

LOCAL ANESTHETIC ANORECTAL CREAM- lidocaine cream
Make Labs LLC

Local Anesthetic Anorectal Cream

Active Ingredients purpose

Lidocaine 5%.....Local Anesthetic

Uses

Temporary relief of pain, itching, and discomfort associated with hemorrhoids

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and ask a doctor if

- rectal bleeding occurs
- condition worsens or does not improve within 7 days
- certain persons can develop allergic reactions to ingredients in this product. If the symptom being treated does not subside or if redness, irritation, swelling, pain, or other symptoms develop or increase, discontinue use and consult a doctor.

Warnings

For external use only

If pregnant or breast-feeding, ask a health professional before use.

Directions

- **Adults:** When practical, clean affected area with mild soap and warm water and rinse thoroughly. Gently dry by patting or blotting with toilet tissue or soft cloth before applying. Apply externally to the affected area up to 6 times a day.
- **Children 12 and under,** consult a physician prior to use.

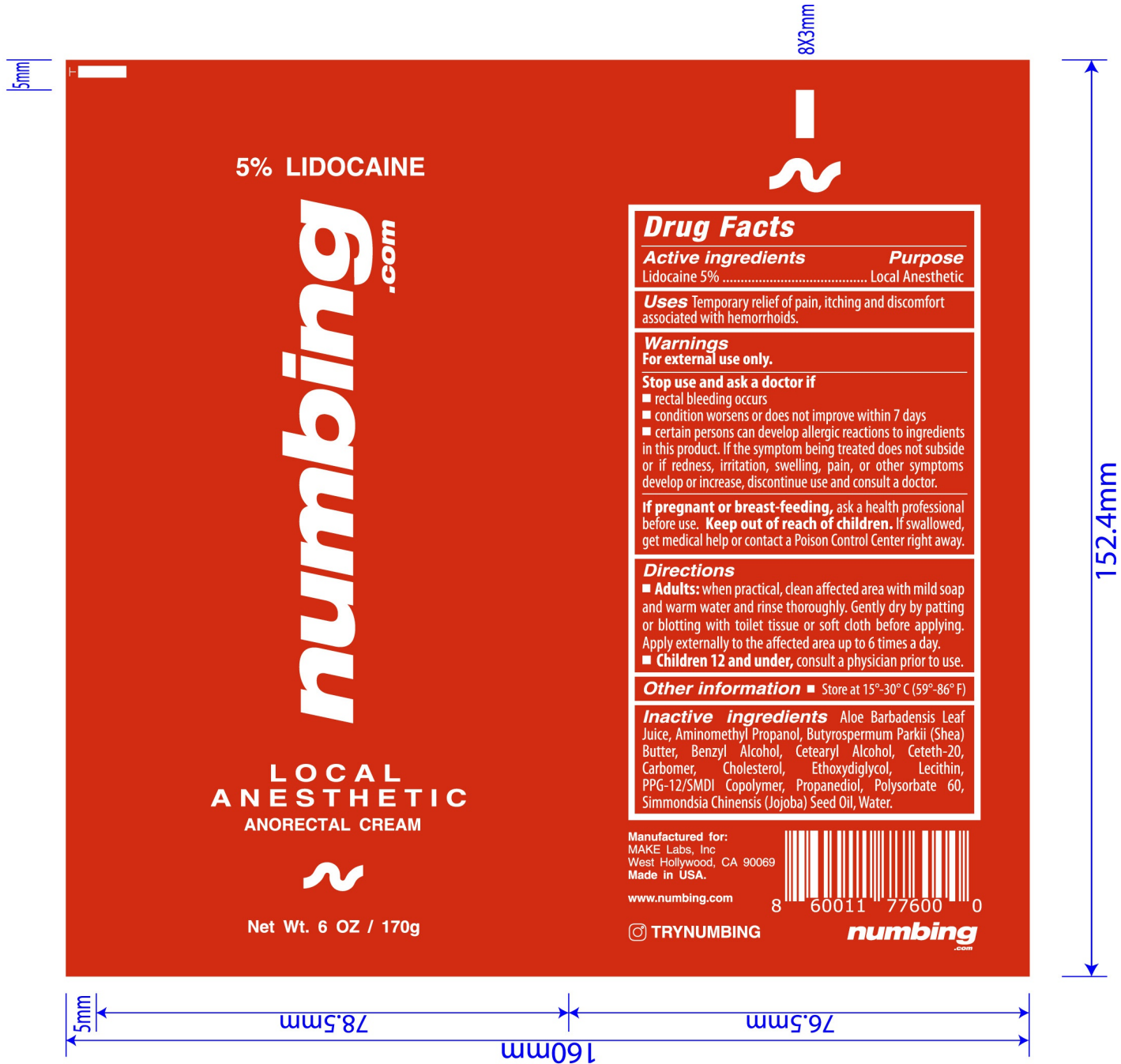
Inactive ingredients Aloe Barbadensis Leaf Juice, Aminomethyl Propanol, Butyrospermum Parkii (Shea) Butter, Benzyl Alcohol, Cetearyl Alcohol, Ceteth-20, Carbomer, Cholesterol, Ethoxydiglycol, Lecithin, PPG-12/SMDI Copolymer, Propanediol, Polysorbate 60, Simmondsia Chinensis (Jojoba) Seed Oil, Water.

numbing.com

Local Anesthetic

Anorectal Cream

Net Wt. 6 OZ / 170g



LOCAL ANESTHETIC ANORECTAL CREAM

lidocaine cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84767-001
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 g

Inactive Ingredients	
Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SHEA BUTTER (UNII: K49155WL9Y)	
CHOLESTEROL (UNII: 97C5T2UQ7J)	
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-20 (UNII: I835H2IHHX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
PPG-12/SMDI COPOLYMER (UNII: 1BK9DDD24E)	
PROPANEDIOL (UNII: 5965N8W85T)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
JOJOBA OIL (UNII: 724GKU717M)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84767-001-01	170 g in 1 TUBE; Type 0: Not a Combination Product	08/01/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	08/01/2024	

Labeler - Make Labs LLC (126167088)