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

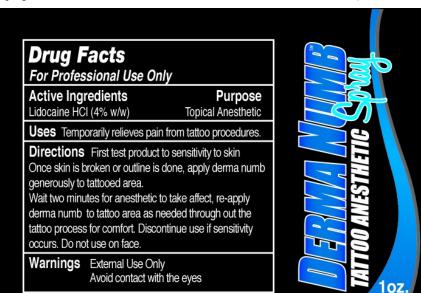
• skin becomes irritated • condition worsens or symptoms last more than 7 days • symptoms clear up and reoccur with a few days

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.



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This product was manufactured for Atlas Tat, Inc. Any questions call 954-492-9898 or check out the web @ dermanumb.com

DERMA NUMB

lidocaine hci spray

Product Information

Product TypeHUMAN PRESCRIPTION DRUGItem Code (Source)NDC:53200-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDO CAINE UNII:98 PI200987)
LIDO CAINE
HYDRO CHLO RIDE
40 mg in 1 g

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
SODIUM METABISULFITE (UNII: 4VON5FNS3C)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)				
ACHILLEA MILLEFO LIUM (UNII: 2FXJ6SW4PK)				
YUCCA GLAUCA ROOT (UNII: 1A15YBH7N1)				
DIAZO LIDINYL 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.