

## **SALICYLIC ACID- medicated callus removers extra thick patch**

### **Target Corporation**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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### **Target Extra Thick Callus Removers**

#### ***Active ingredient***

Salicylic acid 40%

#### ***Purpose***

Callus remover

#### ***Use***

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

#### ***Warnings***

**For external use only.**

#### **Do not use**

- if you are a diabetic
- if you 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 area 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 corn is removed)
- may soak corn 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**

medicated

callus removers

salicylic acid 40%

effective callus removal treatment

extra thick pads cushion & help

protect against pressure & friction

4 PADS/ 4 MEDICATED PATCHES

CLEAR AREA

CRIMP SEAL AREA (NO TEXT)

**Drug Facts**

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Salicylic acid 40%.....Callus remover

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**Directions**

- wash affected area and dry thoroughly
- if necessary, cut medicated patch to fit callus
- apply adhesive side 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° and 30°C (59° and 86°F)

**Inactive ingredients**  
acrylic adhesive, acrylic polymer, polyethylene, polyvinyl alcohol

**Questions? Call 1-800-910-6874**

52-072TG


NDC 11673-631-04

**Compare to Dr. Scholl's® Callus Removers\***

**medicated callus removers**  
salicylic acid 40%

extra-thick cushioned pads  
cushions help protect against pressure and friction  
super adhesive holds cushion firmly in place

CLEAR WINDOW AREA



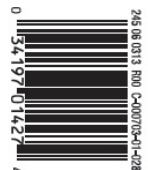
4

EACH

4 CUSHIONS, 4 MEDICATED DISCS

LOT #

EXP DATE:



0 34197 01427 4


245 08 0313 R00 C-000703-01-028

100% satisfaction guaranteed or your money back.

We welcome any questions you may have. Please call our toll-free number for comments or 1-800-910-6874.

Distributed by Target Corporation  
Minneapolis, MN 55403  
Made in U.S.A. with U.S. and imported parts  
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\*This product is not manufactured or distributed by Bayer HealthCare LLC, owner of the registered trademark "up&up".



PLASTIC BAG

EYE MARK

CRIMP SEAL AREA (no text but must have color)

EYE MARK

SALICYLIC ACID			
medicated callus removers extra thick patch			
Product Information			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11673-631
<b>Route of Administration</b>	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		
POLYVINYL ALCOHOL (UNII: 532B59J990)			
VINYL ACETATE (UNII: L9MK238N77)			
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)			
Packaging			
	Marketing Start	Marketing End	

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-631-04	4 in 1 PACKAGE; Type 0: Not a Combination Product	12/01/2017	
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part358F	12/01/2017		

**Labeler** - Target Corporation (006961700)

Revised: 7/2021

Target Corporation