NUPHARMISTO TATTOO NUMBING CREAM 80G- lidocaine cream Orange Lab, Inc

Nupharmisto Lidocaine Tattoo Numbing Cream 4% 80g

Active Ingredient

Drug facts Active Ingredient **Purpose** Lidocaine 4%Topical Analgesic Menthol 0.7%Topical Analgesic

ask doctor

If pregnant or breast-feeding,

ask a health professional before use.

Warning do not use if

- Warnings For external use only. Avoid contact with the eyes. If allergic reaction occurs, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor.

Keep out of reach of children

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In case of accidental ingestion, seek medical attention immediately.

if pregnant or breast feeding

If pregnant or breast-feeding,

ask a health professional before use.

Purpose

DRUG FACTS

Active Ingredient	Purpose
Lidocaine 4%	Topical Analgesic
Menthol 0.7%	Topical Analgesic

questions

Questions or comments? Contact us at service@nupharco.com.

Stop use warnings

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 - If allergic reaction occurs, or if symptoms persist for more than 7 days or clear up and occur again within a few days. discontinue use of this product and consult a doctor.

When using

When using this product Do not use in large quantities, particularly over raw surfaces or blistered areas. Do not use if you are allergic to the ingredients in this product or if the seal is broken/missing.

Directions

- Directions Children under 2 years: Do not use
 - For children under 12: Consult a doctor.
 - For adults: Apply to affected area not more than 3 times daily. 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.

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, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Allantoin, Tocopheryl Acetate, Ethylhexylglycerin, Tetrasodium Edta

warnings sections

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How to use it correctly





How to use it correctly?

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

Note: 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





bottle label



NUPHARMISTO TATTOO NUMBING CREAM 80G lidocaine cream Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration TOPICAL

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

Inactive Ingredients		
Ingredient Name	Strength	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)		
ETHYLHEXYL PALMITATE (UNII: 2865993309)		
HYDROGENATED POLYDECENE (1500 CST) (UNII: 4YI0729529)		
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)		
ALLANTOIN (UNII: 344S277G0Z)		
TRIDECETH-6 (UNII: 3T5PCR2H0C)		
EMU OIL (UNII: 344821WD61)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
EDETATE SODIUM TETRAHYDRATE (UNII: L13NHD21X6)		
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
SUNFLOWER OIL (UNII: 3W1JG795YI)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
WATER (UNII: 059QF0KO0R)		
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71331-116- 04	80 g in 1 PACKAGE; Type 0: Not a Combination Product	05/01/2024	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	05/01/2024		

Labeler - Orange Lab, Inc (004862271)

Revised: 5/2024 Orange Lab, Inc