ADELLINA NUMBING CREAM- lidocaine cream
Guangzhou Haishi Biological Technology Co., Ltd.

Adellina Numbing Cream

Active ingredients

Lidocaine 5%

Purpose

Local Anesthetic

Uses

For the physical or local anesthesia associated with tattoo

For external use only.

Avoid contact with the eyes Keep away from heat or flame. Do not use in large quantities, particularly over raw surfaces or blistered areas.

Stop use and ask a doctor if redness, irritation, swelling, pain, or other symptoms occurs,or If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Do not use this product if Pregnant or breastfeeding, ask a health professional before use. In case of accidental overdose, get medical help or contact the Poison Control Center immediately. Seal is broken or missing.

Do not exceed the recommended daily dosage unless directed by a doctor. certain persons can develop allergic reactions to ingredients in this product. do not put this product into the rectum by using finger.

Keep out of reach of children

In case of accidental ingestion, seek medical help immediately.

Dosage

Squeeze out an appropriate amount of product and spread evenly on skin.

Inactive ingredients

AQUA 71.3%
PROPYLENE 5%
CARBOMER 0.25%

GLYCERIN 5%
DISODIUM EDTA 0.05%
HYDROXYACETOPHENONE 0.4%
1,2-HEXANEDIOL 5%
MENTHA ARVENSIS LEAF EXTRACT 2%
CHRYSANTHELLUM INDICUM EXTRACT 2%
PORTULACA OLERACEA EXTRACT 2%
PEG-40 HYDROGENATED CASTOR OIL 2%

BRAND - ADELLINA NUMBING TATTOO CREAM SIZE : 44.5 X 44.5 X56MM LABEL: 126 X29MM

size:44.5*44.5*56mm

size:29*126mm

ADELLINA NUMBING CREAM

lidocaine cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60771-0011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60771-0011-1	44 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/27/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/27/2025	

Labeler - Guangzhou Haishi Biological Technology Co., Ltd. (421262738)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Haishi Biological Technology Co., Ltd.		421262738	manufacture(60771-0011)

Revised: 3/2025 Guangzhou Haishi Biological Technology Co., Ltd.