CHELIDONIUM CURCUMA THUJA- chelidonium curcuma thuja liquid
Uriel Pharmacy 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.

Chelidonium Curcuma Thuja

Directions: FOR TOPICAL USE ONLY.

Apply topically 3-4 times daily. Under age 2: Consult a doctor.

Active Ingredients: Chelidonium (Greater celandine) 1X, Curcuma (Turmeric) 1X, Thuja (American arborvitae) 1X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol

"prepared using rhythmical processes"

Use: Temporary relief of warts.

KEEP OUT OF REACH OF CHILDREN.

Warnings: FOR EXTERNAL USE ONLY.
Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions, if conditions worsen or persist, or accidental ingestion occurs. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing. Shake Well Before Use.

Questions? Call 866.642.2858 Made with care by Uriel, East Troy, WI 53120 shopuriel.com Lot:

CHELIDONIUM CURCUMA THUJA
chelidonium curcuma thuja liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-3078

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHELIDONIUM MAJUS ROOT (UNII: FLT36UCF0N) (CHELIDONIUM MAJUS ROOT - UNII:FLT36UCF0N)	CHELIDONIUM MAJUS ROOT	1 [hp_X] in 1 mL
TURMERIC (UNII: 856YO1Z64F) (TURMERIC - UNII:856YO1Z64F)	TURMERIC	1 [hp_X] in 1 mL
THUJA OCCIDENTALIS WHOLE (UNII: 5HBV6WCE3N) (THUJA OCCIDENTALIS WHOLE - UNII:5HBV6WCE3N)	THUJA OCCIDENTALIS WHOLE	1 [hp_X] in 1 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:48951-3078-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-3078)

Revised: 4/2024 Uriel Pharmacy Inc.