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

84010-060

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 medicalhelp 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 Methylparaben Chlorhexidine

PRINCIPAL DISPLAY PANEL

138x40x23mm 350克银卡纳米 主体字凹凸



SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)



17 g in 100 g

SALICYLIC ACID

FLOBOW WART REMOVER

salicylic acid 17% wart remover cream

Salicylic acid 17% wart remove	er cream				
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (So	ource)	NDC:84	010-060
Route of Administration	TOPICAL				
Active Ingredient/Active Moiety					
Ingredient Name			Basis of Strength Streng		Strength

Inactive Ingredients			
Ingredient Name	Strength		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
CHLORHEXIDINE (UNII: R4KO0DY52L)			
GLYCERIN (UNII: PDC6A3C0OX)			
WATER (UNII: 059QF0KO0R)			
BORNEOL (UNII: M89NIB437X)			
METHYLPARABEN (UNII: A2I8C7HI9T)			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1	NDC:84010-060- 01	20 g in 1 TUBE; 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(84010-060)	

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