

**WALGREENS MAXIMUM STRENGTH ONE-STEP CORN REMOVER- salicylic acid strip
Walgreen Co.**

Walgreens ONE-STEP Corn Remover Strips

Drug Facts

Active ingredient (in each dosage unit)

Salicylic acid 40% in a plaster vehicle

Purposes

Corn and calluses removal

Uses

- For the removal of corns and calluses
- Relieves pain by removing corns and calluses
- Adhesive bandage holds the medicated pad in position

Warnings

For external use only

Do not use

- If you are a diabetic
- If you have poor blood circulation
- On irritated skin or any area that is infected or reddened

Stop use and ask a doctor if

Discomfort lasts

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
- Apply medicated bandage after 48 hours, remove medicated bandage
- 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 0° to 40°C (32° to 104°F)

Inactive ingredients

Pressure sensitive adhesive fabric

Questions?

1-800-723-2465

Water Resistant

- **Medicated pad**
- **Stays in place**

Pharmoplast S.A.E.

Amria, Free zone 23512,

Alexandria, Egypt.

www.pharmoplast-online.com

†Our pharmacists recommend the Walgreens brand.

DISTRIBUTED BY: WALGREEN CO.

DEERFIELD, IL 60015

100% SATISFACTION GUARANTEED

walgreens.com

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Packaging

NDC 0363-1972-06

Walgreens



ONE-STEP Corn Remover Strips

SALICYLIC ACID 40%

Maximum Strength

Medicated

Water Resistant

- Medicated pad
- Stays in place

6
MEDICATED
STRIPS



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www.pharmaplast-eg.com

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LOT



WALGREENS MAXIMUM STRENGTH ONE-STEP CORN REMOVER

salicylic acid strip

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-1972
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	1.35 mg

Inactive Ingredients

Ingredient Name	Strength
POLYACRYLATE-13 (UNII: FS2D4T67EA)	

Packaging

Medication Start Medication End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-1972-06	6 in 1 CARTON; Type 0: Not a Combination Product	10/18/2023	09/30/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	10/18/2023	09/30/2026

Labeler - Walgreen Co. (008965063)

Revised: 12/2025

Walgreen Co.