POLYHESION FORMULA- sanguinaria canadensis, nitricum acidum, teucrium marum, thuja occidentalis, sanguinarinum nitricum, calcarea carbonica, phosphorus 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:

Sanguinaria Canadensis 3X, Nitricum Acidum 6X, Teucrium Marum 6X, , Thuja Occidentalis 6X, Sanguinarinum Nitricum 15X, Calcarea Carbonica 30X, Phosphorus 30X.

PURPOSE:

Aids in temporary relief of cramping and pain in the abdomen.†

†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 cramping and pain in the abdomen.†

†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 83027-0104-1
POLYHESION
FORMULA
2 FL. OZ (60 ml)

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NDC 83027-0104-1

POLYHESION FORMULA

C50 (reorder code)

Active Ingredients: Sanguinaria Canadensis 3x. Nitricum Acidum 6x. Teucrium Marum 6x. Thuja Occidentalis 6x. Sanguinarinum Nitricum 15x. Calcarea Carbonica 30x. Phosphorus 30x.

Inactive Ingredients: Alcohol USP 20%,

Purified Water USP.

2 FL. OZ. (60 ml)

Indication: Aids in temporary relief of cramping and pain in the abdomen.[†]

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.



POLYHESION FORMULA

sanguinaria canadensis, nitricum acidum, teucrium marum, thuja occidentalis, sanguinarinum nitricum, calcarea carbonica, phosphorus spray

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83027-0104		
Route of Administration	ORAL				

Active Ingredient/Active Moiety						
Ingredient Name	Basis of Strength	Strength				
SANGUINARIA CANADENSIS ROOT (UNII: N9288CD508) (SANGUINARIA CANADENSIS ROOT - UNII:N9288CD508)	SANGUINARIA CANADENSIS ROOT	3 [hp_X] in 1 mL				
NITRIC ACID (UNII: 411VRN1TV4) (NITRIC ACID - UNII:411VRN1TV4)	NITRIC ACID	6 [hp_X] in 1 mL				
TEUCRIUM MARUM WHOLE (UNII: 10464S0TAA) (TEUCRIUM MARUM - UNII:10464S0TAA)	TEUCRIUM MARUM WHOLE	6 [hp_X] in 1 mL				
THUJA OCCIDENTALIS LEAFY TWIG (UNII: 1NT28V9397) (THUJA OCCIDENTALIS LEAFY TWIG - UNII:1NT28V9397)	THUJA OCCIDENTALIS LEAFY TWG	6 [hp_X] in 1 mL				
SANGUINARINE NITRATE (UNII: BOP122S791) (NITRATE ION - UNII:T93E9Y2844)	SANGUINARINE NITRATE	15 [hp_X] in 1 mL				
OYSTER SHELL CALCIUM CARBONATE, CRUDE (UNII: 2E32821G6I) (OYSTER SHELL CALCIUM CARBONATE, CRUDE - UNII: 2E32821G6I)	OYSTER SHELL CALCIUM CARBONATE, CRUDE	30 [hp_X] in 1 mL				
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	30 [hp_X] in 1 mL				

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

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:83027- 0104-1	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/12/2023			
Marketing Information						
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
	approved meopathic		09/12/2023			

Labeler - Nutritional Specialties, Inc. (032744609)

Revised: 9/2023 Nutritional Specialties, Inc.