ATOZCUP TATTOO NUMBING- lidocaine cream
Dongguan City Bobiter Electronic Technology Co., Ltd

Active Ingredient(s)

Lidocaine 5%

Purpose

local anesthetic

Use

For temporary relief of local discomfort, itching, pain,soreness or burning of the skin

Warnings

For external use only.

Do not use

Do not exceed the recommended daily dosage unless directed by a doctor.

When using Avoid contact with eyes. If this happens, rinse thoroughly with water.

If allergy occurs such as redness,irritation,swelling,pain or other symptoms become severe Stop use and get to a doctor.If symptoms persist for more than 7days or subside but occur again within3 days,stop use and consult aphysician.

Keep out of reach of children.

Directions

1.Wash and exfoliate the area to be treated.
2.Dry the area completely.

3.Apply this numbing cream(2mm thick) to the area and rub in thoroughly

4.Cover with cling film.

5.leave the cream and wrap in place for 60-90 minutes before procedure.

6.After the wrapping time,remove thecling film and wipe the cream away

Storage and handling

*Always keep tihgtly closed

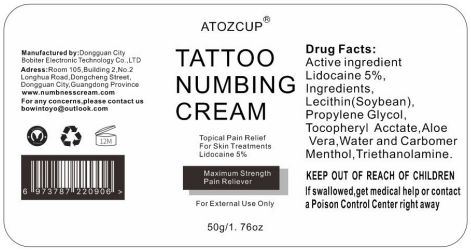
*Store at temperatures between 59°F-86°F(15°C-30°C)

*Protectfrom freezing

Inactive ingredients

Lecithin(Soybean)、Propylene Glycol、Tocopheryl Acctate,Aloe Vera、Water、Carbomer,Menthol、Triethanolamine

Package Label - Principal Display Panel



ATOZCUP TATTOO NUMBING

lidocaine cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83706-001
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
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL SUCROSE CROSSLINKED) (UNII: Z135WT9208)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83706-001-03	1 in 1 PACKAGE	09/22/2023	
1		50 g in 1 BOTTLE; 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	09/22/2023	

Labeler - Dongguan City Bobiter Electronic Technology Co., Ltd (411899304)

Registrant - Dongguan City Bobiter Electronic Technology Co., Ltd (411899304)

Establishment

Name	Address	ID/FEI	Business Operations
Dongguan City Bobiter Electronic Technology Co., Ltd		411899304	manufacture(83706-001)

Revised: 12/2023 Dongguan City Bobiter Electronic Technology Co., Ltd