SCHOLLS WELLNESS COMPANY LLC LIQUID CORN AND CALLUS REMOVERsalicylic acid liquid Scholls Wellness Company LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Liquid Corn & Callus Remover

Drug Facts

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Active Ingredient

Active Ingredient Purpose

Salicylic Acid 17%w/w......Corn and callus remover

Purpose

Corn and callus remover

Uses

Uses

for the removal of corns and calluses relieves pain by removing corns and calluses

Warnings

Warnings

For external use only

Flammable: Keep away from fire or flames.

Do not use

Do not use

if you are a diabetic if you have poor blood circulation on irritated skin or any area that is infected or reddened

When using this product

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

Stop use and ask a doctor

Stop use and ask a doctor if discomfort lasts

Keep out of reach of children. If swallowed,

get medical help or contact a Poison Control Center right away.

Directions

Directions

wash affected area and dry thoroughly apply one drop at a time with applicator to sufficiently cover each corn/callus let dry

repeat procedure once or twice daily as needed for up to 14 days (until corn/callus is removed)

may soak corn/callus in warm water for 5 minutes to assist in removal protective pads may be used to cushion corn/callus during treatment

Other Information

Other information

store between 20°C to 25°C (68° to 77°F)

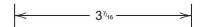
keep tightly closed

retain card for full labeling

Inactive ingredients castor oil, ethyl lactate, flexible collodion, polybutene. (Contains alcohol 18% v/v and ether 53% v/v from flexible collodion)

Questions? 1-866-360-3226

Liquid Corn & Callus Remover Salicylic Acid



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SCHOLLS WELLNESS COMPANY LLC LIQUID CORN AND CALLUS REMOVER

salicylic acid liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73469-0464
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	1.7 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
ETHER (UNII: 0F5N573A2Y)			
ALCOHOL (UNII: 3K9958V90M)			
ETHYL LACTATE (UNII: F3P750VW8I)			

Product Characteristics			
Color	yellow	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73469- 0464-1	1 in 1 BLISTER PACK	01/01/2021	
1		1 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358F	01/01/2021	

Labeler - Scholls Wellness Company LLC (117174744)

Registrant - Scapa Tapes North America LLC (079995435)

Revised: 1/2022 Scholls Wellness Company LLC