BLUEDOT CORN REMOVERS- salicylic acid patch CREST MEDICAL LIMITED

BLUEDOTTM Corn Removers

Drug Facts

Active ingredient (in each dosage unit)

Salicylic acid 40% in a plaster vehicle

Purposes

Corn and calluses removal

Uses

- For the removal of corns and calluses
- Relieves pain by removing corns and calluses
- Adhesive bandage holds the medicated pad in position

Warnings

For external use only

Do not use

- If you are a diabetic
- If you have poor blood circulation
- On irritated skin or any area that is infected or reddened

Stop use and ask a doctor if

Discomfort lasts

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
- Apply medicated bandage after 48 hours, remove medicated bandage
- 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 0° to 40°C (32° to 104°F)

Inactive ingredients

Acrylic adhesive, Titanium dioxide, Ferric hydroxide

Questions?

1-800-723-2465

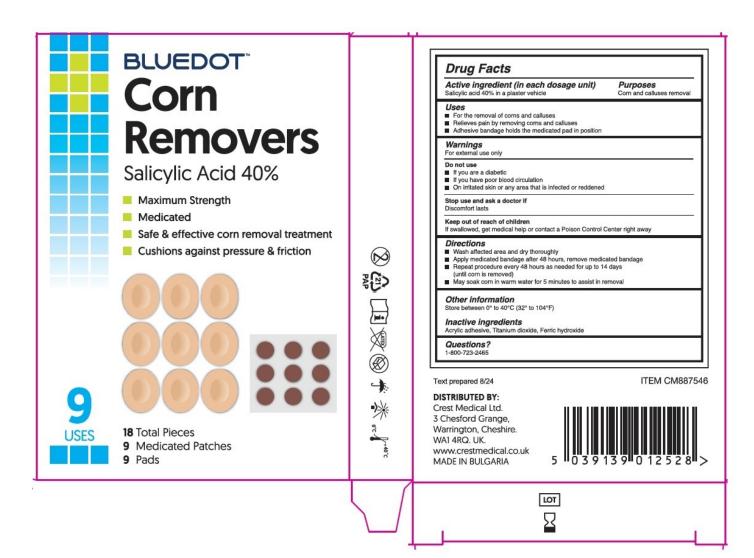
- Maximum Strength
- Medicated
- Safe & effective corn removal treatment
- Cushions against pressure & friction

DISTRIBUTED BY:

Crest Medical Ltd.
3 Chesford Grange,
Warrington, Cheshire.
WA1 4RQ. UK.

www.crestmedical.co.uk MADE IN BULGARIA

Packaging



BLUEDOT CORN REMOVERS

salicylic acid patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84713-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	40 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
FERRIC HYDROXIDE (UNII: 2UA751211N)		

Packaging				
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84713- 102-09	1 in 1 BOX	09/15/2024	
1		9 in 1 BAG		
1		0.0044 g in 1 PATCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M030	09/15/2024		

Labeler - CREST MEDICAL LIMITED (238858539)

Establishment			
Name	Address	ID/FEI	Business Operations
KRE EOOD		565504983	manufacture(84713-102) , label(84713-102) , pack(84713-102)

Revised: 9/2024 CREST MEDICAL LIMITED