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

84010-087

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 using only. Non-edible. Keep away from heat and open flames, Cap tightly and store at indoor temperature. away from heat. Avoid contact with eyes, if product gets into the eyes, flush with water for 15 minutes.

Do not use

Areas of skin irritation, infection or redness. On warts with hair, birthmarks, or moles.

Genital warts, warts near the eyes or on mucous membranes. If you have diabetes or poor circulation.

When Using

.Apply on intact skin; avoid bleeding or inflamed areas. .Avoid excessive water exposure after application. Discontinue if severe irritation occurs; consult a doctor. .For use on common warts (hands) and plantar warts (feet) only.

Stop Use

Stop use and ask a doctor if discomfort persists.

Ask Doctor

Stop use and ask a doctor if discomfort persists.

Keep Oot Of Reach Of Children

If swallowed, get medical help or contact the Poison Control Center (1-800-222-1222) right away.

Directions

Soak wart in warm water for 5 minutes. Wash and dry thoroughly before application. 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

Water | Polyethylene Glycol | Glycerin | Borneol | Podofilox

PRINCIPAL DISPLAY PANEL

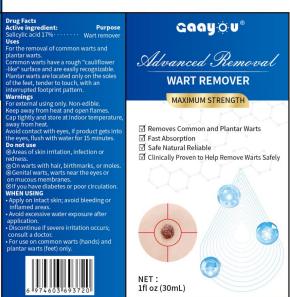




38*50*88mm







GAAYOU WART REMOVER

salicylic acid 17% wart remover liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84010-087
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
BORNEOL (UNII: M89NIB437X)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
GLYCERIN (UNII: PDC6A3C0OX)			
PODOFILOX (UNII: L36H50F353)			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-087- 01	10 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2025	
2	NDC:84010-087- 02	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	03/13/2025	

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

Establishment				
Name	Address	ID/FEI	Business Operations	
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-087)	

Revised: 3/2025 Jiangxi Hemei Pharmaceutical Co., Ltd