

SALICYLIC ACID- medicated callus removers extra thick patch
Topco Associates LLC

Topcare Medicated 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

TopCare

health

Extra Thick • Medicated

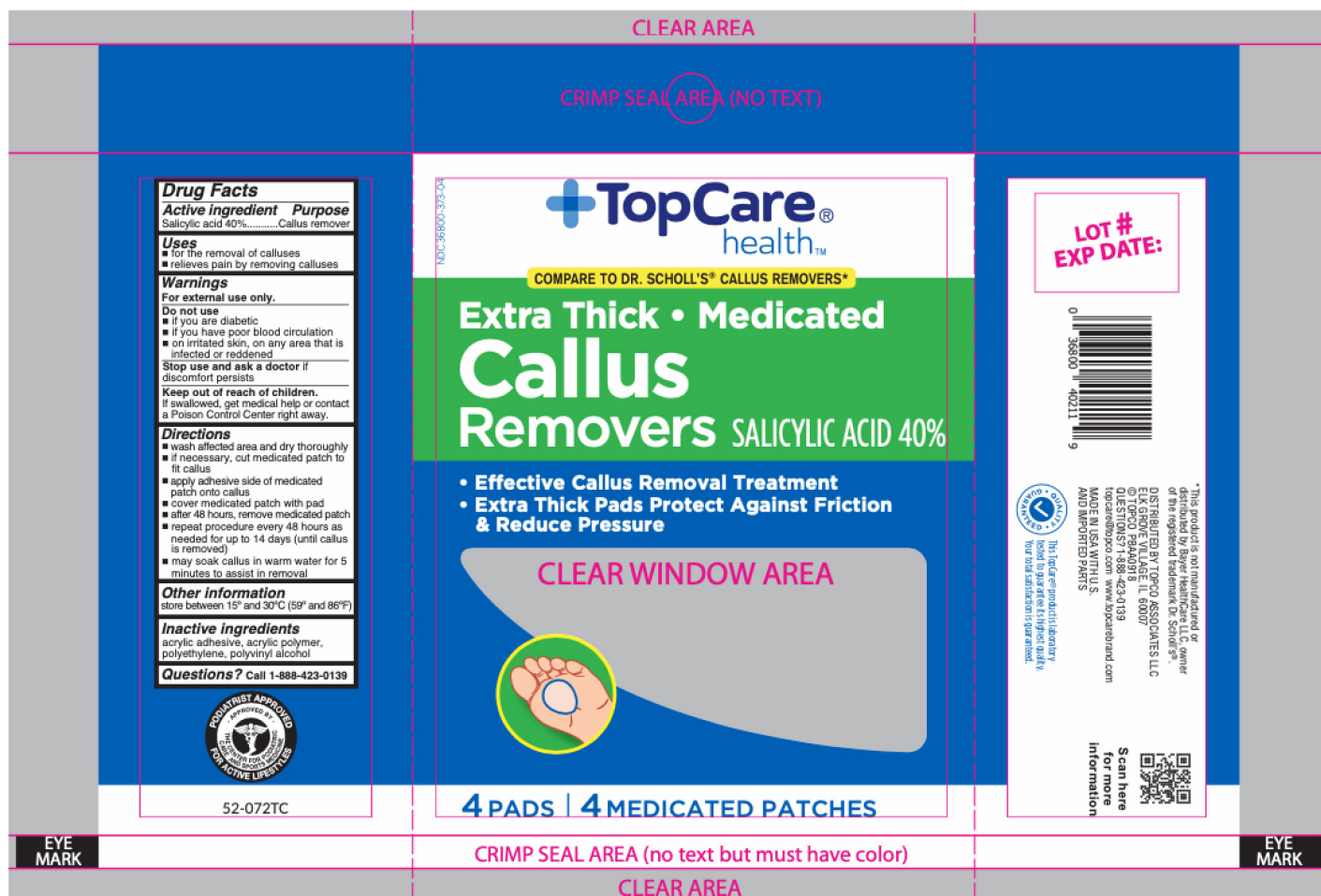
Callus

Removers SALICYLIC ACID 40%

- Effective Callus Removal Treatment
- Extra Thick Pads Protect Against Friction

& Reduce Pressure

4 PADS | 4 MEDICATED PATCHES



SALICYLIC ACID

medicated callus removers extra thick patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-373
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
POLYVINYL ALCOHOL (UNII: 532B59J990)	
VINYL ACETATE (UNII: L9MK238N77)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	

Packaging

	Marketing Start	Marketing End
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-373-04	4 in 1 PACKAGE; Type 0: Not a Combination Product	12/06/2017	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M030	12/06/2017	

Labeler - Topco Associates LLC (006935977)