# DR. LUKE MOLLUSCUM WART REMOVER- molluscum wart remover liquid Dr.luke Healthcare 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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83176-008-01 DR. LUKE MOLLUSCUM WART REMOVER

# **Active Ingredient**

Salicylic Acid 1% Chlorhexidine Acetate 1%

## **Purpose**

Wart Remover Antivira

#### Use

For genital wart, molluscum contagiosum, and small size skin tag,common wart, flat wart, filiform wart

## **Warnings**

For external use only.

Children need to use it under adult supervision.

Please tighten the cap immediately after using it.

Avoid contact with eyes, flush with water immediately if it getsinto eyes.

A small number of users may experience a slight burning ortingling sensation after use..

Minor rash and swelling reactions may occur, needn't stop using

#### Do not use

If you are pregnant, breastfeeding
If you are allergic to this product
On children under 10 years
On unhealed wounds after surgery

#### WHEN USING SECTION

For external use only, don't contact with eyes or swallow.

Children should use it under the supervision of adults.

Keep out of reach of children.

The liquid can get easily volatilized & crystallized, tighten the cap after use.

Avoid long-term contact with air while using.

### **STOP USE section**

you are experiencing severerash, edema, or severe painallergic reactions, local ulcers, heavy

#### ASK DOCTOR

you are experiencing severerash, edema, or severe painallergic reactions, local ulcers, heavy

### **KEEP OUT OF REACH OF CHILDREN**

Keep out of reach of children.

#### **Directions**

- 1. Wash wart and surrounding skin with rubbing alcohol orastringent solution (e.g.: Potassium Permanganate solution), anddry the area thoroughly.
- 2. Apply a small amount of liquid solution onto the wart with acotton swab, and press the wart 2-3 times. Avoid applying onhealthy skin.
- 3. Do not apply the liquid over 1 ml onto the affected area. Afterapplying, let the affected area dry.
- 4. Use once in the morning and evening continuously for 7 daysThe wart will begin to fall off after the last day of treatment. stopusing, and observe for 3 days.Repeat the treatment p rocess if the wart has not yet fallen off.lf swallowed,rinse immediately with water, and get medicalhelp or contact a poison control center right away

# **Inactive ingredients**

Cortex Phellodendri, Chinensis, Radix Isatidis, Radix Arnebiae. Semen Coicis Carthamus Tinctorius L, Angelica Sinensis, Rhizoma Chuanxiong, Crassostrea Gigas.

# **QUESTIONS?**

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#### PRINCIPAL DISPLAY PANEL

83176-008-01



# DR. LUKE MOLLUSCUM WART REMOVER

molluscum wart remover liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83176-008
Route of Administration	TOPICAL		

	Active Ingredient/Active Moiety			
	Ingredient Name	<b>Basis of Strength</b>	Strength	
	CHLORHEXIDINE (UNII: R4KO0DY52L) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE	1 mg in 100 mL	
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)   SALICYLIC ACID   1 mg in 100 mL	SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	1 mg in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength
ISATIS TINCTORIA ROOT (UNII: 8S10GFI6DX)	
COIX LACRYMA-JOBI VAR. MA-YUEN SEED (UNII: 8DW23817Z1)	
PHELLODENDRON CHINENSIS BARK (UNII: 2866QMZ434)	
CARTHAMUS TINCTORIUS WHOLE (UNII: 5EMV416J82)	
ANGELICA SINENSIS ROOT (UNII: B66F4574UG)	
PACIFIC OYSTER (UNII: FX2S0D3781)	
ARNEBIA GUTTATA ROOT (UNII: 9XN41J8M2D)	
COPTIS DELTOIDEA ROOT (UNII: 1UIP402HTI)	

l	P	ackaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:83176-008- 01	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2023	

Marketing In	formation				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	M028	04/17/2023			

# Labeler - Dr.luke Healthcare LLC (118868014)

<b>Establishment</b>			
Na me	Address	ID/FEI	Business Operations
Dr.luke Healthcare LLC		118868014	label(83176-008) . manufacture(83176-008)

Revised: 4/2023 Dr.luke Healthcare LLC