

SALICYLIC ACID- medicated callus removers patch
CVS Pharmacy

CVS Medicated Callus Removers

Active ingredient

Salicylic acid 40%

Purpose

Callus remover

Uses

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

Warnings

For external use only.

Do not use

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

Stop use and ask a doctor if discomfort persists

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

CVS Health

Medicated

Callus

Removers

Salicylic Acid 40%

- Effective callus

removal treatment

- Cushioning pads help

protect against pressure

and friction

4 MEDICATED PATCHES

6 PROTECTIVE PADS

CRIMP SEAL AREA (NO TEXT)

Drug Facts
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Regular

NDC 59779-569-10

Medicated
Callus Removers

SALICYLIC ACID 40%

- Effective callus removal treatment
- Cushioning pads help protect against pressure and friction



4 MEDICATED PATCHES
 6 PROTECTIVE PADS

52-005CV-01

LOT #
 EXP DATE:



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CRIMP SEAL AREA (no text but must have color)

SALICYLIC ACID

medicated callus removers patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-569
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	40 mg in 4

Inactive Ingredients

Ingredient Name	Strength
POLYVINYL ALCOHOL (UNII: 532B59J990)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-569-10	4 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	01/01/2014	

Labeler - CVS Pharmacy (062312574)

Revised: 12/2025

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