WALGREENS MEDICATED CALLUS REMOVERS- salicylic acid disc WALGREENS Co.

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.

Walgreens Medicated Gel Callus Removers

Drug Facts

Active Ingredient

Salicylic acid 40%

Purpose

Callus remover

Uses

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

Warnings

For external use only Use only as directed

Do Not Use

- if you are diabetic
- on irritated, broken or infected skin
- if you have poor blood circulation

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 immediately.

Directions

- Wash affected area. Soak in warm water for 5 minutes if desired. Dry thoroughly
- Apply medicated disk directly over callus
- Cover medicated disk with gel pad
- After 48 hours, remove medicated disk
- Repeat procedure every 48 hours as needed (until callus is removed) for up to 14 days

Other Information

Store between 59° and 86°F (15° and 30°C)

Inactive Ingredients

Polyvinyl Alcohol, Acrylic Copolymer

Questions or Comments?

· · · · · ·

CALL: 1-800-925-4733

DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015

PRINCIPAL DISPLAY PANEL - 4 Disk Blister Pack

Well at **Walgreens**

WALGREENS PHARMACIST RECOMMENDED [‡]

NEW

MEDICATED

Gel Callus Removers

Salicylic Acid 40%

- Callus removal treatment
- Cushions against pressure & pain
- Self-sticking, washable & reusable
- Effective for weeks

4

MEDICATED DISKS | 4 CUSHIONS



WALGREENS MEDICATED CALLUS REMOVERS

salicylic acid disc

Product Information						
Product T ype	HUMAN OTC DRUG	Item Code (Source)		NDC:0363-0819		
Route of Administration	TOPICAL					
Active Ingredient/Active Moiety						
Ingre	Strength					
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ) SALICYLIC ACID 17			17.67 mg in 44.18 mg			
Inactive Ingredients						
Ingredient Name				Strength		

PO	LYVINYL ALCOHO	DL (UNII: 532B59J990)		
HIC	GH DENSITY POLY	ETHYLENE (UNII: UG00KM4WR7)		
VIN	NYL ACETATE (UN	II: L9MK238N77)		
Pa	ckaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1 N	NDC:0363-0819-01	1 in 1 CONTAINER	06/01/2013	
1		4 in 1 CELLO PACK		
1		44.18 mg in 1 PATCH; Type 0: Not a Combination Product		
M	arketing Info	ormation		
M	arketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
1410		part358 F	06/01/2013	

Labeler - WALGREENS Co. (008965063)

Registrant - PEDIFIX, INC. (122271935)

Establishment

Name	Address	ID/FEI	Business Operations
PEDIFIX		122271935	pack(0363-0819), label(0363-0819)

Establishment

Name	Address	ID/FEI	Business Operations
AKRON		186569323	manufacture(0363-0819)

Revised: 3/2020

WALGREENS Co.