PRINUMB TATTOO NUMBING CREAM- tattoo numbing cream cream Shijiazhuang Auro Technology Co., Ltd.

84555-001

Active Ingredient

Lidocaine 5%

Purpose

Local Anesthetic

Use

For temporarily relieves pain and itching associated with minor burns, minor cuts or skin irritations

Warnings

For External Use Only

Do not use

Pregnant or breastfeeding, ask your doctor before use. in case of accidental overdose, seek immediate medical help or contact the Poison Control Center. seal is broken or missing.

When Using

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 fingers or any medical device or applicator.

Stop Use

The symptom being treated does not improve, or if redness, irritation, swelling, pain, or other symptoms appear or worsen. Seek medical help right away.

Ask Doctor

discomfort persists for 7 days.

Keep Oot Of Reach Of Children

Avoid contact with the eyes. If swallowed, seek medical attention immediately.

Directions

Apply up to 3 times a day.

Children (under 12 years of age): consult a doctor.

Adults: before applying this product, clean the affected area with mild soap and warm water and rinse thoroughly. Pat dry the area with a soft cloth or tissue.

Other information

Store at room temperature and out of direct sunlight or heat.

Inactive ingredients

.Alpha.-Tocopherol Acetate
Carbomer 940
Chrysanthemum Extract
Glycerol
Lecithin, Soybean
Portulaca Oleracea Extract
Prilocaine
Propylene Glycol
Sodium Hyaluronate
Water

PRINCIPAL DISPLAY PANEL

DRUG FACTS

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NUMBING CREAM







5% LIDOCAINE 3.4oz/100ml

Control of the second DRUG FACTS (Continued)

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DRUG FACTS (Continued)

Active Ingredient

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Warnings (For External Use Only)
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5% LIDOCAINE | 1.7oz/50ml

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STOP USE AND ASK A DOCTOR IF • the symptom being treated



TATTOO NUMBING CREAM

DRUG FACTS(Continued)

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KEEP OUT OF REACH OF CHILDREN - Avoid contact

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OTHER INFORMATION - Store at room temperature. - Out

INACTIVE INGREDIENTS Alpha-Tocopherol Acetate, Carbomer 940, Chrysanthemum Extract, Glycerol, Lecithin, Soybean, Portulaca Oleracea Extract, Prilocaine, Propylene Glycol, Sodium Hyaluronate, Water

CORRECT USE STEPS BEFORE TATTOOING

1.Clean hands and treatment area with mild soap and warm wate

rinse thoroughly and gently pat dry,

2.Apply a thick numbing cream to the treatment area, making sure

Zappy a linck numbing cream to the treatment area, making sure to cover the tattoo and its surroundings.

3. Wrap tightly withe plastic wrap and waif for 40-60minutes.

4. Wipe away any excess residue and allow an additional 15-25 minutes for the numbing effect to reach its peak then strat working.

NOTICE the duration of numbness may vary depending on factors such as skin type and temperature, it is recommended to perform a samil test area first to determine the optimal numbness effect.

MANUFACTURED BY Shijiazhuang Auro Te Room 1501, Building B, Dongsheng Plaza, NO.508 Zhongshan East Road, Chang'an District, Shijiazhuang, Hebei, China.

Lot No. : Mfg Date :







PRINUMB TATTOO NUMBING CREAM

tattoo numbing cream cream

Product Information

Product Type HUMAN OTC DRUG NDC:84555-001 **Item Code (Source)**

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
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CHRYSANTHEMUM INDICUM WHOLE (UNII: O9ECF2PL1F)	
PRILOCAINE (UNII: 046O35D44R)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
GLYCERIN (UNII: PDC6A3C0OX)	

.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
WATER (UNII: 059QF0KO0R)	
PURSLANE (UNII: M6S840WXG5)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:84555- 001-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/17/2024		
2	NDC:84555- 001-02	100 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package	07/17/2024		
3	NDC:84555- 001-03	150 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/17/2024		
4	NDC:84555- 001-04	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/17/2024		
5	NDC:84555- 001-05	30 mL in 1 TUBE; Type 0: Not a Combination Product	07/17/2024		
6	NDC:84555- 001-06	60 mL in 1 TUBE; Type 0: Not a Combination Product	07/17/2024		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M017	07/17/2024			

Labeler - Shijia*z*huang Auro Technology Co., Ltd. (406846461)

Establishment				
Name	Address	ID/FEI	Business Operations	
Shijiazhuang Auro Technology Co., Ltd.		406846461	manufacture(84555-001)	

Revised: 7/2024 Shijiazhuang Auro Technology Co., Ltd.