

PROINNATC NUMBING CREAM- menthol cream
Guangzhou Haishi Biological Technology Co., Ltd.

PROINNATC NUMBING CREAM

Active ingredients

Menthol 10%

Purpose

For the temporary alleviation of localized skin pain/discomfort, pruritus, or burning sensation.

Uses

- 1, Clean your hands and treatment area thoroughly with mild soap and warm water. Dry them afterward.
- 2, Apply a thick layer of numbing cream to the treatment area, making sure to cover both the tattooed area and its surroundings.
- 3, Wrap the area securely with plastic wrap and leave it on for 40-60 minutes for the best results (keeping it on longer gives better effects).
- 4, Remove any remaining cream and wait an additional 5 minutes for the numbing sensation to reach its peak.

Warnings

For external use only.

Stop use and ask a doctor if

rash occurs.

Do not use

if you are allergic to the ingredients in this product or if the seal is broken or missing.

When using this product

keep out of eyes. Rinse with water to remove.

Keep out of reach of children.

3. Children under 2 years: Do not use

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

Dosage

Apply to affected area not more than 3 times daily.

Inactive ingredients

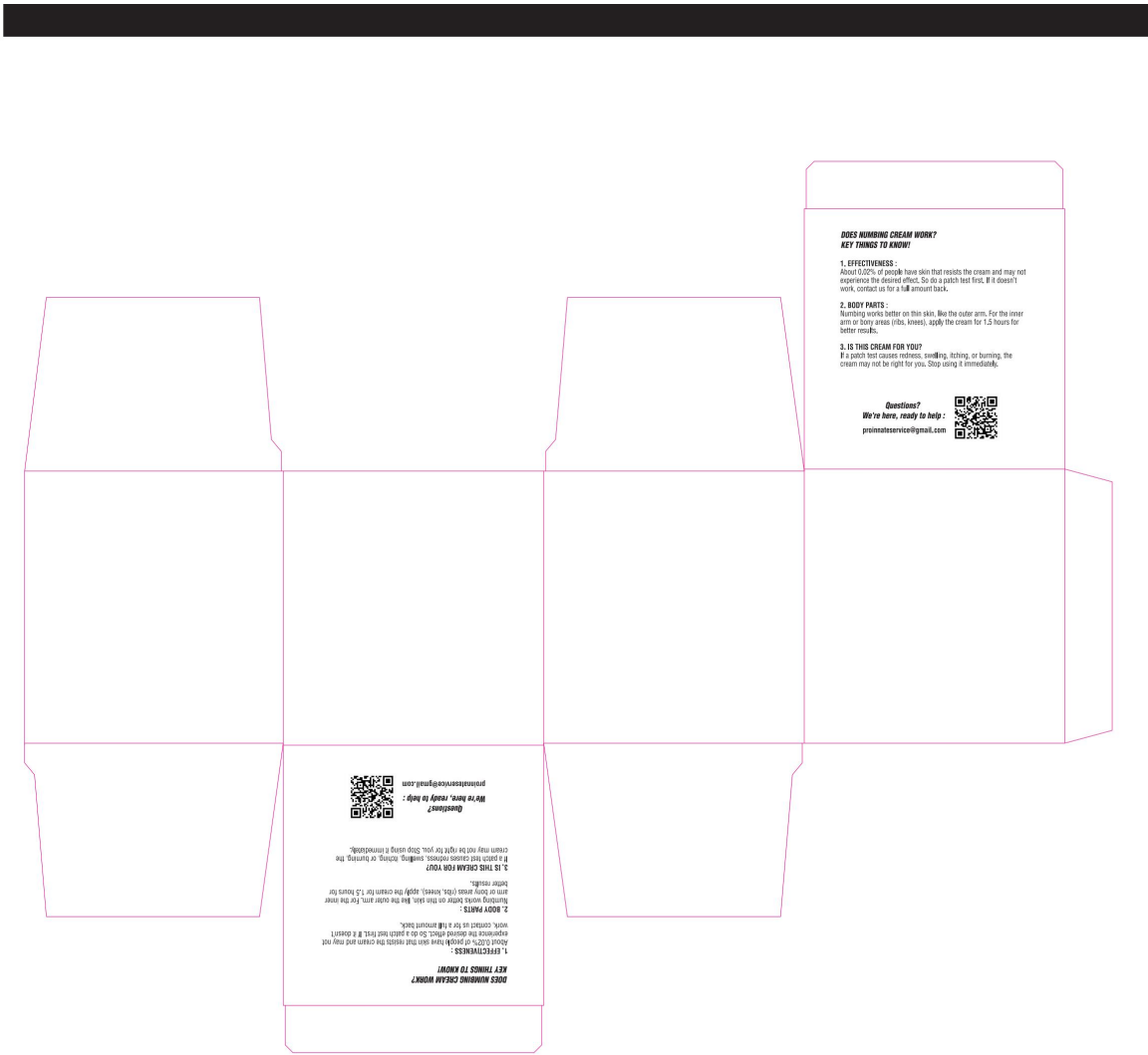
- Aqua 69%
- Mentha Arvensis Leaf Extract, 3%
- Zanthoxylum Bungeanum Fruit Extract, 5%
- Arnica Montana, 3%
- Emu Oil, 2%
- Chrysanthellum Indicum Extract, 1%
- Propylene Glycol, 5%
- Disodium Edta, 1%
- Portulaca Oleracea Extract, 1%



182*38mm



66*66*69mm
含内托



PROINNATC NUMBING CREAM			
menthol cream			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60771-0023
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)		MENTHOL	10 g in 100 g
Inactive Ingredients			
Ingredient Name			Strength

ARNICA MONTANA (UNII: O80TY208ZW)				
EMU OIL (UNII: 344821WD61)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60771-0023-1	120 g in 1 BOTTLE; Type 0: Not a Combination Product	07/08/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M017	07/08/2025	

Labeler - Guangzhou Haishi Biological Technology Co., Ltd. (421262738)

Establishment			
Name	Address	ID/FEI	Business Operations
Guangzhou Haishi Biological Technology Co., Ltd.		421262738	manufacture(60771-0023)