

AEMERRY NUMBING CREAM- menthol 5% cream
Da Capo Prima LLC

Initial Drug Listing - Aemerry Numbing Cream-171

Menthol 5%

Local Anesthetic

Temporarily relieves local and anorectal discomfort associated with the anorectal disorder or swelling.

For external use only

Do not use this product if

- you are allergic to lidocaine.
- in case of accidental overdose, seek immediate medical help or contact the Poison Control Center.
- seal is broken or missing.

When using this product

- do not exceed the recommended daily dosage unless directed by a doctor. Certain sensitive individuals may have an allergic reaction to some of the ingredients of this product.
- do not put this cream into the rectum using your fingers, applicators, or any medical devices.

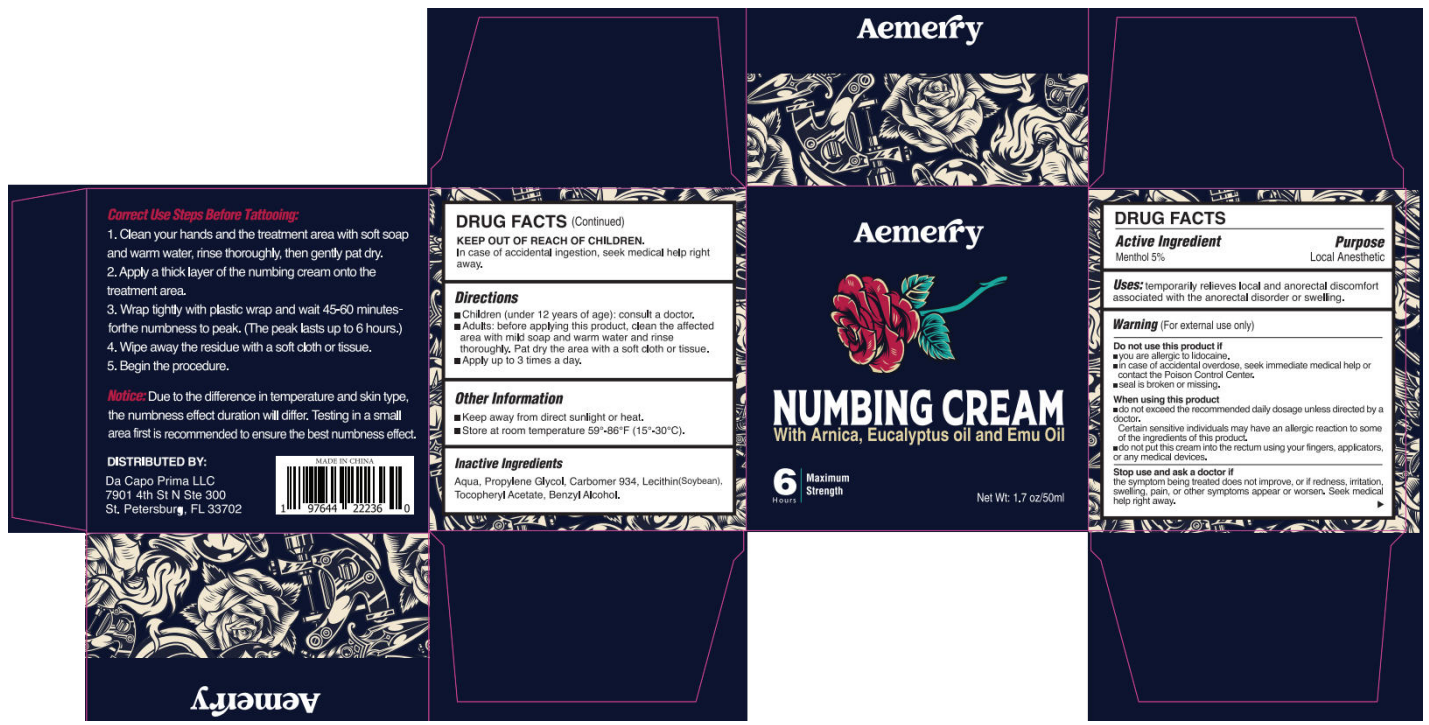
Stop use and ask a doctor if

the symptom being treated does not improve, or if redness, irritation, swelling, pain, or other symptoms appear or worsen. Seek medical help right away.

In case of accidental ingestion, seek medical help right away.

- Children (under 12 years of age): consult a doctor.
- Adults: before applying this product, clean the affected area with mild soap and warm water and rinse thoroughly. Pat dry the area with a soft cloth or tissue.
- Apply up to 3 times a day.
- Keep away from direct sunlight or heat.
- Store at room temperature 59°-86°F (15°-30°C).

Aqua, Propylene Glycol, Carbomer 934, Lecithin (Soybean), Tocopheryl Acetate, Benzyl Alcohol.



AEMERRY NUMBING CREAM

menthol 5% cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87615-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.5 g in 50 g

Inactive Ingredients

Ingredient Name	Strength
AQUA (UNII: 059QF0KOOR)	
CARBOMER 934 (UNII: Z135WT9208)	
LECITHIN, SOYBEAN (UNII: 1D156QDM62)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WP17EW8)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87615-002-01	50 g in 1 CARTON; Type 0: Not a Combination Product	05/12/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/12/2026	

Labeler - Da Capo Prima LLC (144987090)

Revised: 4/2026

Da Capo Prima LLC