# SALICYLIC ACID- corn removers patch Walgreen Company

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## **Walgreens Medicated Corn Removers**

### Active ingredient

Salicylic acid 40%

## 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
- if you 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 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**

Walgreens

Corn

Removers

**SALICYLIC ACID 40%** 

MAXIMUM STRENGTH

#### **MEDICATED**

- Safe & effective corn removal treatment
- Cushioned pads protect against friction & reduced pressure

#### 9 MEDICATED PATCHES

9 PADS



## **SALICYLIC ACID**

corn removers patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9757
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 9

Inactive Ingredients		
Ingredient 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:0363-9757- 09	9 in 1 PACKAGE; Type 0: Not a Combination Product	07/22/2008	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	07/22/2008	

## Labeler - Walgreen Company (008965063)

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