

RHUS TOXICODENDRON 200C- 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.

Rhus Toxicodendron 200C

ACTIVE INGREDIENTS (HPUS*): Rhus Toxicodendron 200C

USES: Temporarily relieves irritation from red, itchy, blistering skin rash.**

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DIRECTIONS: Adults & 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 product are officially monographed in the Homeopathic Pharmacopeia of the United States.

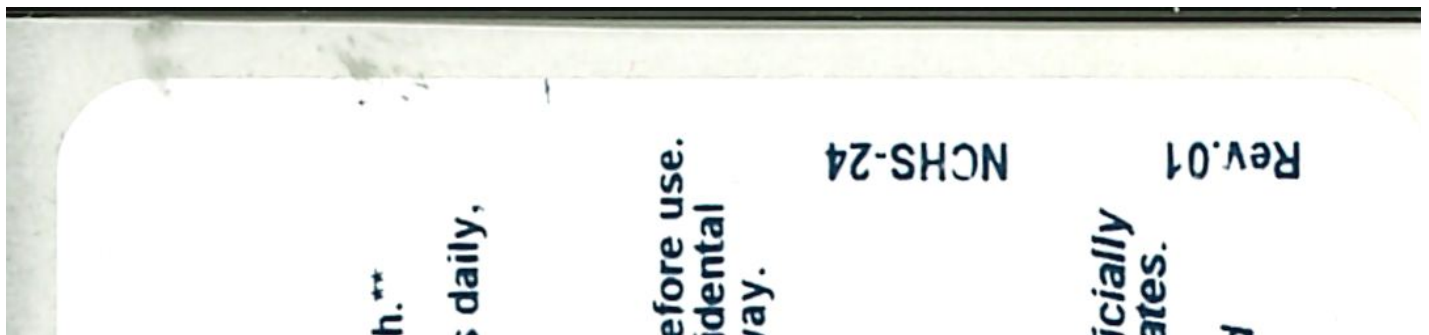
**Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

NDC:43406-0255-1

Rhus Toxicodendron 200C

HOMEOPATHIC

1 fl oz (30mL) / 20% Alcohol





55-1

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DRON



HIC

Alcohol

natural
crea

NDC: 43406-02

RHUS
TOXICODENI
200C

HOMEOPAT

1 fl oz (30 mL) / 20%

RHUS TOXICODENDRON 200C

rhus toxicodendron liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43406-0255
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TOXICODENDRON PUBESCENS LEAF (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)	TOXICODENDRON PUBESCENS LEAF	200 [hp_C] in 1 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43406-0255-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/18/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/18/2012	

Labeler - Natural Creations, Inc (018022074)

Establishment

Name	Address	ID/FEI	Business Operations
OHM Pharma, Inc		030572478	manufacture(43406-0255)

Revised: 12/2018

Natural Creations, Inc