

TATTOO NUMBING CREAM- lidocaine paste
Shenzhen Yangan Technology Co., Ltd.

Tattoo Numbing Cream

Tattoo Numbing Cream

Lidocaine 5%

Topical Anesthetic

For temporary relief of local discomfort, itching, pain, soreness, or burning in the perianal area associated with anorectal disorders.

For external use only

Pregnant or breastfeeding, ask a health professional before use.

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

Do not put into the rectum by using fingers or any medical device or applicator.

Condition worsens, or does not improve within 7 days. Certain persons can develop allergic reactions to ingredients in this product. If the symptom being treated does not subside or if redness, irritation, swelling, pain, or other symptoms develop or increase, discontinue use.

In case of accidental ingestion, seek medical attention immediately.

Adults and Children 12 years and older: When practical, cleanse the affected area with mild soap and warm water and rinse thoroughly. Gently dry by patting or blotting with toilet tissue or a soft cloth, before application of this product. Apply externally to the affected area up to 6 times daily.

Keep away from direct sunlight or heat.

Store in room temperature (59-86°F/15-30°C).

Aloe barbadensis leaf juice (Aloe vera)

Water

Lecithin (Soybean)

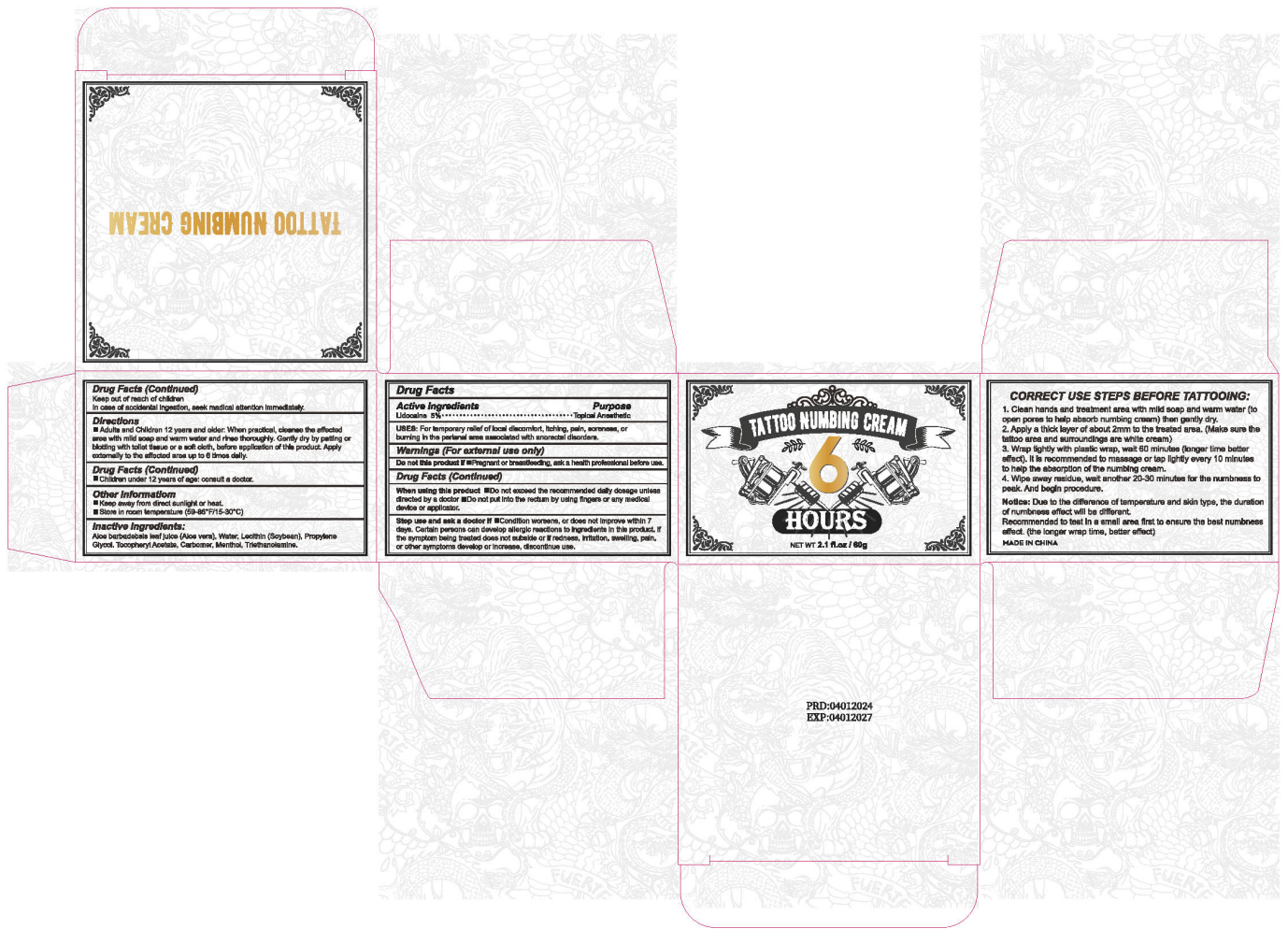
Propylene Glycol

Tocopheryl Acetate

Carbomer

Menthol

Triethanolamine



TATTOO NUMBING CREAM

lidocaine paste

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84023-201
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

ALOE (UNII: V5VD430YW9)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84023-201-01	1 in 1 BOX	01/30/2024	
1		60 g in 1 BOTTLE, PLASTIC; 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	01/30/2024	

Labeler - Shenzhen Yangan Technology Co., Ltd. (419283765)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Yangan Technology Co., Ltd.		419283765	manufacture(84023-201)

Revised: 4/2024

Shenzhen Yangan Technology Co., Ltd.