

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

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**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 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**

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## Keep Out 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 brush 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
<b>SALICYLIC ACID</b> (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	17 g in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>BORNEOL</b> (UNII: M89NIB437X)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>PODOFILOX</b> (UNII: L36H50F353)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

**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