# SALICYLIC ACID- callus removers patch Premier Brands of America Inc.

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#### **Premier Brands Callus Removers**

## Active ingredient

Salicylic acid 40%

## **Purpose**

Callus remover

#### Use

- for the removal of calluses
- relieves pain by removing calluses

# Warnings

For external use only.

#### Do not use

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

**If discomfort persists** see your doctor or podiatrist

# Keep out of reach of children.

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

#### **Directions**

- wash affected area and dry thoroughly
- if necessary, cut medicated patch to fit callus
- apply adhesive side down of medicated patch onto callus
- cover medicated patch with pad
- after 48 hours, remove medicated patch
- repeat procedure every 48 hours as needed for up to 14 days (until callus is removed)
- may soak callus in warm water for 5 minutes to assist in removal
- continued wearing of pad (without patch) will help prevent recurrence of calluses

#### Other information

store between 15°C to 30°C (59°F to 86°F)

## Inactive ingredients

acrylic adhesive, acrylic polymer, polyethylene, polyvinyl alcohol

# Questions?

call 1-866-964-0939

# Principal Display Panel Premier CALLUS REMOVERS

Salicylic Acid
Effective callus removal treatment
Protects against pressure & friction
MEDICATED
6 PADS/4 MEDICATED PATCHES



#### SALICYLIC ACID

callus removers patch

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56104-013	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	40 mg in 4

Inactive Ingredients		
Ingredient Name	Strength	
POLYVINYL ALCOHOL (UNII: 532B59J990)		
VINYL ACETATE (UNII: L9MK238N77)		
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)		

Pa	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:56104-013- 04	4 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2013	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	01/01/2013	

# **Labeler - Premier Brands of America Inc. (117557458)**

Revised: 2/2024 Premier Brands of America Inc.