# SALICYLIC ACID- medicated corn removers patch Chain Drug Consortium, LLC

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#### **Premier Value Corn Removers**

#### Active ingredient

Salicylic acid

#### Purpose

Corn Remover

#### Uses

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

#### Warnings

For external use only.

#### Do not use

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

## Ask your doctor

**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 area thoroughly
- if necessary, cut medicated patch to fit corn
- apply adhesive side down of medicated patch onto corn
- cover medicated patch with pad
- after 48 hours, remove medicated patch
- repeat procedure every 48 hours as needed for up to 14 days (until corn is removed)
- may soak corn 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 Value** 

## **Medicated Corn Removers**

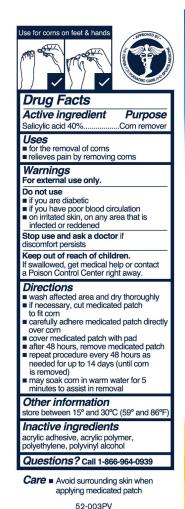
Salicylic Acid 40%

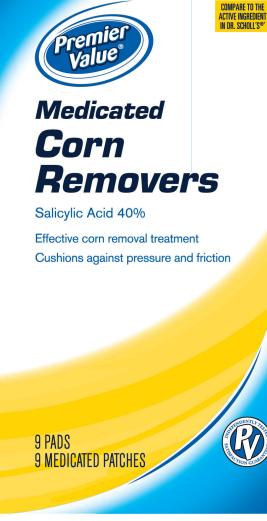
Effective corn removal treatment

Cushions against pressure and friction

9 PADS

9 MEDICATED PATCHES







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#### SALICYLIC ACID

medicated corn removers patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-606
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
	Ingredient Name	<b>Basis of Strength</b>	Strength
SALICYLIC A	ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	40 mg in 9

Inactive Ingredients		
Ingred	ient Name	Strength

POLYVINYL ALCOHOL (UNII: 532B59J990)	
VINYL ACETATE (UNII: L9MK238N77)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:68016-606- 00	9 in 1 PACKAGE; Type 0: Not a Combination Product	05/23/2003	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	05/23/2003	

## Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 12/2024 Chain Drug Consortium, LLC