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

84010-061

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. Flammable. Keep away from fire and flame.

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 diabetesor poor blood circulation.

When Using

Avoid contact with eyes. If product gets into the eye, flush with Water for 15minutes. 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 Oot Of Reach Of Children

If swallowed, get medical help or contact a 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 bursh 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.

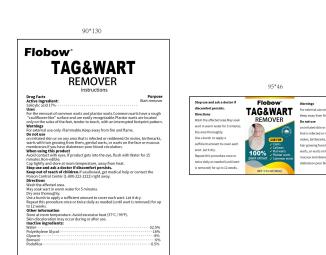
Other information

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





FLOBOW TAG WART REMOVER salicylic acid 17% tag wart remover liquid Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:84010-061 Route of Administration TOPICAL Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	17 g in 100 mL

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

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:84010-061- 01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/24/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	12/24/2024	

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

Establishment				
Name	Address	ID/FEI	Business Operations	
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-061)	

Revised: 12/2024 Jiangxi Hemei Pharmaceutical Co., Ltd