## WART REMOVER- zitalon wart removal treatment liquid liquid Ehy Holdings LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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81799-004

**Wart Remover** 

## **Active Ingredient(s)**

Salicylic Acid 12%

### **Purpose**

Wart remover

#### Use

use

for the removal of common and plantar warts

the common war is easy ily recognized by the rough macauli flowerlike "appearance of the surface

the plantar wart is recognized by its location only on the bottom of the foot . its tenderness and the interru option of the foot pró nt pattern

# Warnings

Warnings For externall use

#### Do not use

Do not use

on irritated skin or on any area that is infected or reddened

Do not use on moles skin tags , birthmarks , warts with hair growing from them , genital warts , or warts on the face or mucous membranes

Ask a doctor before you use if you have diabetes or poor blood circulation Stop use and ask a doctor if discomfort persists

Keep out of reach of children

If product gets into eyes a flush with water for

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IF swallowed, get medical help or contact a Poison Control Center right away

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### **Directions**

Directions:

Wash the affected area and dry it
Apply this product with a cotton swab to the wart
Repeat this procedure twice daily as needed ( until wart is removed ) for up to 12 weeks

#### Other information

Other information Store at room temperature and out of direct sunlight skin discoloration may occur during or affter use

## **Inactive ingredients**

Water

BRUCEA JAVANICA FRUIT

Coptis Chinensis Root Extract

Phellodendron Amurense Bark Extract

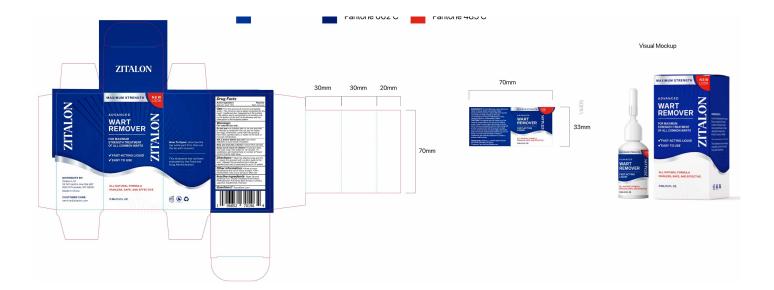
LONICERA JAPONICA TOP

Triclosan

Borneol

# Package Label - Principal Display Panel

81799-004-01



# **WART REMOVER**

zitalon wart removal treatment liquid liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81799-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	12 mg in 100 mL	

Inactive Ingredients			
Ingredient Name	Strength		
COPTIS CHINENSIS ROOT (UNII: CXS4LJR7EL)			
WATER (UNII: 059QF0KO0R)			
PHELLODENDRON AMURENSE BARK (UNII: PBG27B754G)			
BRUCEA JAVANICA FRUIT (UNII: 50268139XV)			
BORNEOL (UNII: M89NIB437X)			
TRICLOSAN (UNII: 4NM5039Y5X)			
LONICERA JAPONICA TOP (UNII: 60Q6UP226V)			

Packaging				
#	Item Code Package Description		Marketing Start Date	Marketing End Date
1	NDC:81799-004- 01	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/25/2022	

# **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358B	10/25/2022	

# Labeler - Ehy Holdings LLC (117322715)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Ehy Holdings LLC		117322715	manufacture(81799-004)	

Revised: 10/2022 Ehy Holdings LLC