

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

Adellina Tattoo Numbing Cream

Active Ingredients

Lidocaine 5%

Purpose

Use

For the physical or local anesthesia associated with tattoo

For external use only.

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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

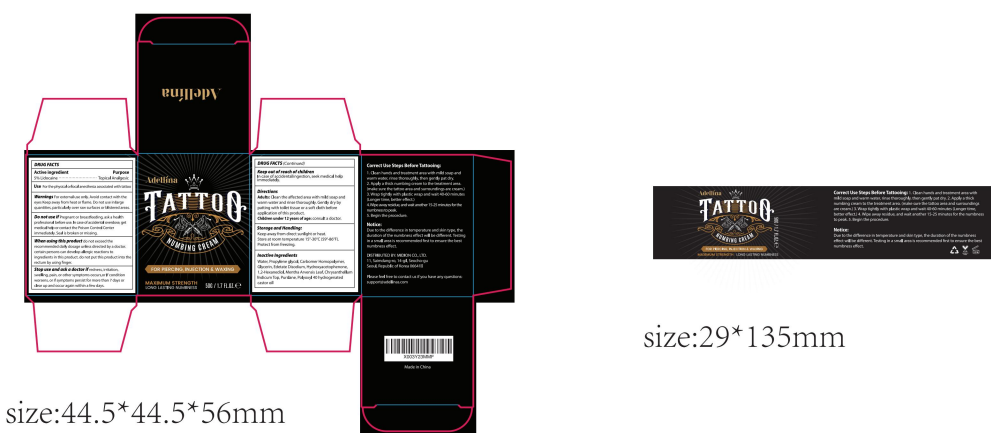
Squeeze out an appropriate amount of product and spread evenly on skin

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: 135 X29MM



ADELLINA TATTOO NUMBING CREAM
lidocaine cream, augmented

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60771-0010
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 g
Inactive Ingredients			
Ingredient Name			Strength

1,2-HEXANEDIOL (UNII: TR046Y3K1G)				
GLYCERIN (UNII: PDC6A3C0OX)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60771-0010-1	50 g 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(60771-0010)