THUJA- thuja occidentalis leafy twig liquid
PEKANA Naturheilmittel GmbH

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.

Thuja™

Indications

For skin tags and warts. Application of this homeopathic remedy for the designated usage is exclusively based on homeopathic experience. With severe forms of this disease, a clinically proven therapy is indicated.

Dosage

Unless otherwise prescribed, adults take 15-20 drops, 3 times per day. For pediatric dosages, consult your practitioner.

Warning

If symptoms persist, contact a licensed practitioner. If you have known sensitivity to any of the ingredients, please consult your licensed practitioner before use. If you are pregnant or nursing a baby, seek the advice of a health care professional before use.

Keep out of the reach of children.

Protect from light and heat.

Tamper Evident

Do not use this product if tamper evident strip is broken or removed from base of cap.

Ingredients

Thuja occidentalis	3X

Contains 69% alcohol by volume.

To report adverse events, contact BioResource at 321B Blodgett Street, Cotati, CA 94931

Manufactured by:
PEKANA® GmbH
D-88353 Kisslegg

Distributed by: BioResource Inc.

Cotati, CA 94931

PRINCIPAL DISPLAY PANEL - 50 ml Bottle Box

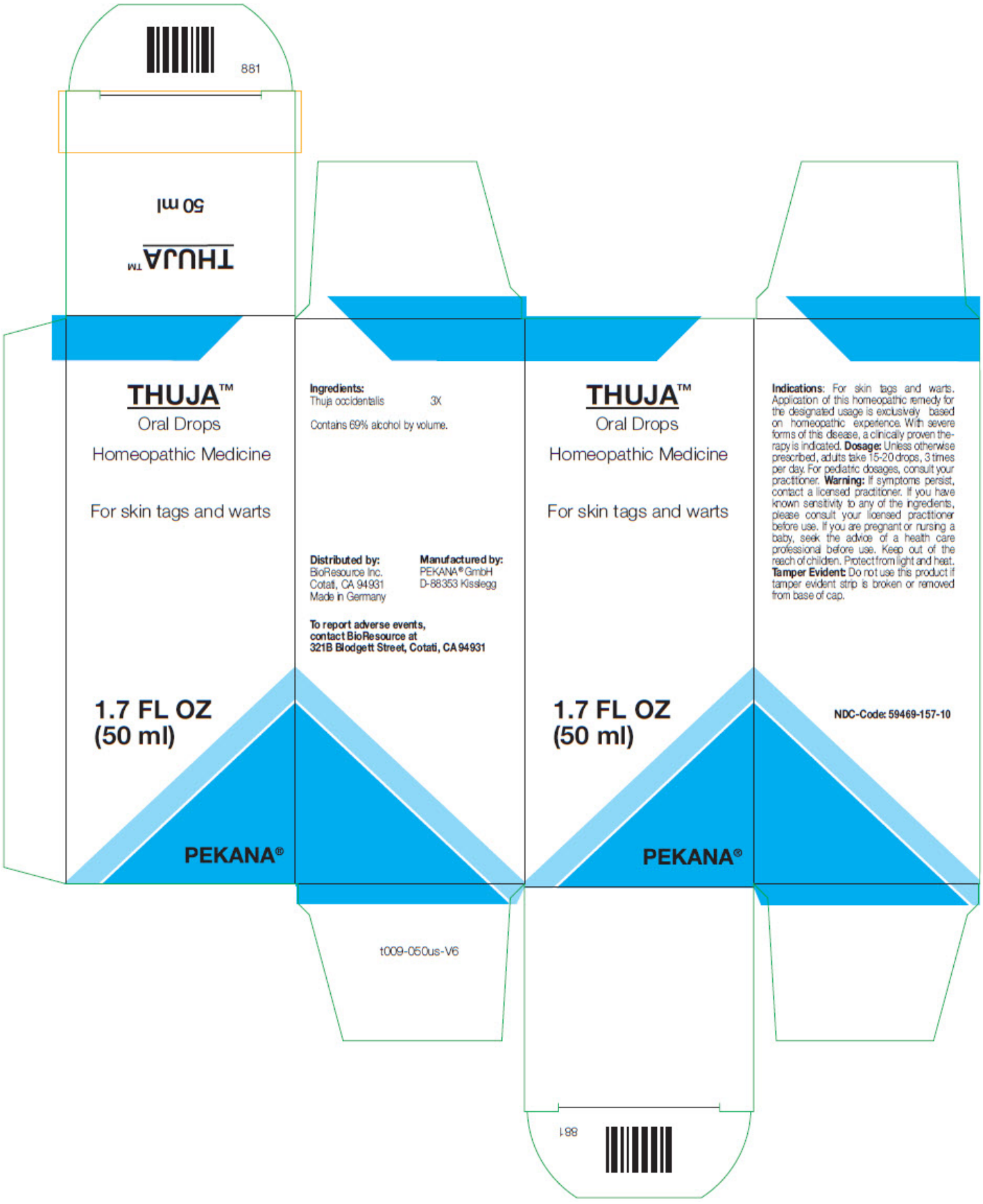
THUJA™
Oral Drops

Homeopathic Medicine

For skin tags and warts

1.7 FL OZ
(50 ml)

PEKANA®



THUJA

thuja occidentalis leafy twig liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59469-157
Route of Administration	ORAL		

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	3 [hp_X] in 50 mL

Inactive Ingredients

Ingredient Name	Strength
Alcohol (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59469-157-10	1 in 1 BOX	05/01/2005	
1		50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED HOMEOPATHIC		05/01/2005	

Labeler - PEKANA Naturheilmittel GmbH (320344542)

Establishment

Name	Address	ID/FEI	Business Operations
PEKANA GmbH		320344542	MANUFACTURE(59469-157)

Revised: 12/2022 PEKANA Naturheilmittel GmbH