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

Premier Brands Extra Thick Callus Remover

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, or 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

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

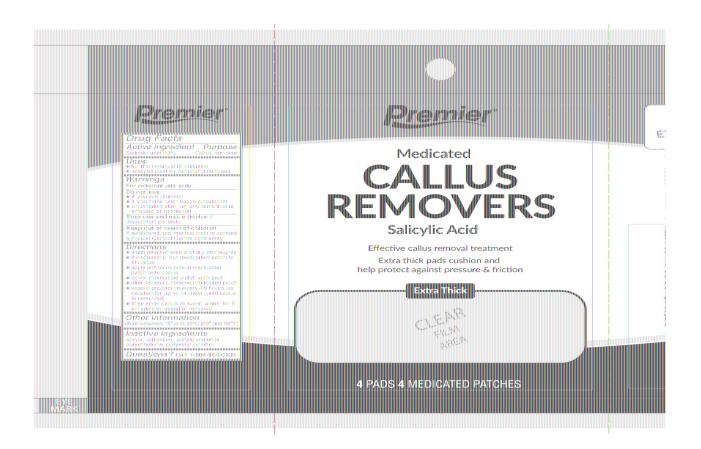
Extra Thick

Callus Removers

Salicylic acid

NEW!

- Medicated Callus removal treatment
- Protects against pressure & pain
- 4 Pads
- 4 Medicated Patches



SALICYLIC ACID

extra thick callus removers patch

D	Information
Product	Intormation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:56104-014

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)

SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)

Inactive Ingredients

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

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:56104-014- 04	4 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2013		
Marketing Information					

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.