RHUS TOX 6X/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 6x/12x

ACTIVE INGREDIENT (HPUS*):

Rhus Toxicodendron 6X/12X

*The letters "HPUS" indicate that the components in this 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.

USES: For Temporary relief of red, swollen, itchy skin.**

SUGGESTED DOSAGE:

3 droppers full 1 time per week for 3 weeks, then 1 time per month thereafter. Hold solution in mouth for 30 seconds then swallow.

Can be taken any time of year but ideal to begin administering during Poision Ivy dormant phase (winter) often starting in February

in the Midwest.

WARNINGS:

If pregnant or nursing, ask a health care professional before using.

Keep out of reach of children. In case of an accidental overdose, seek professional assistance or contact a poison control center immediately.

Keep out of reach of children.

Alcohol

QUESTIONS & COMMENTS?

Preckshot Pharmacy / Peoria, IL 61614 / 309-649-2047

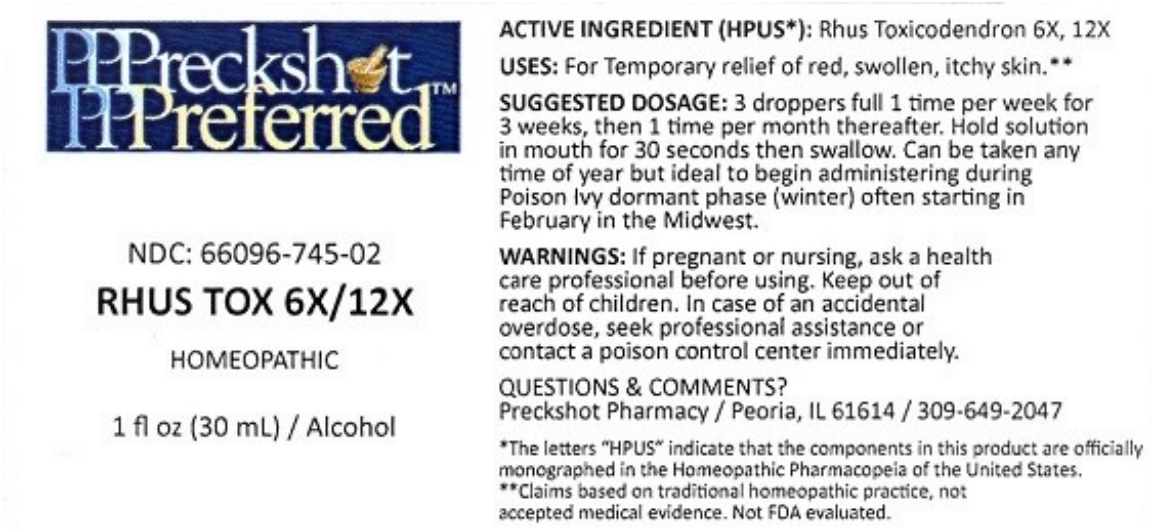
Preckshot Preferred ™

NDC: 66096-745-02

RHUS TOX 6X/12X

HOMEOPATHIC

1 fl oz (30 mL) / Alcohol



For Temporary relief of red, swollen, itchy skin.**

RHUS TOX 6X/12X

rhus toxicodendron. liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66096-745
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	6 [hp_X] in 30 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66096-745-02	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	10/30/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/30/2018	

Labeler - OHM PHARMA INC. (030572478)

Registrant - OHM PHARMA INC. (030572478)

Establishment

Name	Address	ID/FEI	Business Operations
OHM PHARMA INC.		030572478	manufacture(66096-745)

Revised: 10/2018 OHM PHARMA INC.