QUITA CALLOS CORN AND CALLUS REMOVER- salisylic acid 17% liquid Germa Products, LLC

Quita Callos Analgesic Liquid

Active Ingredient

Salicylic Acid 17%

Uses

for the removal of common warts. The common wart is easily recognized by the rough "cauliflower-like" appearance of the surface.

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Purpose

Wart Remover Liquid

Warnings

For external use only. Flammable: Keep away from fire or flame.

Do not use: in or near eyes or mucous membranes. On irritated skin. If prone to allergic reaction to any ingredient in this product. On moles, birthmarks, warts with hair growing from them, genital warts, or warts on the face or mucous membranes. On any area that is infected or reddened. If you are a diabetic, or have poor blood circulation n on large area of the body, on wounds or damaged skin.

When using this product: If product gets in eyes, flush with water for 15 minutes. Do not inhale vapors. Cap bottle tightly and store at room temperature away from heat. If discomfort persists, see your doctor.

Stop use and ask a doctor if; Irritation occurs n condition worsens or does not improve or discomfort persists. If pregnant or breast-feeding. Ask a health professional before use. Keep out of reach of children. In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Warnings

• Keep out of reach of children. In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Other Information

Store between 20^o to 25^oC (68^o to 77^oF). Cap bottle tightly. Use product in well ventilated area. Avoid inhaling vapors. Retain packaging for full labeling.

Directions

- Wash affected area and dry thoroughly.
- Apply one drop at a time with applicator to sufficiently cover each wart.
- Let dry.
- Repeat this procedure once or twice daily as needed for up to 2 weeks.

Inactive Ingredients

Flexible Collodion (Alcohol, Camphor, Mineral Oil), and Isopropyl Alcohol.





Quita Callos

QUITA CALLOS CORN AND CALLUS REMOVER salisylic acid 17% liquid								
Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:73635-5811				
Route of Administration	TOPICAL							
Active Ingredient/Active Moiety								
Ingredient Name			Basis of Strength		Strength			
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414			SALICYLIC ACID		170 mg in 1 mL			
Inactive Ingredients								
Ingredient Name					Strength			
ISOPROPYL ALCOHOL (UNII: ND2M416302)								
ALCOHOL (UNII: 3K9958V90M)								

CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)								
CASTOR OIL (UNII: D5340Y2I9G)								
Packaging								
#	ltem Code		Package Description		Marketing Star Date	t Marketing End Date		
1	NDC:73635- 5811-5	1 in	in 1 BOX		03/29/2019			
1			mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not bination Product	а				
Marketing Information								
Marketing Category			Application Number or Monograph Citation	М	arketing Start Date	Marketing End Date		
OTC Monograph Drug		Drug	M030	03/2	9/2019			

Labeler - Germa Products, LLC (116626935)

Revised: 10/2023

Germa Products, LLC