SKIN TAG RELIEF- thuja occidentalis leafy twig oil
Quest Products, 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.

Skin Tag Relief

Drug Facts

Active Ingredients

Thuja Occidentalis 6X (HPUS) 1%

The letters HPUS indicate the component(s) in this product is (are) officially monographed in the Homeopathic Pharmacopeia of the United States.

Purpose

Cutaneous Skin Tags Relief

Uses

Symptomatic treatment of Skin Tags.

Warnings

For external use only.

If pregnant or breast-feeding, ask a health professional before use.

Avoid using near the eyes or mouth.

Keep out of the reach of children. In case of ingestion call Poison Control Center hotline immediately at 1-800-222-1222.

Do not use if tamper evident seal is broken.

Directions

Apply 3 times daily to affected area. Skin tags will dry and flake away over several week period. Some individuals may be sensitive to essential oil. Skin test for tolerability. If irritation or reaction occurs, discontinue use.

Other ingredients

Cedar Leaf Oil, Melaleuca Alternifollia Leaf Oil (Tea Tree Oil), Ricinus Communis Seed Oil (Castor Oil).

There is no scientific evidence that this product works. The product's claims are based only on theories of homeopathy from the 1700's that are not acceptert by most modern medical experts.

PRINCIPAL DISPLAY PANEL - 10 mL Bottle Carton

SKIN TAG RELIEF

thuja occidentalis leafy twig oil

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68229-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
THUJA OCCIDENTALIS LEAFY TWIG (UNII: 1NT28V9397) (THUJA OCCIDENTALIS LEAFY TWIG - UNII:1NT28V9397)	THUJA OCCIDENTALIS LEAFY TWIG	6 [hp_X] in 10 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CEDAR LEAF OIL (UNII: BJ169U4NLG)	
TEA TREE OIL (UNII: VIF565UC2G)	
CASTOR OIL (UNII: D5340Y2I9G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68229-100-02	1 in 1 CARTON	05/21/2013	
1	NDC:68229-100-01	10 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		
2	NDC:68229-100-04	1 in 1 CARTON	05/23/2022	
2	NDC:68229-100-03	12.5 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/21/2013	

Labeler - Quest Products, Inc. (075402441)

Establishment

Name	Address	ID/FEI	Business Operations
Fill Tech USA		926433855	manufacture(68229-100)

Revised: 11/2023 Quest Products, Inc.