

**RHUS TOX 4X / 8X / 12X- rhus toxicodendron liquid**  
**Natural Creations, 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.*

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**RHUS TOX 4X/8X/12X**

ACTIVE INGREDIENTS (HPUS) - equal amounts of: Rhus Toxicodendron 4X, 8X, 12X.

USES: Temporarily relieves intense itching & skin irritation from contact with Poison Ivy &/or Poison Oak.\*\*

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DIRECTIONS: Adults and children above 12 years: 10 drops orally 3 times daily, or as directed by a health care professional.

**KEEP OUT OF THE REACH OF CHILDREN.** In case of overdose (or accidental ingestion) get medical help or contact a Poison Control Center right away.

**WARNINGS:**

- Consult a physician for use in children under 12 years of age.
- **IF PREGNANT OR BREAST-FEEDING**, ask a health care professional before use.
- **KEEP OUT OF THE REACH OF CHILDREN.** In case of overdose (or accidental ingestion) get medical help or contact a Poison Control Center right away.
- Do not use if **TAMPER EVIDENT** seal is broken or missing.

INACTIVE INGREDIENTS: Ethyl Alcohol USP, Purified Water.

**QUESTIONS & COMMENTS?**

Natural Creations, Inc. / Woodbine, IA 51579 / 712-647-1600

\*The letters "HPUS" indicate the components in the products are officially monographed in the Homeopathic Pharmacopeia of the United States.

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

NDC: 43406-0216-1

RHUS TOX 4X / 8X / 12X

Homeopathic

1 fl oz (30 mL) / 20% Alcohol



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NCHS-1887

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rhus toxicodendron liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:43406-0216
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>TOXICODENDRON PUBESCENS LEAF</b> (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)	TOXICODENDRON PUBESCENS LEAF	12 [hp_X] in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>WATER</b> (UNII: 059QF0KO0R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43406-0216-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	02/20/2009	

**Marketing Information**

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
unapproved homeopathic		02/20/2009	

**Labeler** - Natural Creations, Inc (018022074)

### Establishment

Name	Address	ID/FEI	Business Operations
OHM Pharma, Inc		030572478	manufacture(43406-0216)

Revised: 3/2022

Natural Creations, Inc