

NUMB 100- lidocaine cream
Clinical Resolution Laboratory, Inc.

Numb 100

DRUG FACTS

Active Ingredient

Lidocaine 5%

Purpose

Local Anesthetic

Uses:

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

Warnings

(For external use only)

Do not use

this product if

Pregnant or breast-feeding,

- ask a health professional before use.
- In case of accidental overdose, contact a doctor or Poison Control Center immediately.
- Tamper Evident "Warranty Void...Seal..."label atop the container is broken.

When using this product

- Do not exceed the recommended daily dosage unless 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 fingers or any medical device or applicator.

Stop use and ask a doctor if

The symptom being treated does not subside, or if redness, irritation, swelling, pain, or other symptoms develop or increase.

Keep out of reach of children

In case of accidental ingestion, seek medical attention immediately.

Directions

- Adults: When practical, cleanse the affected area with mild soap and warm water and rise thoroughly. Gently dry by patting or blotting 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 in room temperature (59-86°F / 15-30°C).

Inactive Ingredients

Water, Triethanolamine, Carbomer, Propylene Glycol, Benzyl Alcohol, Ethoxydiglycol, Lecithin, Neopentyl Glycol Dicaprylate/Dicaprate, Sodium Polacrylate, Hydrogenated Polycene, Trideceth-10, Cholesterol, Allantoin, Tocopheryl Acetate, Polysorbate-80

Package Labeling:

Manufactured for Laser Jet, Inc. Brooklyn NY 11206 - Made in USA	DOCTOR RECOMMENDED	NUMB 100	
			
Topical Anesthetic Cream 1.35 fl. oz e 38g		DRUG FACTS Active Ingredient Lidocaine 5% ----- Purpose Local Anesthetic Uses: For the temporary relief of local and anorectal discomfort associated with anorectal discomfort or inflammation. Warnings (For external use only) Do not use this product if <ul style="list-style-type: none"> ▪ Pregnant or breast-feeding, ask a health professional before use. ▪ In case of accidental overdose, contact a doctor or Poison Control Center immediately. ▪ Tamper Evident "Warranty Void...Seal..." label atop the container is broken. When using this product <ul style="list-style-type: none"> ▪ Do not exceed the recommended daily dosage unless 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 fingers or any medical device or applicator. Stop use and ask a doctor if The symptom being treated does not subside, or if redness, irritation, swelling, pain, or other symptoms develop or increase.	DRUG FACTS (Continued) Keep out of reach of children In case of accidental ingestion, seek medical attention immediately. Directions <ul style="list-style-type: none"> ▪ Adults: When practical, cleanse the affected area with mild soap and warm water and rise thoroughly. Gently dry by patting or blotting 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 <ul style="list-style-type: none"> ▪ Keep away from direct sunlight or heat. ▪ Store in room temperature (59-86°F / 15-30°C). Inactive Ingredients Water, Triethanolamine, Carbomer, Propylene Glycol, Benzyl Alcohol, Ethoxydiglycol, Lecithin, Neopentyl Glycol Dicaprylate/Dicaprate, Sodium Polacrylate, Hydrogenated Polycene, Trideceth-10, Cholesterol, Allantoin, Tocopheryl Acetate, Polysorbate-80

NUMB 100

lidocaine cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63742-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	50 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A18X02B)	
NEOPENTYL GLYCOL DICAPRYLATE/DICAPRATE (UNII: VLW429K27K)	
TRIDECETH-10 (UNII: G624N6MSBA)	
CHOLESTEROL (UNII: 97C5T2UQ7J)	
ALLANTOIN (UNII: 344S277G0Z)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63742-003-01	38 g in 1 BOTTLE; Type 0: Not a Combination Product	06/18/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	06/18/2016	

Labeler - Clinical Resolution Laboratory, Inc. (825047942)

Revised: 12/2023

Clinical Resolution Laboratory, Inc.