## NUMBING CREAM MAXIMUM STRENGTH- lidocaine cream BLOOMWELL GARDENS LLC

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Lidocaine 4% --- Topical Anesthetic

Purified Water, Glycerin, Allantoin, Tocopheryl Acetate

(Vitamin E), Carbomer, Propylene Glycol,

Sodium Hydroxide, Phenoxyetha-nol, Ethylhexylglycerin

Local anesthetic

Temporarily helps reduce discomfort from: Minor burns, insect bites, minor skin irritations, Post-shaving or grooming sensitivity External itching or rash due to dry skin

For external use only

- For external use only
- Do not use on large areas of the body or on broken, blistered, or abraded skin.
- Avoid contact with eyes, mouth, and mucous membranes.
- If pregnant or breastfeeding, consult a healthcare provider before use

## Stop use and ask a doctor if:

Condition worsens or symptoms persist for more than 7 days Skin irritation, rash or allergic reaction occurs.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Tamper evident: Do not use if seal is broken or missing.

- Adults and children over 12 years:
- Apply a thin layer to affected area not more than 3-4 times daily
- Children under 12 years: Consult a doctor before use
- Wash hands after application



57\*57\*46mm

171\*22mm



## **NUMBING CREAM MAXIMUM STRENGTH**

lidocaine cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85284-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
<b>LIDOCAINE</b> (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
VITAMIN E POLYETHYLENE GLYCOL SUCCINATE (UNII: 003S90U1F2)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
ALLANTOIN (UNII: 344S277G0Z)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
GLYCERIN (UNII: PDC6A3C0OX)		
WATER (UNII: 059QF0KO0R)		
CARBOMER (UNII: 0A5MM307FC)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
<b>1</b> NDC:85284-006-01	50 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/29/2025		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/29/2025	

## Labeler - BLOOMWELL GARDENS LLC (124839891)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Jiangxi Hemei Pharmaceutical Co.,Ltd		724892056	manufacture(85284-006)	

Revised: 7/2025 BLOOMWELL GARDENS LLC