# HEALSURE WART REMOVER- salicylic acid 17% wart remover cream Jiangxi Hemei Pharmaceutical Co., Ltd

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84010-086

## **Active Ingredient**

Salicylic acid 17%

### **Purpose**

Wart remover

#### Use

For the removal of common warts, which have a rough, cauliflower-like appearance.

#### **Warnings**

For external use only- Avoid contact with eyes. If the product gets into eyes, rinse with water for 15 minutes. Avoid inhaling vapors.

#### Do not use

Don't use on irritated or infected skin, or reddened areas. If you are diabetic or have poor blood circulation, consult a doctor before use. Don't use on birthmarks, warts with hair growth or moles.

## When Using

If pregnant or breast feeding, consult a health professional before use.

## Stop Use

Stop use and ask a doctor if discomfort persists.

#### **Ask Doctor**

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## Keep Oot Of Reach Of Children

If swallowed, seek immediate medical help or contact a Poison Control Center.

#### **Directions**

- 1.Wash the affected area. (Soaking in warm water for 5 minutes may help.)
- 2. Dry thoroughly.
- 3. Apply a small amount to cover each wart. Let it dry.
- 4. Repeat twice daily as needed, for up to 12 weeks.

#### **Inactive ingredients**

Water, Polyethylene Glycol, Glycerin, Borneol Podofilox, Methylparaben Chlorhexidine

#### PRINCIPAL DISPLAY PANEL



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salicylic acid 17% wart remover cream

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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84010-086
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	17 g in 100 g		

Inactive Ingredients		
Ingredient Name	Strength	
BORNEOL (UNII: M89NIB437X)		
CHLORHEXIDINE (UNII: R4KO0DY52L)		
METHYLPARABEN (UNII: A218C7HI9T)		
WATER (UNII: 059QF0KO0R)		
PODOFILOX (UNII: L36H50F353)		
GLYCERIN (UNII: PDC6A3C0OX)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		

Packaging						
	#	# Item Code Package Description		Marketing Start Date	Marketing End Date	
		NDC:84010-086- 01	20 g in 1 TUBE; Type 0: Not a Combination Product	03/11/2025		

Marketing Information				
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M028	03/11/2025		

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

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-086)	

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