SALICYLIC ACID- medicated plantar wart remover patch
Chain Drug Marketing Association

Quality Choice 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.

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

QC Quality Choice
Medicated

PLANTAR WART

REMOVER

Salicylic Acid

For Feet:

Cushions, conceals and protects while removing warts

Maxium strength formula without the mess of liquids

24 Medicated Patches

24 Cushioning Pads

SALICYLIC ACID

medicated plantar wart remover patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-042
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	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:63868-042-24	24 in 1 BOX; Type 0: Not a Combination Product	12/18/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	12/18/2017	

Labeler - Chain Drug Marketing Association (011920774)

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