DERMA NUMB- lidocaine hci spray
Atlas Tat Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Derma Numb Tattoo Anesthetic Spray

Active Ingredients

Lidocaine HCI

Purpose

Topical Anesthetic

Uses

Temporarily relieves pain from tattoo procedures.

Directions

First test product to sensitivity to skin.

Once skin is broken or outline is done, apply derma numb generously to tattooed area.

Wait two minutes for anesthetic to take affect, re-apply derma numb to tattoo area as needed through out the tattoo process for comfort. Discontinue use if sensitivity occurs. Do not use on face.

Warnings

External Use Only

Avoid contact with the eyes

Do Not Use

If you have any known allergy or sensitivity to any of these ingredients in this product. Discontinue use and seek medical attention.

Stop Use and ask a doctor if

• skin becomes irritated • condition worsens or symptoms last more than7 days • symptoms clear up and reoccur with a few days

Inactive ingredients

Water, Propylene Glycol, Achillea Millefolium (Yarrow) Extract, Yucca Glauca Root Extract, Citric Acid, Sodium Metabisulfate, Caprylyl Glycol, Disodium EDTA, Methylisothiazolinone.

Other Information

This product was manufactured for Atlas Tat, Inc.

Any questions call 954-492-9898 or check out the web @ dermnumb.com

Drug Facts
For Professional Use Only

Active Ingredients — **Purpose**
Lidocaine HCl (4% w/w) — Topical Anesthetic

Uses Temporarily relieves pain from tattoo procedures.

Directions First test product to sensitivity to skin
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Warnings External Use Only
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DERMA NUMB™ Spray
TATTOO ANESTHETIC
1oz.

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DERMA NUMB
lidocaine hci spray

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:53200-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ACHILLEA MILLEFOLIUM (UNII: 2FXJ6SW4PK)	
YUCCA GLAUCA ROOT (UNII: 1A15YBH7N1)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53200-001-01	28.35 g in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	07/15/2012	

Labeler - Atlas Tat Inc. (061275633)

Revised: 10/2012 Atlas Tat Inc.