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

Quality Choice Medicated Plantar Wart Remover

Active ingredient

Salicylic acid 40%

Purpose

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

- 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
- if you are diabetic
- if you have poor blood circulation

Stop use and ask a doctor if

discomfort persists

Keep out of reach of children.

If swallowed, get medical help or contact 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 patch to fit wart
- carefully adhere medicated patch directly over the wart
- cover medicated patch with pad to conceal area
- repeat procedure every 48 hours as needed (until wart is removed) for up to 12 weeks

Other information

- store between 15° and 30°C (59° and 86°F)
- avoid surrounding skin when applying medicated product

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 40% / Plantar Wart Remover

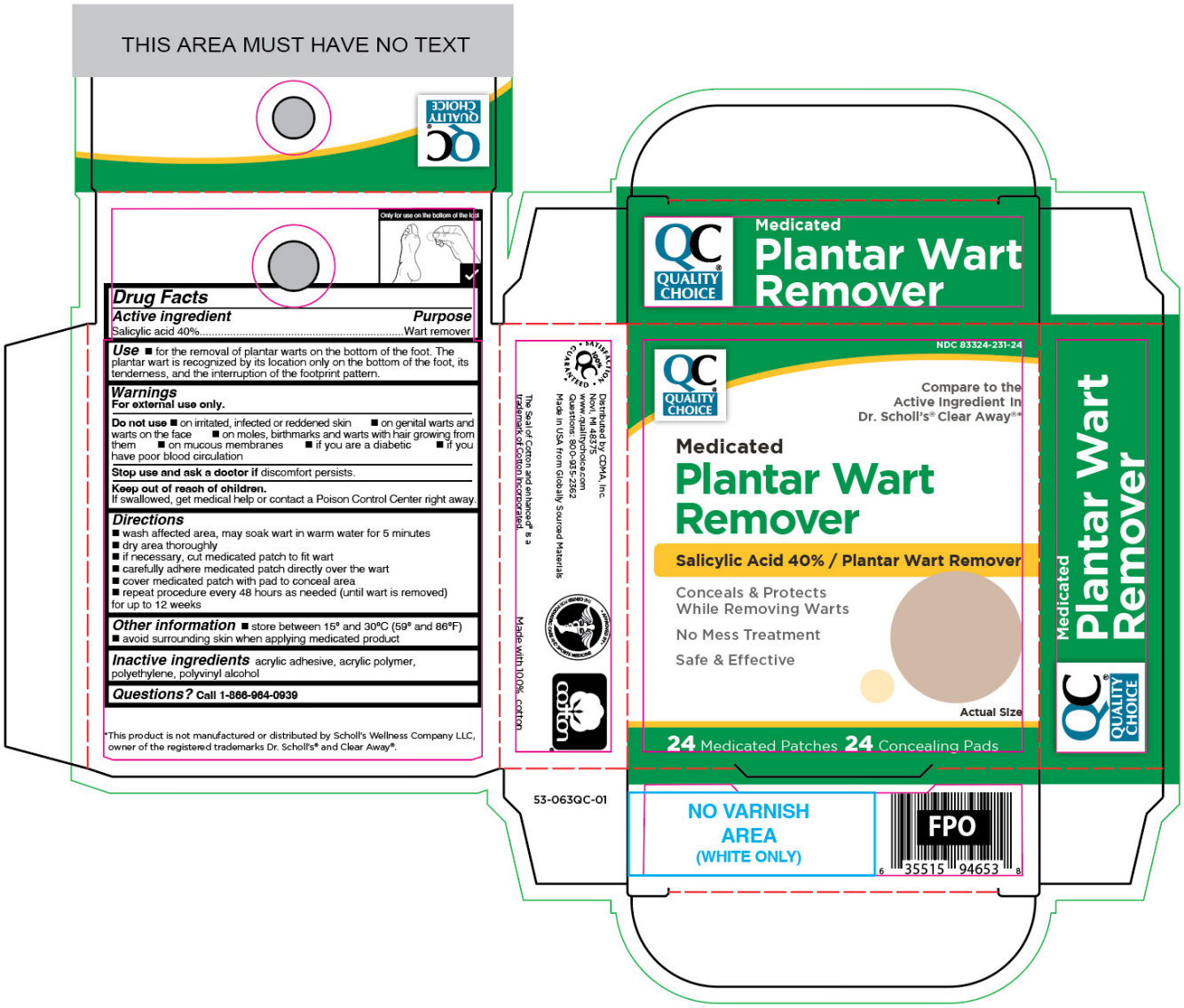
Conceals & Protects

While Removing Warts

No Mess Treatment

Safe & Effective

24 Medicated Patches 24 Concealing Pads



SALICYLIC ACID

medicated plantar wart remover patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-231
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	400 mg in 1 g

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:83324-231-24	24 in 1 BOX	04/29/2025	
1		1 g in 1 PATCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	04/29/2025	

Labeler - Chain Drug Marketing Association (011920774)

Revised: 5/2025 Chain Drug Marketing Association