

HEALMUSZ WART REMOVER- salicylic acid 17% wart remover liquid
Jiangxi Hemei Pharmaceutical Co., Ltd

84010-235

Active Ingredient

Salicylic acid 17%

Purpose

Wart remover

Use

For the removal of common warts and plantar warts. Common warts have a rough "cauliflower-like" surface and are easily recognizable. Plantar warts are located only on the soles of the feet, tender to touch, with an interrupted footprint pattern.

Warnings

For external use only.

Do not use

on irritated skin or on any area that is infected or reddened. On moles, birthmarks, warts with hair growing from them, genital warts, or warts on the face or mucous membranes. If you have diabetes or poor blood circulation.

When Using

Avoid contact with eyes. If product gets into the eye, flush with Water for 15 minutes. Non-edible. Cap tightly and store at room temperature, away from heat.

Stop Use

Stop use and ask a doctor if discomfort persists.

Ask Doctor

Stop use and ask a doctor if discomfort persists.

Keep Out Of Reach Of Children

If swallowed, get medical help or contact the Poison Control Center (1-800-222-

1222)right away.

Directions

Wash the affected area. May soak wart in warm water for 5 minutes. Dry area thoroughly. Use a cotton swab to apply a sufficient amount to cover each wart. Let it dry. Repeat this procedure once or twice daily as needed (until wart is removed) for up to 12 weeks.

Other information

Store at room temperature. Avoid excessive heat (37°C/99°F). Skin discoloration may occur during or after use.

Inactive ingredients

Purified Water □ Polyvinyl Alcohol □ Chitosan □ Ethanol □ Glycerin. Laurocapram □ Collodion □ nitrocellulose

PRINCIPAL DISPLAY PANEL



HEALMUSZ WART REMOVER

salicylic acid 17% wart remover liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84010-235
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)		SALICYLIC ACID	17 g in 100 mL
Inactive Ingredients			

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	
LAUROCAPRAM (UNII: 1F3X9DRV9X)	
LYTTA VESICATORIA (UNII: 3Q034RO3BT)	
NITROCELLULOSE (UNII: KYR8BR2X6O)	
POLIGLUSAM (UNII: 82LKS4QV2Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-235-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/17/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	12/17/2025	

Labeler - Jiangxi Hemei Pharmaceutical Co., Ltd (724892056)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-235)

Revised: 12/2025

Jiangxi Hemei Pharmaceutical Co., Ltd