Water USP.

QUESTIONS:

MANUFACTURED EXCLUSIVELY FOR NUTRITIONAL SPECIALTIES, INC. PO BOX 97227 PITTSBURG, PA 15229 www.phpltd.com

PACKAGE LABEL DISPLAY:

Professional
Health Products
HOMEOPATHIC
NDC 83027-0105-1
SINUS RELIEF
FORMULA
2 FL. OZ (60 ml)

Professional Use Only

WARNINGS: If pregnant or breastfeeding, ask a health professional before use. In case of overdose, get medical help or contact a Poison Control Center right away. If condition worsens, seek medical attention.

KEEP OUT OF REACH OF CHILDREN

Do not use if tamper evident seal is broken or missing.

Store in a cool place after opening.

[†] Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.





NDC 83027-0105-1



Active Ingredients: Pulsatilla 4x. Sinapis Nigra 4x. Luffa Operculata 5x. Euphorbium Officinarum 8x. Guaiacum 12x. Mercurius Sulphuratus Ruber

12x.

Inactive Ingredients: Alcohol USP 20%, Purified Water USP.

2 FL. OZ. (60 ml)

Indication: Aids in temporary relief of sinus congestion and irritation.

Directions For Use: Adults: 2 to 3 sprays orally, three times daily. Children under twelve one half adult dosage. Do not take within 15 minutes of consuming food, beverage or brushing teeth. Consult a physician for use in children under 12 years of age.



SINUS RELIEF FORMULA

pulsatilla (pratensis), sinapis nigra, luffa operculata, euphorbium officinarum, guaiacum, mercurius sulphuratus ruber spray

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83027-0105		
Route of Administration	ORAL				

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
PULSATILLA PRATENSIS WHOLE (UNII: 8E272251DI) (PULSATILLA PRATENSIS WHOLE - UNII:8E272251DI)	PULSATILLA PRATENSIS WHOLE	4 [hp_X] in 1 mL	
BLACK MUSTARD SEED (UNII: 8LTY55LQ8D) (BLACK MUSTARD SEED - UNII:8LTY55LQ8D)	BLACK MUSTARD SEED	4 [hp_X] in 1 mL	
LUFFA OPERCULATA FRUIT (UNII: C4MO6809HU) (LUFFA OPERCULATA FRUIT - UNII:C4MO6809HU)	LUFFA OPERCULATA FRUIT	5 [hp_X] in 1 mL	
EUPHORBIA RESINIFERA RESIN (UNII: 1TI109028K) (EUPHORBIA RESINIFERA RESIN - UNII:1TI109028K)	EUPHORBIA RESINIFERA RESIN	8 [hp_X] in 1 mL	
GUAIACUM OFFICINALE RESIN (UNII: N0K2Z502R6) (GUAIACUM OFFICINALE RESIN - UNII:N0K2Z502R6)	GUAIACUM OFFICINALE RESIN	12 [hp_X] in 1 mL	
MERCURIC SULFIDE (UNII: ZI0T668SF1) (MERCURIC CATION - UNII:ED30FJ8Y42)	MERCURIC SULFIDE	12 [hp_X] in 1 mL	

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

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:83027- 0105-1	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/28/2023	

Marketing Information				
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
	09/28/2023			
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date		

Labeler - Nutritional Specialties, Inc. (032744609)