SALICYLIC ACID- medicated callus removers extra thick patch
HEB

Hill Country Essentials 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

HILL COUNTRY ESSENTIALS

Extra Thick

Callus Removers

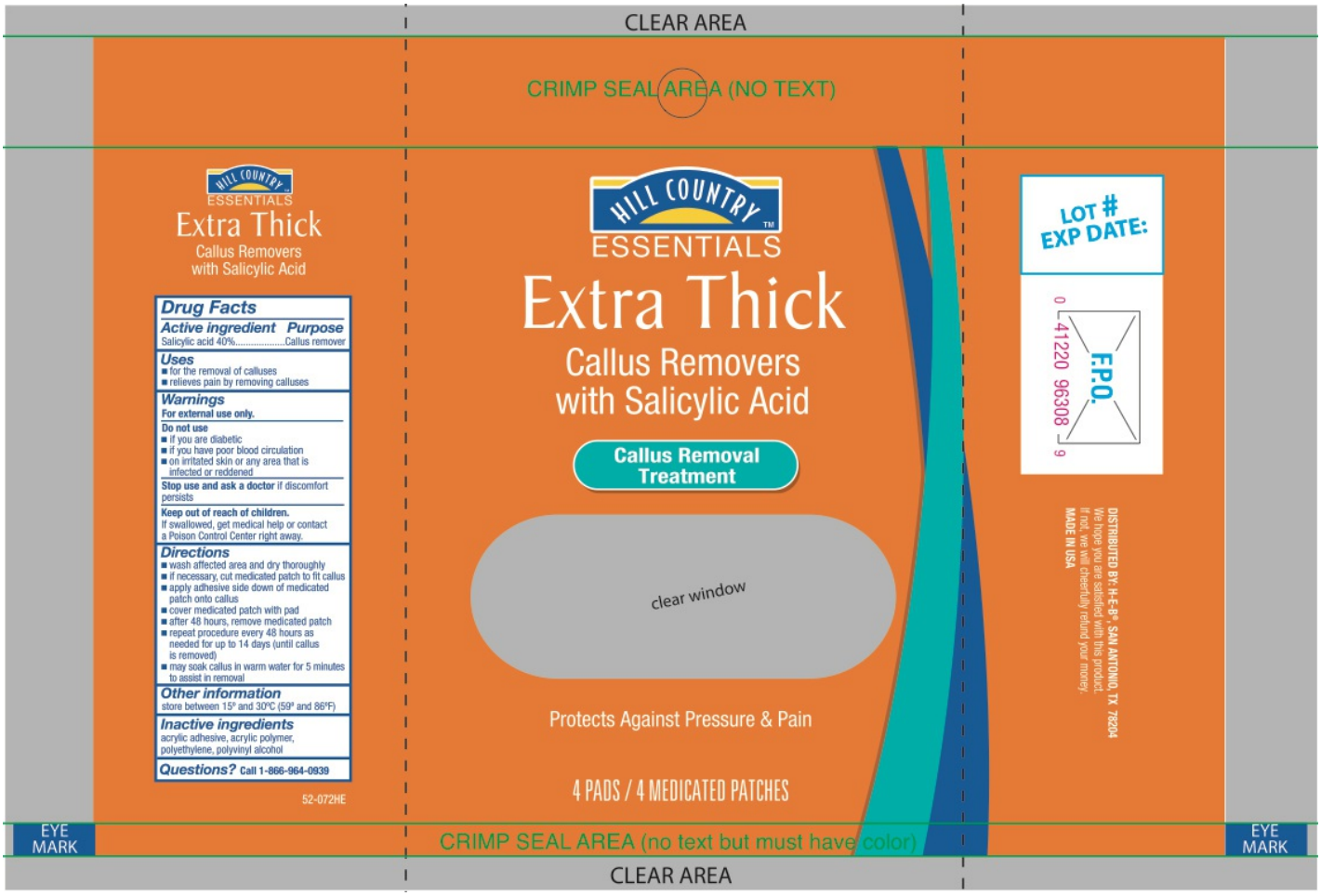
with Salicylic Acid

Callus Removal

Treatment

Protects Against Pressure & Pain

4 PADS / 4 MEDICATED PATCHES



SALICYLIC ACID

medicated callus removers extra thick patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-522
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)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:37808-522-04	4 in 1 PACKAGE; Type 0: Not a Combination Product	12/22/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	12/22/2017	

Labeler - HEB (007924756)

Revised: 1/2024 HEB