

TATTOO NUMBX NUMBING PAIN RELIEVING- lidocaine hydrochloride cream
Scyberco Ltd

TATTOO NUMBX NUMBING CREAM PAIN RELIEVING

Drug Facts

Active ingredients

Lidocaine hydrochloride 4%

Purpose

Topical Analgesic

Uses:

- For the temporary relief of pain and itching associated with minor burns, sunburn, minor cuts, scrapes, Insect bites, minor skin irritations, rashes due to poison ivy, poison oak, or poison sumac.

Warnings:

For external use only

Do not use

- in large quantities, particularly over raw surfaces or blistered areas.

When using this product

- Avoid contact with the eyes

Stop use and ask a doctor if

- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

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

If pregnant or breast-feeding,

ask a health professional before use

Directions:

- 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: consult a doctor.

Other information:

- Store at 20-25°C (68-77°F) and protect from moisture

Inactive ingredients:

Aloe Barbadensis Leaf Juice, Carbomer, Cetearyl Alcohol, Dimethyl Sulfone, Ethylhexylglycerin, Glyceryl Stearate SE, Helianthus Annuus (Sunflower) Seed Oil, Phenoxyethanol, Polysorbate 20, Polysorbate 60, Simmondsia Chinensis (Jojoba) Seed Oil, Stearic Acid, Triethanolamine, Water (Aqua)

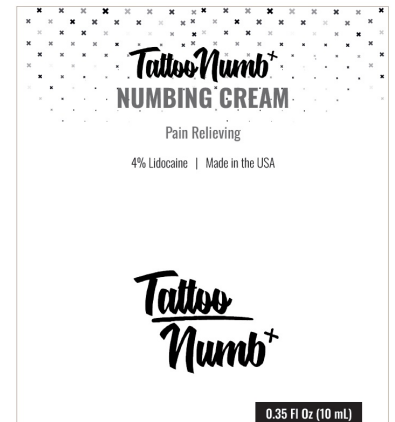
Package Labeling:




Tube



Zoomed in 200% without guides (printable area)





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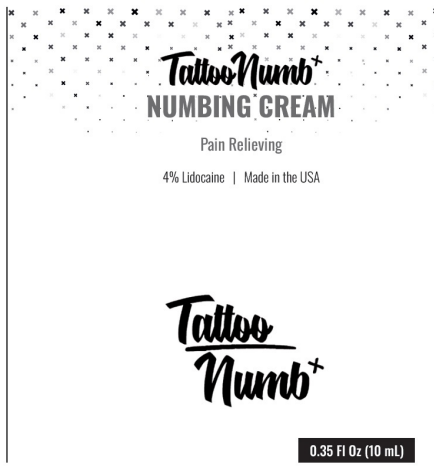
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TATTOONUMBX





TATTOO NUMBX NUMBING PAIN RELIEVING

lidocaine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87460-905
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

POLYSORBATE 20 (UNII: 7T1F30V5YH)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87460-905-00	1 in 1 BOX	01/23/2026	
1		10 mL 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	01/23/2026	

Labeler - Scyberco Ltd (234099080)

Registrant - Pure Source, LLC (080354456)

Revised: 4/2026

Scyberco Ltd