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

84010-083

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

On irritated or infected skin, or reddened areas.

If you are diabetic or have poor blood circulation. 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

Stop use and ask a doctor if discomfort persists.

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

NATUHEAL WART REMOVER

salicylic acid 17% wart remover cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84010-083
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 g

Inactive Ingredients

Ingredient Name	Strength
METHYLPARABEN (UNII: A2I8C7HI9T)	
CHLORHEXIDINE (UNII: R4KO0DY52L)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
BORNEOL (UNII: M89NIB437X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PODOFILOX (UNII: L36H50F353)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-083-01	20 g in 1 TUBE; Type 0: Not a Combination Product	03/07/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	03/07/2025	

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

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-083)

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