

WALGREENS CORN REMOVER- salicylic acid strip

WALGREENS Co

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Walgreens Corn Remover Strips

Drug Facts

Active Ingredient

Salicylic acid 40%

Purpose

Corn remover

Uses

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

Warnings

For external use only

Use only as directed

Do Not Use

- if you are diabetic
- on irritated, broken or infected skin
- if you have poor blood circulation

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 immediately.

Directions

- Wash affected area. Soak in warm water for 5 minutes if desired. Dry thoroughly.
- Remove bandage from backing
- Apply with medicated area directly over corn
- After 48 hours, remove medicated bandage
- Repeat procedure every 48 hours as needed (until corn is removed) for up to 14 days

Other Information

Store between 59° and 86°F (15° and 30°C)

Inactive Ingredients

Polyvinyl Alcohol, Acrylic Copolymer

Questions or Comments?

CALL: 1-800-925-4733

DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015

PRINCIPAL DISPLAY PANEL - 6 Strip Box

Well at

Walgreens

WALGREENS PHARMACIST RECOMMENDED ‡

NEW

MAXIMUM STRENGTH • ONE-STEP

Corn Remover

Strips

Salicylic Acid 40 %

- Removes corns in as few as two treatments*
- Water resistant
- Medicated pad
- Stays in place

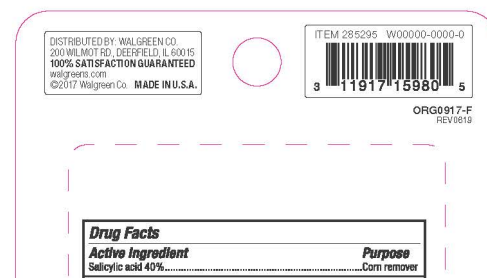
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MEDICATED

STRIPS

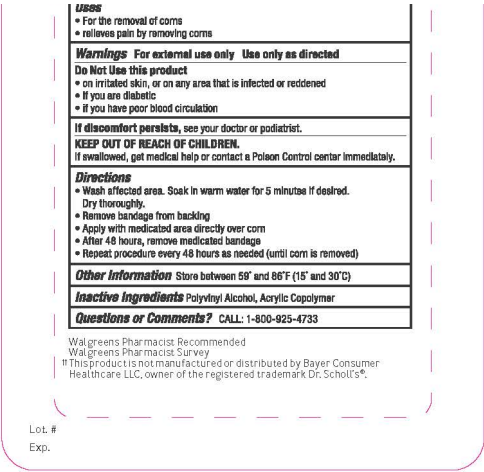
Compare to Dr. Scholl's® OneStep

Corn Removers active ingredient ‡‡





FRONT



WALGREENS CORN REMOVER

salicylic acid strip

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-3063
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	4.42 mg

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-3063-02	2 in 1 BOX	01/01/2021	
1	NDC:0363-3063-01	3 in 1 BAG; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358F	06/01/2013	

Labeler - WALGREENS Co (008965063)

Registrant - PEDIFIX, INC. (122271935)

Establishment			
Name	Address	ID/FEI	Business Operations
PEDIFIX		122271935	pack(0363-3063) , label(0363-3063)

Establishment			
Name	Address	ID/FEI	Business Operations
AKRON		186569323	manufacture(0363-3063)

Revised: 1/2021

WALGREENS Co