

BLACK COBRA 4% LIDOCAINE NUMBING CREAM- lidocaine cream
BLACKCOBRASUPPLY, LLC

Purpose

Topical anesthetic

Uses

Temporarily relieves pain and itching due to minor cuts, scrapes, burns, insect bites, and skin irritations.

Warnings

For external use only.

Avoid contact with eyes.

Do not use:

- on large areas of the body
- on broken or blistered skin
- for more than 7 days unless directed by a doctor

Stop use and ask a doctor if:

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days
- redness, irritation, swelling, or pain persists or increases

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

Dosage and Administration

Adults and children 12 years and older:

- Apply a thin layer to affected area not more than 3 to 4 times daily.

Children under 12 years:

- Consult a doctor before use.

Inactive Ingredients

Purified Water, Propylene Glycol, Glycerin, Cetyl Alcohol, Stearyl Alcohol, Carbomer 1342, Benzyl Alcohol, Sodium Hydroxide.

Keep Out of Reach of Children

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Black Cobra 4% Lidocaine Numbing Cream - Principal Display Panel

See image below.



BLACK COBRA 4% LIDOCAINE NUMBING CREAM

lidocaine cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85924-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CARBOMER 1342 (UNII: 809Y72KV36)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics

Color	pink	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85924-001-02	1 in 1 CARTON	07/08/2025	
1	NDC:85924-001-01	20 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/08/2025	

Labeler - BLACKCOBRASUPPLY, LLC (119545644)

Registrant - BLACKCOBRASUPPLY, LLC (119545644)