CLEAR GEL CORN REMOVER- salicylic acid patch Chain Drug Consortium, LLC

Premier Value Clear Gel Wart Removers

Active Ingredient

Salicylic acid 40%

Purpose

Wart Remvoer

Uses

• for the removal of common warts. the common wart is easily recognized by the rough 'cauliflower-like' appearance of the surface.

Warnings

For external use only.

Do not use

- if you are diabetic
- have poor blood circulation
- on irritated skin, on any area that is infected or reddened
- on genital warts and warts on the face
- on moles, birthmarks and warts with hair growing from them
- on mucous membranes

Stop use and ask doctor

if discomfort persist

Keep out of reach of children.

If, swallowed, get medical help or cotnact a poison Control Center right away.

Directions

- wash affected area, may soak wart in warm water for 5 minutes
- dry area thoroughly
- if necessary, cut medicated disc to fit wart
- carefully apply adhesive side down of disc onto the wart
- cover disc with cushion to conceal area
- repeart this procedure every 48 hours as needed (until wart is removed) for up to 12 weeks

Other Information

store between 59°F to 86°F (15°F to 30°F)

Inactive Ingredients

acrylic adhesive, acrylic polymer, polyethylene, polyvinyl alcohol

Questions

Call 1-866-964-0939

Principal Display Panel

Premier Value

Medicated

Clear Gel Wart Removers

SALICYLIC ACID 40%

Maximum strength

- Effective wart removal treatment
- Clear gel cushions affected area
- Thin, flexible bandage conceals and protects

9 CUSHIONS

12 MEDICATED DISCS



CLEAR GEL CORN REMOVER

salicylic acid patch

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Prod	IUCT	Information	

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68016-444

Route of Administration TOPICAL

Active Ingredient/Active Moiety

ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	40 mg in 6

Inactive Ingredients

Ingredient Name

Strength

HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
VINYL ACETATE (UNII: L9MK238N77)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59 990)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-444- 12	6 in 1 PACKAGE; Type 0: Not a Combination Product	01/19/2023	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M030	01/01/2018		

Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 2/2024 Chain Drug Consortium, LLC