

**TATTOOGIRL NUMBING- lidocaine hcl cream**  
**Prodigy Media Inc**

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**Active Ingredient**

Lidocaine HCL 4%

**Purpose**

External Analgesic

**Uses**

For the temporary relief of pain and itching associated with sunburns, minor cuts, insect bites, and skin irritations

**Warnings**

- **For external use only.**
- **Avoid contact with eyes**

**Stop use and ask a doctor if**

- Condition worsens or symptoms persist for more than 7 days
- Symptoms clear up and occur again within a few days
- Do not use in large quantities, particularly over raw surfaces or blistered areas. Do not exceed the recommended daily dosage unless directed by a doctor.

**Do not use**

- On wounds or damaged skin.

**Keep out of reach of children**

If product is swallowed, get medical help or contact a Poison Control Center right away.

***Direction***

Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. Children under 2 years of age: do not use, consult a physician.

***Other information***

Do not use if seal is broken.

***Inactive ingredients***

Aqua (Deionized Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Chondroitin Sulfate, Dimethyl Sulfone (MSM), Emu Oil, Ethoxydiglycol, Ethylhexylglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Phenoxyethanol, Polyacrylamide, Propylene Glycol, Stearic Acid, Tea Tree Oil, Triethanolamine.

***Questions?* 866-488-0066**

**Product label**



TATTOO\_GIRL\_1oz Label.jpg





TATTOO\_GIRL\_1oz BOX.jpg

## TATTOOGIRL NUMBING

lidocaine hcl cream

### Product Information

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:70171-0021

Route of Administration	TOPICAL
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**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ARNICA MONTANA FLOWER</b> (UNII: OZ0E5Y15PZ)	
<b>C13-14 ISOPARAFFIN</b> (UNII: E4F12ROE70)	
<b>ELOSULFASE ALFA</b> (UNII: ODJ69JZG85)	
<b>DIMETHYL SULFONE</b> (UNII: 9H4PO4Z4FT)	
<b>EMU OIL</b> (UNII: 344821WD61)	
<b>DIETHYLENE GLYCOL MONOETHYL ETHER</b> (UNII: A1A1I8X02B)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>GLUCOSAMINE SULFATE</b> (UNII: 1FW7WLR731)	
<b>ISOPROPYL PALMITATE</b> (UNII: 8CRQ2TH63M)	
<b>LAURETH-7</b> (UNII: Z95S6G8201)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>POLYACRYLAMIDE (CROSSLINKED; 2 MOLE PERCENT BISACRYLAMIDE)</b> (UNII: 9FPL31B58Q)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TEA TREE OIL</b> (UNII: VIF565UC2G)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70171-0021-1	1 in 1 CARTON	07/27/2025	
1		28 g in 1 TUBE; 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	07/27/2025	

**Labeler** - Prodigy Media Inc (080011622)