BLUEDOT ONE-STEP CORN REMOVER STRIPS- salicylic acid plaster
CREST MEDICAL LIMITED

BLUEDOT™ One-Step Corn Remover Strips

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 Pad**
- **Water Resistant**
- **Stays in place**

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

www.crestmedical.co.uk
MADE IN BULGARIA

Packaging

BLUEDOT ONE-STEP CORN REMOVER STRIPS

salicylic acid plaster

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84713-101
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 g in 100 g

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84713-101-06	1 in 1 BOX	09/15/2024	
1		6 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(84713-101) , label(84713-101) , pack(84713-101)

Revised: 9/2024 CREST MEDICAL LIMITED