

SALICYLIC ACID- medicated callus removers patch
Chain Drug Marketing Association

Quality Choice Medicated Callus Removers

Active ingredient

Salicylic acid 40%

Purpose

Callus remover

Uses

- for the removal of calluses
- relieves pain by removing calluses

Warnings

For external use only.

Do not use

- if you are a diabetic
- have poor blood circulation
- on irritated skin, on any area that is infected or reddened

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 and dry thoroughly if necessary, cut medicated patch to fit callus
- apply adhesive side down of medicated patch onto callus
- cover medicated patch with pad
- after 48 hours, remove medicated patch
- repeat procedure every 48 hours as needed for up to 14 days (until callus is removed)
- may soak callus in warm water for 5 minutes to assist in removal

Other information

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

Inactive ingredients

acrylic adhesive, acrylic polymer, polyethylene, polyvinyl alcohol

Questions?

call 1-866-964-0939

Principal Display Panel

QC Quality Choice

Medicated

Callus

Removers

Treatment & Protection

Salicylic Acid

Callus Removal Treatment

Protect Against Pressure & Pain

6 Pads

4 Medicated Patches

TOP		CLEAR AREA	
		CRIMP SEAL AREA (NO TEXT)	
<p>Medicated</p> <p>Callus Removers</p> <p>Treatment & Protection</p> <p>Salicylic Acid</p> <hr/> <p>Drug Facts</p> <p>Active ingredient Purpose</p> <p>Salicylic acid 40%.....Callus remover</p> <p>Uses</p> <ul style="list-style-type: none"> ■ for the removal of calluses ■ relieves pain by removing calluses <p>Warnings</p> <p>For external use only.</p> <p>Do not use</p> <ul style="list-style-type: none"> ■ if you are diabetic ■ if you have poor blood circulation ■ on irritated skin or any area that is infected or reddened <p>Stop use and ask a doctor if discomfort persists</p> <p>Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.</p> <p>Directions</p> <ul style="list-style-type: none"> ■ wash affected area and dry thoroughly ■ if necessary, cut medicated patch to fit callus ■ apply adhesive side down of medicated patch onto callus ■ cover medicated patch with pad ■ after 48 hours, remove medicated patch ■ repeat procedure every 48 hours as needed for up to 14 days (until callus is removed) ■ may soak callus in warm water for 5 minutes to assist in removal <p>Other information</p> <p>store between 15° and 30°C (59° and 86°F)</p> <p>Inactive ingredients</p> <p>acrylic adhesive, acrylic polymer, polyethylene, polyvinyl alcohol</p> <p>Questions? Call 1-248-449-9300</p>	<p>QC QUALITY CHOICE</p> <p>*Compare to DR. SCHOLL'S®</p> <p>Medicated</p> <p>Callus Removers</p> <p>Treatment & Protection</p> <p>Salicylic Acid</p> <p>Callus Removal Treatment Protects Against Pressure & Pain</p> <p>6 Pads 4 Medicated Patches</p> <p>made in the USA</p>	<p>LOT # EXP DATE:</p> <p>6 35515 90598 6</p> <p>F.P.O.</p> <p>Made in USA LFC/ROC-2</p> <p>SALES GUARANTEED</p> <p>© Distributed by C.D.M. A., Inc. 43157 W. Nine Mile Novi, MI 48376-0995 www.qualitychoice.com Questions: 248-449-9300</p> <p>*This product is not manufactured or distributed by MSD Consumer Care, Inc., owner of the registered trademark Dr. Scholl's®.</p>	
		<p>Medicated</p> <p>Callus Removers</p> <p>Treatment & Protection</p> <p>Salicylic Acid</p> <p>Callus Removal Treatment Protects Against Pressure & Pain</p> <p>6 Pads 4 Medicated Patches</p> <p>made in the USA</p>	
BOTTOM		CLEAR AREA	
EYE MARK	CRIMP SEAL AREA (no text but must have color)		EYE MARK

SALICYLIC ACID

medicated callus removers patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-041
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 4

Inactive Ingredients

Ingredient Name	Strength
VINYL ACETATE (UNII: L9MK238N77)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-041-04	4 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	01/01/2014	

Labeler - Chain Drug Marketing Association (011920774)

Revised: 2/2024

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