

NUPHARMISTO TATTOO NUMBING CREAM- tattoo numbing cream cream
Ehy Holdings LLC

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.

81799-008
Nupharmisto Tattoo Numbing Cream

Active Ingredient(s)

Lidocaine 5%

Purpose

Local Anesthetic

Use

For the temporary relief of local and anorectal discomfort associated with anorectal disorder or inflammation.

Warnings

For external use only

Do not use

Pregnant or breastfeeding, ask a health professional before use.

In case of accidental overdose, get medical help or contact the Poison Control Center immediately.

Seal is broken or missing

directed by a doctor.

certain persons can develop allergic reactions to ingredients in this product.

do not put this product into the rectum by using finger or any medical device or applicator

the symptom being treated does not subside, or if redness

Adults: When practical, cleanse the affected area with mild soap and warm water and rise thoroughly.

Gently dry by patting with toilet tissue or a soft cloth, before application of this product.

Apply up to 6 times a day.

Children under 12 years of age: consult a doctor

Directions

Adults: When practical, cleanse the affected area with mild soap and warm water and rise thoroughly.

Gently dry by patting with toilet tissue or a soft cloth, before application of this product.

Apply up to 6 times a day.

Children under 12 years of age: consult a doctor

Other information

Keep away from direct sunlight or heat.

Store at room temperature 15°-30°C (59°-86°F)

Inactive ingredients

Carbomer 934, Emu Oil, Arnica, Lecithin(Soybean), Propylene Glycol, Tocopheryl Acetate, Benzyl Alcohol, Water

irritation, swelling, pain, or other symptoms develop or increase

service@nupharco.com

Package Label - Principal Display Panel

81799-008-01 1



81799-008-01 2



NUPHARMISTO TATTOO NUMBING CREAM

tattoo numbing cream cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81799-008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	5 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
WATER (UNII: 059QF0KO0R)	
CARBOXPOLYMETHYLENE (UNII: 0A5MM307FC)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81799-008-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/15/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	02/15/2023	

Labeler - Ehy Holdings LLC (117322715)**Establishment**

Name	Address	ID/FEI	Business Operations
Ehy Holdings LLC		117322715	manufacture(81799-008)

Revised: 2/2023

Ehy Holdings LLC