

DR. NUMB- lidocaine and benzethonium chloride cream cream
Shinpharma Inc

Drug Facts

Active Ingredient

Lidocaine 4%

benzethonium Chloride 0.25%

Purpose

External Analgesic and First Aid Antiseptic

Uses

- Can be used instead of soap and water to help clean minor cuts, scrapes, and burns
- For the temporary relief of discomfort and pain associated with dermal procedures such as tattoo removal, dermarolling, electrolysis, microblading, and piercing
- Temporarily relieves pain and itch while helping to prevent infection.

Warnings

- **For external use only**
- avoid contact with eyes

Do not use

- do not use in large quantities particularly over raw surfaces or blistered area
- do not exceed the recommended dosage unless directed by a doctor
- in the eyes or apply over large areas of body
- longer than one week unless directed by a doctor

Ask a doctor before use if you have

- deep or puncture wounds, animal bites or serious burns

Stop the use and consult doctor if:

- condition worsens or symptoms persist for more than 7 days or clear up and occur again within few days.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- Use Dr.Numb® Topical Anesthetic Foam Soap to cleanse the targeted area.
- Shake well before each use.

- Apply 2 to 3 pumps of foam soap and gently rub into the skin. Leave it on for 5 to 10 minutes. Rinse off gently with running water or wipe with clean paper towel.
- Use before, during and after the procedure.
- Make sure to cover the bottle tightly when not in use, otherwise, the anesthetic effect will be lessened.

Adults and children two years old and older: Use to clean minor cuts, scrapes, and burns by thoroughly washing with water. Rinse and air dry. Use no more than three times daily.

Children under two years of age, ask a doctor.

Other information

- Store at controlled room temperature 59 °-86 °F (15 °-30 °C)
- Do not expose to temperature above 120 °F (49 °C)
- Protect from freezing

Inactive Ingredients

Allantoin, Aloe Barbadensis Leaf Juice, Arnica Montana Flower Extract, Benzyl Alcohol, Glycerin, Lauramidopropyl Betaine, Leuconostoc/Radish Root Ferment Filtrate, Menthoxypentanediol, Phenoxyethanol, Selaginella Lepidophylla Extract, Sodium Hydroxide, Water.

Question or comments?

Call weekdays 9 AM to 6 PM PST at 1-877-786-2001 or email us at support@drnumb.com

Principal Display Panel

NDC 69903-002-50

Dr.Numb

TOPICAL ANESTHETIC FOAM SOAP

4% LIDOCAINE

0.25% BENZETHONIUM CHLORIDE

DISINFECT DESENSITIZE and RELAX

1.7 oz

Dr. Numb®
TOPICAL ANESTHETIC
DISINFECTANT FOAM SOAP

4% LIDOCAINE
0.25% BENZETHONIUM CHLORIDE

NDC 69903-002-50

FOR EXTERNAL USE ONLY

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Dr. Numb®
TOPICAL ANESTHETIC DISINFECTANT FOAM SOAP

Drug Facts (continued)

Directions

Use Dr.Numb® Topical Anesthetic Disinfectant Foam Soap to cleanse the targeted area. Shake well before each use. Apply 2 to 3 pumps of foam soap and gently rub into the skin. Leave it on for 5 to 10 minutes. Rinse off gently with running water or wipe with clean paper towel. Use before, during and after the procedure. Make sure to cover the bottle tightly when not in use, otherwise, the anesthetic effect will be lessened.

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Dr. Numb®

Distributed by:
Shirpharma Inc.
1124 Pitt Avenue, Blaine, Washington, USA 98230

www.DrNumb.com

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at 1-877-786-2001 or
email us at support@drnumb.com



Dr. Numb®
TOPICAL ANESTHETIC DISINFECTANT FOAM SOAP

4% LIDOCAINE
0.25% BENZETHONIUM CHLORIDE 1.7oz

Disinfect,
Desensitize,
& Relax

Batch No.:

Exp. Date:

DR. NUMB

lidocaine and benzethonium chloride cream cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69903-002	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)		LIDOCAINE	4 mg in 50 g	
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)		BENZETHONIUM CHLORIDE	0.25 mg in 50 g	
Inactive Ingredients				
Ingredient Name			Strength	
ALLANTOIN (UNII: 344S277G0Z)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)				
BENZYL ALCOHOL (UNII: LKG8494WBH)				
GLYCERIN (UNII: PDC6A3C0OX)				
LAURAMIDOPROPYL BETAINE (UNII: 23D6XVI233)				
LEUCONOSTOC/RADISH ROOT FERMENT FILTRATE (UNII: D2QHA03458)				
3-((L-MENTHYL)OXY)PROPANE-1,2-DIOL (UNII: KD6TZ2QICH)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
SELAGINELLA LEPIDOPHYLLA (UNII: 02JQ564P1G)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
WATER (UNII: 059QF0KOOR)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69903-002-50	1 in 1 CARTRIDGE	09/12/2017	
1		50 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		09/12/2017	

Labeler - Shinpharma Inc (248552403)

Registrant - Shinpharma Inc (248552403)