

**GEL CORN REMOVER- salicylic acid patch**  
**HEB**

-----  
**HEB Gel Corn Remover**

***Active Ingredient***

Salicylic acid 40%

***Purpose***

Corn remover

***Uses***

- for the removal of corns
- relieves pain by removing corns

***Warnings***

**For external use only.**

**Do not use**

- if you are a diabetic
- if you have poor blood circulation
- on irritated skin or 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 thoroughly
- carefully apply adhesive side of medicated disc onto corn
- cover medicated disc with gel cushion
- after 48 hours remove the medicated disc
- repeat this 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° and 30°C (59° and 86°F)

- avoid surrounding skin when applying medicated disc

### ***Inactive ingredients***

acrylic adhesive, acrylic polymer, polyethylene, polyvinyl alcohol

### **Questions?**

call 1-866-964-0939

### **Principal Display Panel**

HEB

MAXIMUM STRENGTH

MEDICATED

CLEAR GEL

CORN

REMOVERS

SALICYLIC ACID 40% / CORN REMOVER

- EFFECTIVE CORN REMOVAL TREATMENT
- THIN, FLEXIBLE BANDAGE CONCEALS & PROTECTS
- CLEAR GEL CUSHIONS AFFECTED AREA

**6 CUSHIONS**

**6 MEDICATED DISCS**



## GEL CORN REMOVER

salicylic acid patch

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:59640-802
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	400 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	

**POLYVINYL ALCOHOL, UNSPECIFIED** (UNII: 532B59J990)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59640-802-01	6 in 1 BOX	05/21/2026	
1		1 g in 1 PATCH; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M030	05/21/2026	

**Labeler** - HEB (007924756)

Revised: 5/2026

HEB