# THUJA OCCIDENTALIS - thuja occidentalis pellet Newton Laboratories, 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.

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#### Thuja occidentalis

#### INDICATIONS & USAGE SECTION

Warts; Brown spots on hand & arms; Ovarian cysts; Vaccination effects; Dandruff; Diarrhea.

#### **DOSAGE & ADMINISTRATION SECTION**

**Directions:** Ages 12 and up, take 6 pellets by mouth (ages 0 to 11, give 3 pellets) as needed or as directed by a health professional. Under age 2, crush/dissolve pellets in purified water. Sensitive persons begin with 1 pellet and gradually increase to full dose.

#### **OTC - ACTIVE INGREDIENT SECTION**

Thuja occidentalis 15x, 10x, 200c, 30c.

#### **OTC - PURPOSE SECTION**

Warts; Brown spots on hand & arms; Ovarian cysts; Vaccination effects; Dandruff; Diarrhea.

#### INACTIVE INGREDIENT SECTION

**Inactive Ingredients:** Gluten-free, non-GMO, organic beet-derived sucrose (lactose free) pellets.

#### QUESTIONS SECTION

www.newtonlabs.net Newton Laboratories, Inc. FDA Est # 1051203 - Conyers, GA 30012 Questions? 1.800.448.7256

#### WARNINGS SECTION

**WARNINGS: Keep out of reach of children. Do not use** if tamper-evident seal is broken or missing. If symptoms worsen or persist for more than a few days, consult a doctor. If **pregnant or breast-feeding**, ask a doctor before use.

#### OTC - PREGNANCY OR BREAST FEEDING SECTION

If **pregnant or breast-feeding**, ask a doctor before use.

#### OTC - KEEP OUT OF REACH OF CHILDREN SECTION

Keep out of reach of children.

#### **PACKAGE LABEL**



### THUJA OCCIDENTALIS

thuja occidentalis pellet

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55714-7540

Route of Administration ORAL

## Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength		
Thuja Occidentalis Leafy Twig (UNII: 1NT28 V9397) (Thuja Occidentalis Leafy Twig -	Thuja Occidentalis Leafy	15 [hp_X]		
UNII:1NT28 V9 39 7)	Twig	in 1 g		

# **Inactive Ingredients**

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Ingredient Name	Strength

Sucrose (UNII: C151H8M554)

#### **Packaging**

	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:55714-7540-1	28 g in 1 BOTTLE, GLASS		

# **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2011	

# Labeler - Newton Laboratories, Inc. (788793610)

# **Registrant -** Newton Laboratories, Inc. (788793610)

Establishment				
Name	Address	ID/FEI	Business Operations	
Newton Laboratories, Inc.		788793610	MANUFACTURE(55714-7540)	

Revised: 9/2011 Newton Laboratories, Inc.