RHUS TOX 4X, 8X, 12X- rhus toxicodendron. liquid
OHM PHARMA 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 TOX 4X, 8X, 12X

ACTIVE INGREDIENTS (HPUS*):

Rhus Toxicodendron 4X, 8X, 12X

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

USES: Temporarily relieves blistering rash.**

*** These statements have not been reviewed by the FDA. They are based on traditional homeopathic practice.*

Temporarily relieves blistering rash.**

DIRECTIONS: Adults & children above 12 years: 3 dropperfulls in mouth and hold for 1 minute before swallowing. Repeat once a week for 3 weeks, or as directed by a health care professional.

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.

Keep out of reach of children.

INACTIVE INGREDIENTS: Purified Water, Ethyl Alcohol USP.

QUESTIONS & COMMENTS?:

Little Drug / Sweetwater, TN 37874 / 423-337-7933

LITTLE DRUG

FAMILY WELLNESS CENTER

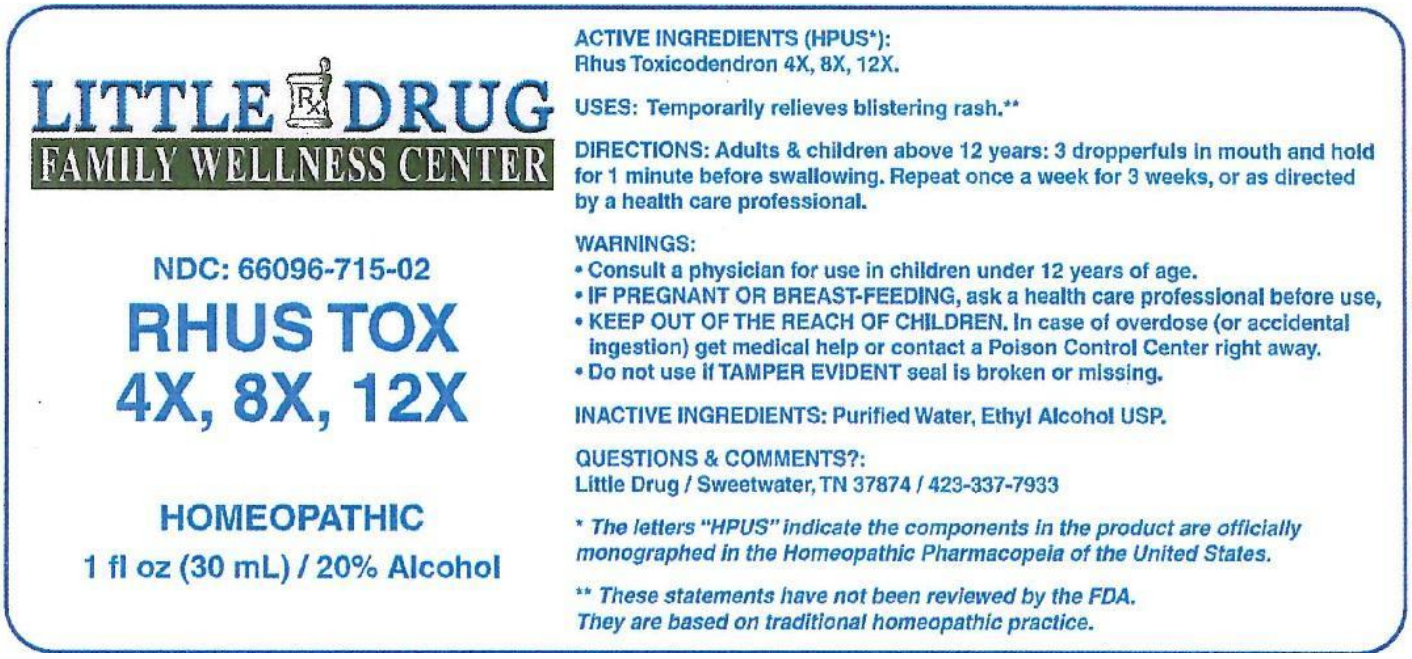
NDC: 66096-715-02

RHUS TOX

4X, 8X, 12X

HOMEOPATHIC

1 fl oz (30 mL) / 20% Alcohol



RHUS TOX 4X, 8X, 12X

rhus toxicodendron. liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66096-715
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	4 [hp_X] in 30 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:66096-715-02	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/06/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/06/2017	

Labeler - OHM PHARMA INC. (030572478)

Registrant - OHM PHARMA INC. (030572478)

Establishment

Name	Address	ID/FEI	Business Operations
OHM PHARMA INC.		030572478	manufacture(66096-715)

Revised: 3/2017 OHM PHARMA INC.