

**POISON IVY AND OAK FORMULA- glandula suprarenalis (bovine), hepar suis, rhus tox, urtica dioica, rhus aromatica, rhus glabra, histaminum hydrochloricum, rhus diversiloba, adrenocorticotrophin 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:

Glandula Suprarenalis (Bovine) 6X, 12X, Hepar Suis (Liver) 6X, 12X, Rhus Tox (Poison Ivy) 6X, 12X, 60X, 100X, Urtica Dioica (Stinging Nettle) 6X, 12X, 60X, 100X, Rhus Aromatica (Fragrant Sumac) 8X, 12X, 60X, 100X, Rhus Glabra 8X, 12X, 60X, 100X, Histaminum Hydrochloricum 12X, Rhus Diversiloba (Pacific Poison Oak) 16X, 30X, 60X, 100X, Adrenocorticotrophin 15C, 30C.

PURPOSE:

Provides factors that mitigate hypersensitivity to poison ivy and poison oak.

†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

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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:

Provides factors that mitigate hypersensitivity to poison ivy and poison oak.†
†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 83027-0039-1

POISON IVY &

OAK FORMULA

2 FL. OZ (60 ml)

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POISON IVY & OAK FORMULA

A263 (reorder code)

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

2 FL. OZ. (60 ml)

Indication: Provides factors that mitigate hypersensitivity to poison ivy and poison oak.†

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.

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POISON IVY AND OAK FORMULA

glandula suprarenalis (bovine), hepar suis, rhus tox, urtica dioica, rhus aromatica, rhus glabra, histaminum hydrochloricum, rhus diversiloba, adrenocorticotrophin spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83027-0039
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BOS TAURUS ADRENAL GLAND (UNII: M2776SWB29) (BOS TAURUS ADRENAL GLAND - UNII:M2776SWB29)	BOS TAURUS ADRENAL GLAND	6 [hp_X] in 1 mL
PORK LIVER (UNII: 6EC706HI7F) (PORK LIVER - UNII:6EC706HI7F)	PORK LIVER	6 [hp_X] in 1 mL
TOXICODENDRON PUBESCENS LEAF (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)	TOXICODENDRON PUBESCENS LEAF	6 [hp_X] in 1 mL
URTICA DIOICA WHOLE (UNII: 710FLW4U46) (URTICA DIOICA - UNII:710FLW4U46)	URTICA DIOICA WHOLE	6 [hp_X] in 1 mL
RHUS AROMATICA ROOT BARK (UNII: Q3H36W0J42) (RHUS AROMATICA ROOT BARK - UNII:Q3H36W0J42)	RHUS AROMATICA ROOT BARK	8 [hp_X] in 1 mL
RHUS GLABRA TOP (UNII: RHH784E0K6) (RHUS GLABRA TOP - UNII:RHH784E0K6)	RHUS GLABRA TOP	8 [hp_X] in 1 mL
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - UNII:820484N8I3)	HISTAMINE DIHYDROCHLORIDE	12 [hp_X] in 1 mL
TOXICODENDRON DIVERSILOBUM LEAF (UNII: V727AWD6ZZ) (TOXICODENDRON DIVERSILOBUM LEAF - UNII:V727AWD6ZZ)	TOXICODENDRON DIVERSILOBUM LEAF	16 [hp_X] in 1 mL
CORTICOTROPIN (UNII: K0U68Q2TXA) (CORTICOTROPIN - UNII:K0U68Q2TXA)	CORTICOTROPIN	15 [hp_C] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)

ALCOHOL (UNII: 3K9958V90M)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83027-0039-1	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/29/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/29/2023	

Labeler - Nutritional Specialties, Inc. (032744609)

Revised: 3/2023

Nutritional Specialties, Inc.