

SALICYLIC ACID- plantar wart remover patch
Chain Drug Consortium, LLC

Premier Value Medicated Plantar Wart Remover

Active ingredient

Salicylic acid 40%w/w

Purpose

Plantar wart remover

Uses

- for the removal of plantar warts on the bottom of the foot. The plantar wart is recognized by its location only on the bottom of the foot, its tenderness, and the interruption of the footprint pattern.

Warnings

For external use only.

Do not use

- if you are diabetic or have poor blood circulation, except under the advise and supervision of a doctor or podiatrist
- on irritated, infected or reddened skin
- on genital warts and warts on the face
- on moles, birthmarks and warts with hair growing from them
- on mucous membranes

Keep this and all drugs out of the reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Ask a doctor

If discomfort persists see your doctor or podiatrist.

Directions

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

Other information

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

Inactive ingredients

acrylic adhesive, acrylic polymer, polyethylene, polyvinyl alcohol

Questions?

call 1-866-964-0939

Principal Display Panel**Premier Value****Medicated PLANTAR WART REMOVER**

Salicylic acid 40%

Conceals and protects while removing warts

No mess treatment

Safe and effective

FOR FEET

24 MEDICATED PATCHES

24 CONCEALING PADS

THIS AREA MUST HAVE NO TEXT



SALICYLIC ACID

plantar wart remover patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-618
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 24

Inactive Ingredients

Ingredient Name	Strength
POLYVINYL ALCOHOL (UNII: 532B59J990)	

VINYL ACETATE (UNII: L9MK238N77)

HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-618-00	24 in 1 PACKAGE; Type 0: Not a Combination Product	02/19/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	02/19/2009	

Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 12/2024

Chain Drug Consortium, LLC