DR. NUMB- lidocaine cream Shinpharma Inc

Dr. Numb

Active ingredient

Lidocaine 5%

Purposes

Anorectal (Hemorrhoidal)

Uses

For temporary relief of local discomfort, itching, soreness or burning in the perianal area associated with anorectal disorders

Warnings

For external use only

When using this product

- do not exceed the recommended daily dosage unless directed by a doctor
- do not put into the rectum by using fingers or any mechanical device or applicator

Stop use and ask a doctor if

- allergic reaction occurs
- rectal bleeding occurs
- redness, irritation, swelling, pain or other symptoms begin or increase
- condition worsens or does not improve within 7 days

KEEP OUT OF REACH OF CHILDREN

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

Directions

Adults and children 12 years and older: When practical, cleanse the affected area with mild soap and warm water and rinse thoroughly. Gently dry by patting or blotting with toilet tissue or a soft cloth before application of this product. Apply to the affected area up to 6 times a day.

Children under 12 years: Consult a doctor

Other information

- always keep the tube tighlty closed
- store at temperatures not exceeding 15°C-30°C (59°F 86°F)
- protect from freezing

Inactive ingredients

Benzyl Alcohol, Carbopol, Letcithin, Propylene Glycol, Tocopheryl Acetate, Water

Questions?

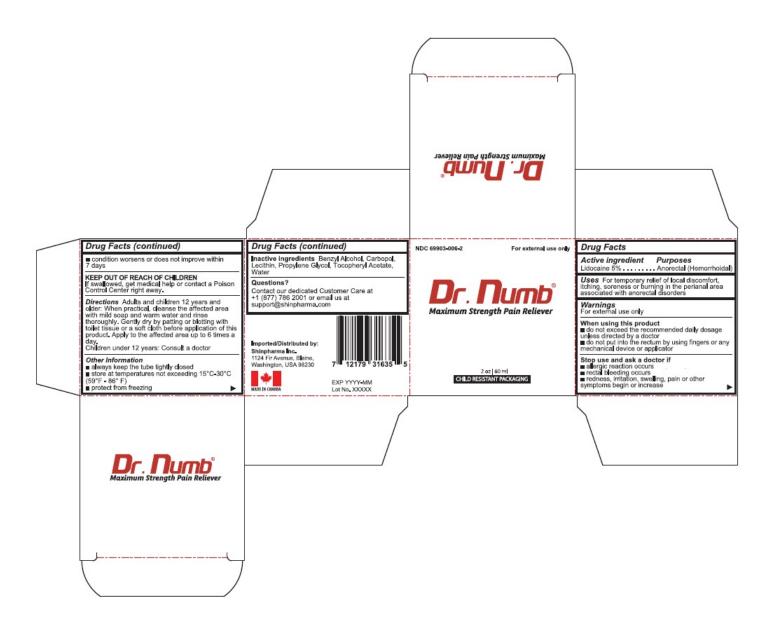
Contact our dedicated Customer Care at + (877) 786 2001 or email us at support@shinpharma.com

Principal Display Panel

Dr. Numb

Maximum Strength Pain Reliever

2 oz | ml



DR. NUMB

lidocaine cream

D		Inform	:
Prod	uct	INTOFM	ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69903-006

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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

Inactive Ingredients		
Ingredient Name	Strength	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		

LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
WATER (UNII: 0590F0KO0R)	

Packaging				
# Ite	m Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC 006-	:69903- 02	1 in 1 CARTON	09/09/2022	
1		60 mL in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	09/09/2022	

Labeler - Shinpharma Inc (248552403)

Registrant - Shinpharma Inc (248552403)

Revised: 4/2024 Shinpharma Inc