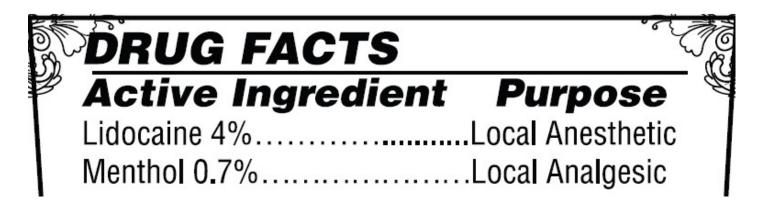
NUPHARMISTO TATTOO NUMBING CREAM 60ML- lidocaine cream Orange Lab, Inc

Nupharmisto Tattoo Numbing Cream 60ml

Drug Facts



Purpose

Uses: For the temporary relief of pain and itching associated withminor burns, sunburn, minor cuts, scrapes, and insect bites.

Warnings

Warning (For external use only)

Stop and ask a doctor if

- an allergic reaction occurs.
- symptom does not improve or if redness, irritation, swelling, pain or other symptoms develop or increase.

When use this product

do not exceed the recommended daily dosage unless directed by a doctor. Avoid contact with the eyes. Do not use if you are allergic to the ingredients in this product or if the seal is broken or missing.

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of accidental ingestion, seek medical advice immediately or contact a Poison Control Center right away.

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If pregnant of breast feeding section

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other information (storage)

Other Information

- Store room temperature 59°-86°F (15°-30°C).
- Keep away from direct sunlight or heat.

Inactive Ingredients

Inactive Ingredients

DI Water, Dimethyl Isosorbide, Ethylhexyl Palmitate, Hydrogenated Polydecene, Propylene Glycol, Arnica Montana Flower Extract, Helianthus Annuus (Sunflower) Seed Oil, Emu Oil, Phenoxyethanol, Sodium Polyacrylate, Trideceth-6, Menthol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Allantoin, Tocopheryl Acetate, Ethylhexylglycerin, Tetrasodium EDTA.

Directions for use

Directions

- For children under 12, consult a doctor.
- For adults: If possible, clean the affected area with mild soap and warm water, making sure to rinse thoroughly. Gently dry the area with toilet tissue or a soft cloth before applying the product.
- Apply up to 4 times daily.

Ouestions? Contact us.

Questions or comments?

Contact us at service@nupharco.com.

dosage and how to use it correctly section





How to use it correctly?

- Clean your hands and treatment area thoroughly with mild soap and warm water. Gently dry them afterward.
- Apply a thick layer of numbing cream to the treatment area, making sure to cover both the tattooed area and its surroundings.
- Wrap the area securely with plastic wrap and leave it on for 40-50 minutes for the best results (keeping iton longer gives better effects).
- Remove any remaining cream and wait an additional 10 minutes for the numbing sensation to reach its peak.
- Begin the procedure.

Notic: The duration of numbness may vary depending on temperature and individual skin types. It's recommended to perform a small patch test for the best numbing effect.





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Label Images

DRUG FACTS Active Ingredient Purpose Lidocaine 4%.....Local Anesthetic Menthol 0.7%.....Local Analgesic **Uses:** For the temporary relief of pain and itching associated withminor burns, sunburn, minor cuts, scrapes, and insect bites.

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Directions

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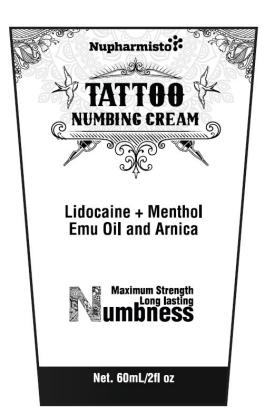
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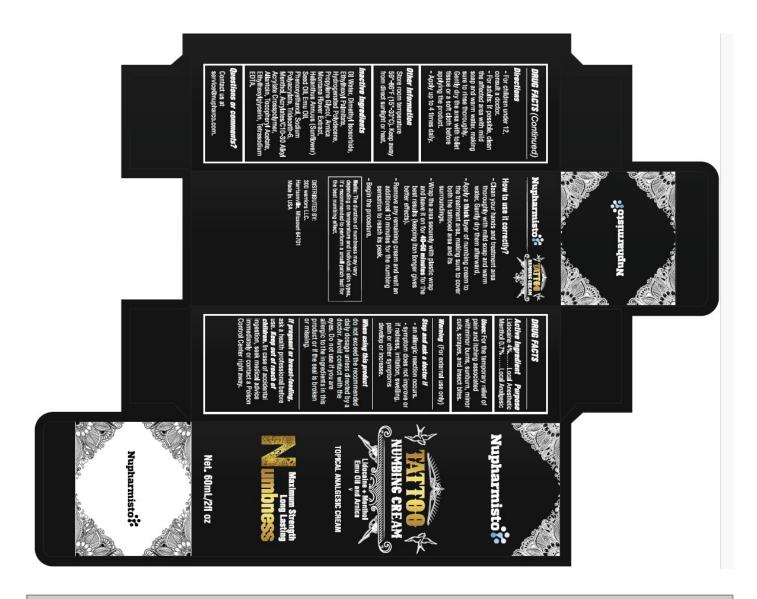
Other Information

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Inactive Ingredients

DI Water, Dimethyl Isosorbide, Ethylnexyl Palmitate, Hydrogenated Polydecene, Propylane Glycol, Arrica Montana Flower Extract, Helanthus Annus (Sunflower) Seed Oil, Emu Oil, Penoxyethanol, Sodium Polyacyylate, Trideceth-6, Menthol, Arrylatas (T.O.3 Allyl Acyylate Crosspolyme, Allandi, Tocopheryl Acetate, Ethylnexylglycerin, Tetrasodium EDTA,





NUPHARMISTO TATTOO NUMBING CREAM 60ML

lidocaine cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71331-119
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 g	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.7 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)		
ALLANTOIN (UNII: 344S277G0Z)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		

WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EMU OIL (UNII: 344821WD61)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
HYDROGENATED POLYDECENE (1500 CST) (UNII: 4YI0729529)	
SUNFLOWER OIL UNSAPONIFIABLES (UNII: T7Z E2WA4MB)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	
EDETATE SODIUM (UNII: MP1J8420LU)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71331-119- 02	1 in 1 BOX	09/01/2024	
1		60 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	09/01/2024	

Labeler - Orange Lab, Inc (004862271)

Revised: 9/2024 Orange Lab, Inc