# SALICYLIC ACID- medicated callus removers patch CVS Pharmacy

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#### **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
- 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 thoroughlyif 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

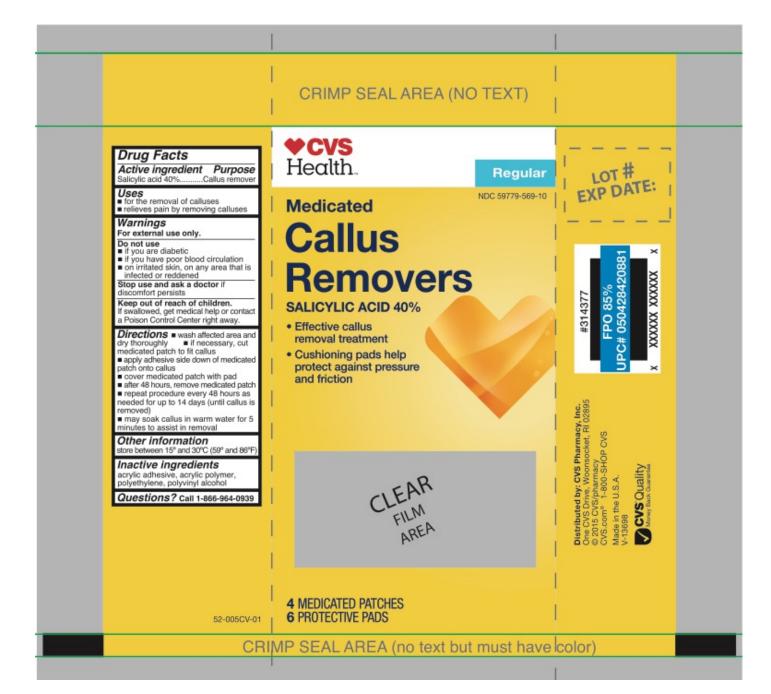
## **Medicated Callus Removers**

Salicylic Acid 40%

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

## **4 Medicated Patches**

**6 Protective Pads** 



## **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	<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)		

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

Marketing In	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: 1/2024 CVS Pharmacy