BASE LABORATORIES BASE NUMB TOPICAL ANORECTAL CREAM- lidocaine cream
Joonem LLC

Base Laboratories Base Numb Topical Anorectal Cream

Drug Facts

Active ingredients

Lidocaine 5%

Purpose

Topical Analgesic

Indications:

For the temporary relief of pain and discomfort associated with anorectal disorders

Warnings:

If condition worsens or does not improve within 7 days, consult a doctor.

- Do not exceed the recommended daily dosage unless directed by a doctor.
- In case of bleeding, consult a doctor promptly.
- Do not put this product into the rectum by using fingers or any mechanical device or applicator.
- Certain persons can develop allergic reactions to ingredients in this product. If the symptoms being treated does not subside or if redness, irritation, swelling, pain, or other symptoms develop or increase, discontinue use and consult a doctor.

Keep out of reach of children

to avoid accidental ingestion!

If swallowed, get medical help or contact a posion control center immediately.

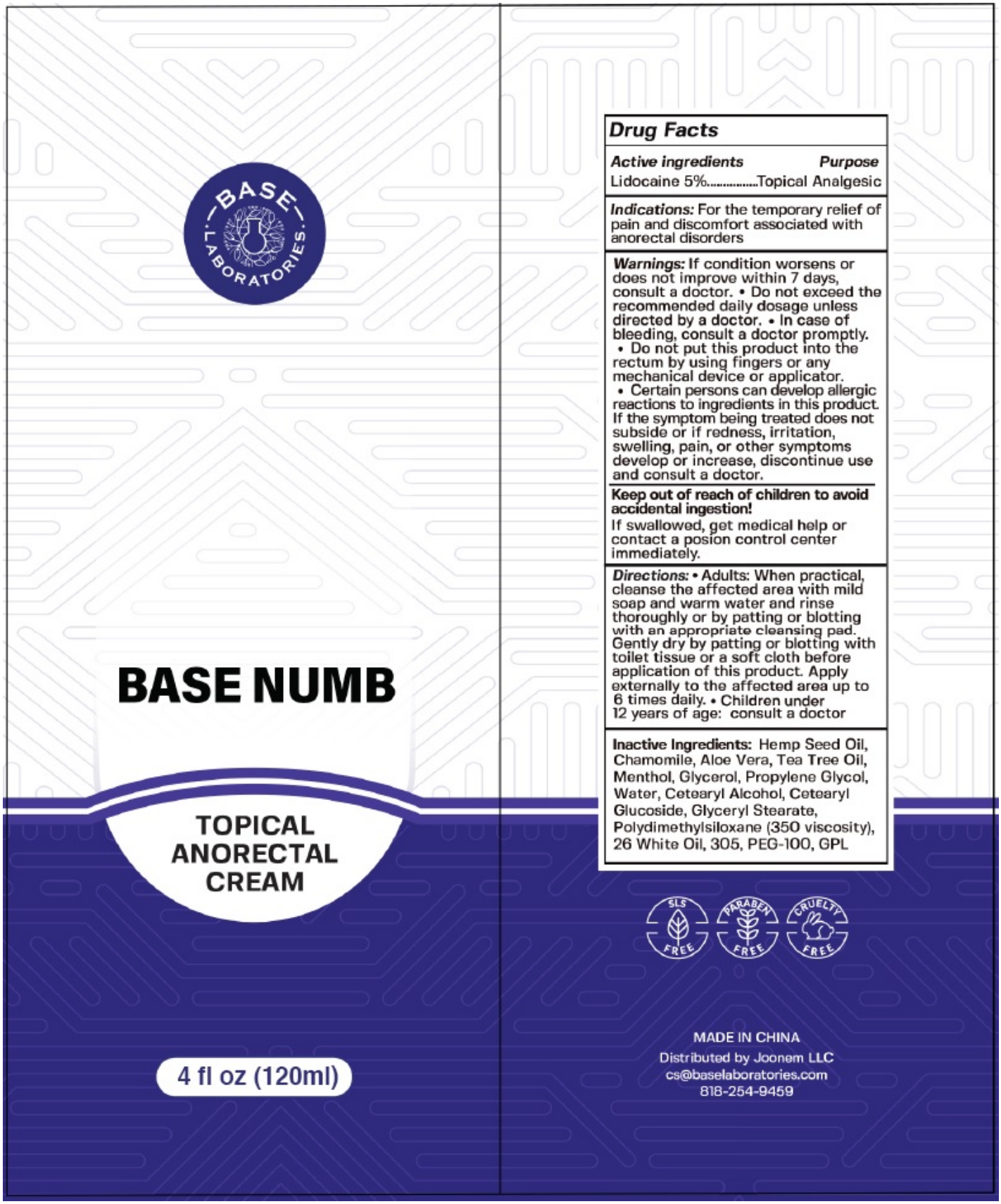
Directions:

- Adults: When practical, cleanse the affected area with mild soap and warm water and rinse thoroughly or by patting or blotting with an appropriate cleansing pad. Gently dry by patting or blotting with toilet tissue or a soft cloth before application of this product. Apply externally to the affected area up to 6 times daily.
- Children under 12 years of age: consult a doctor

Inactive Ingredients:

Hemp Seed Oil, Chamomile, Aloe Vera, Tea Tree Oil, Menthol, Glycerol, Propylene Glycol, Water, Cetearyl Alcohol, Cetearyl Glucoside, Glyceryl Stearate, Polydimethylsiloxane, (350 viscosity), 26 White Oil, 305, PEG-100, GPL

Package Labeling:



BASE LABORATORIES BASE NUMB TOPICAL ANORECTAL CREAM

lidocaine cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80327-008
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 mL

Inactive Ingredients

Ingredient Name	Strength
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
CHAMOMILE (UNII: FGL3685T2X)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TEA TREE OIL (UNII: VIF565UC2G)	
MENTHOL (UNII: L7T10EIP3A)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
POLYETHYLENE GLYCOL 4500 (UNII: TVH7653921)	
LIQUEFIED PETROLEUM GAS (UNII: 5K616HU99V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80327-008-01	1 in 1 CARTON	10/01/2021	
1		120 mL in 1 BOTTLE; 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	10/01/2021	

Labeler - Joonem LLC (117633878)

Revised: 1/2024 Joonem LLC