EXTRA THICK CALLUS REMOVERS- extra thick callus removers with salicylic acid patch
Wal-Mart Stores, Inc.

Equate Extra Thick 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 diabetic
- if you have poor blood circulation
- on irritated skin or any area that is infected or reddened

If discomfort persists

see your doctor or podiatrist

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 patches to fit callus
- apply the 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 59° and 86°F (15° and 30°C)

Inactive ingredients

acrylic adhesive, acrylic polymer, polyethylene, polyvinyl alcohol

Questions?

call 1-888-287-1915

Principal Display Panel

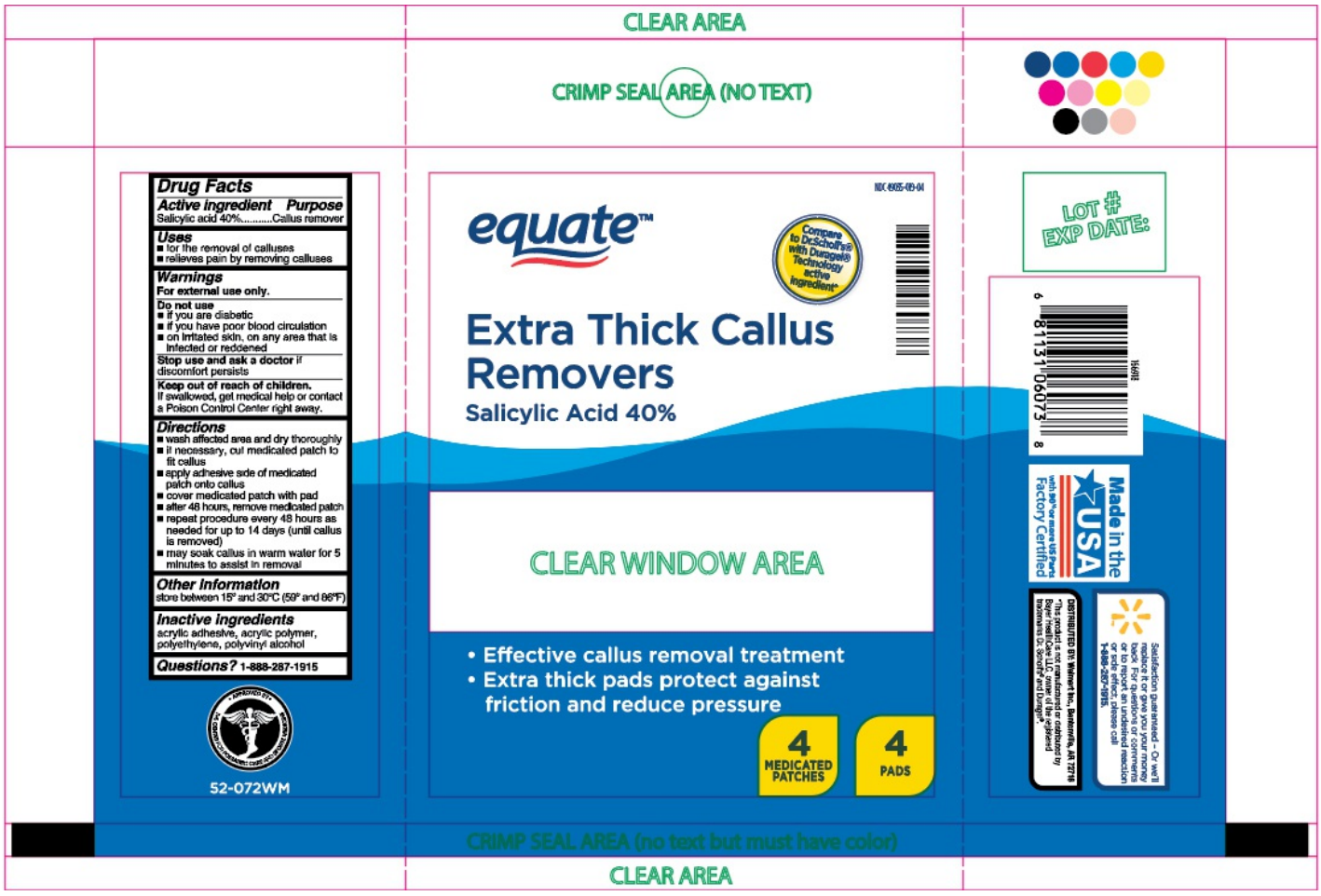
Equate

Extra Thick Callus Removers

Salicylic Acid 40%

- Effective callus removal treatment
- Extra thick pads protect against friction and reduce pressure

4 Medicated Patches/ 4 Pads



EXTRA THICK CALLUS REMOVERS

extra thick callus removers with salicylic acid patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-019
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 100 mg

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:49035-019-04	160 mg in 1 PACKAGE; Type 0: Not a Combination Product	08/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	08/01/2013	

Labeler - Wal-Mart Stores, Inc. (051957769)

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