

ELLOCY WART REMOVER- wart remover liquid

Aramode

83596-004

Active Ingredient

Salicylic acid 1%

Purpose

remove wart

Use

- for removal of common and plantar warts
- the plantar wart is recognized by its location only on the bottom of the foot, its tenderness, and the interruption of the footprint pattern

Warnings

For external use only

Do not use

- On damaged skin (cuts, abrasions, eczema, sunburn).
- If you are pregnant or breastfeeding

When Using

- Avoid contact with eyes. If product gets into the eyes, flush with water for 15 minutes.
- If irritation persists, seek medical attention immediately.

Stop Use

if irritation occurs

if an allergic reaction occurs

Ask Doctor

if you are pregnant or breastfeeding

if you have diabetes

Keep Out Of Reach Of Children

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Directions

- wash the affected area.
- dry thoroughly
- apply 2 times a day (2-3 applications each time).
- continue application for 5-7 days.

Other information

Store between 20 to 25°C (68 to 77F)

Protect from excessive heat above 40°C (104F)

Inactive ingredients

Fritillaria extract, Purslane extract, Safflower, Willow bark extract, Lactic acid, Borneol, Dimethyl sulfoxide, Chlorhexidine acetate

Questions

CUSTOMER SUPPORT Contact us at support@ellocy.com

PRINCIPAL DISPLAY PANEL

PACKAGE

BRAND / ELLOCY WART REMOVER

PACKAGE SIZE / 65×30×95 mm
 LABEL SIZE / 35 x 20mm



ELLOCY WART REMOVER

wart remover liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83596-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WILLOW BARK (UNII: S883J9JDYX)	
FRITILLARIA DELAVAYI BULB (UNII: AG6Q756AT7)	
SAFFLOWER (UNII: 4VBL71TY4Y)	
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)	
PURSLANE (UNII: M6S840WVG5)	
BORNEOL (UNII: M89NIB437X)	
WATER (UNII: 059QF0KO0R)	
LACTIC ACID (UNII: 33X04XA5AT)	
CHLORHEXIDINE ACETATE (UNII: 5908ZUF22Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83596-004-01	5 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/02/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	09/02/2024	

Labeler - Aramode (963192477)**Establishment**

Name	Address	ID/FEI	Business Operations
Aramode		963192477	manufacture(83596-004)

Revised: 9/2024

Aramode