

PUTRIZEN NUMBING CREAM- glycerin cream
Shantou South Moon Biotechnology Co., Ltd.

ARNICA MONTANA FLOWER

For the temporary relief of localized pain, itching, or discomfort from minor skin irritations.

- 1.Clean hands and treatment area with mild soap and warm water, rinse thoroughly,then gently pat dry.
- 2.Apply a thick layer of numbing cream to the treatment area, making sure to cover both the treatment area and its surroundings.
- 3.wrap tightly with, plastic wrap and wait 40-60 minutes (Longer time.batter effect)
- 4.Wipe away residue, and wait another 15-25 minutes for the numbness to peak.

For external use only.

Do not get into the eyes.m Do not exceed recommended dosage unless directed by a doctor. m Do not put this product into the rectum by using fingers or any mechanical device or applicator.

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Keep out of reach of children

Store at room temperature 59°F-86°F(159C-309C).Avoid direct sunlight Do not use if tamper-evident packaging has been breached or is missing!

EUCALYPTUS GLOBULUS LEAF□EMU OIL□AQUA□GLYCERIN□MENTHOL□MINERAL OIL□GLYCERYL MONOSTEARATE□1,2-HEXANEDIOL□CARBOMER HOMOPOLYMER TYPE A□TROLAMINE

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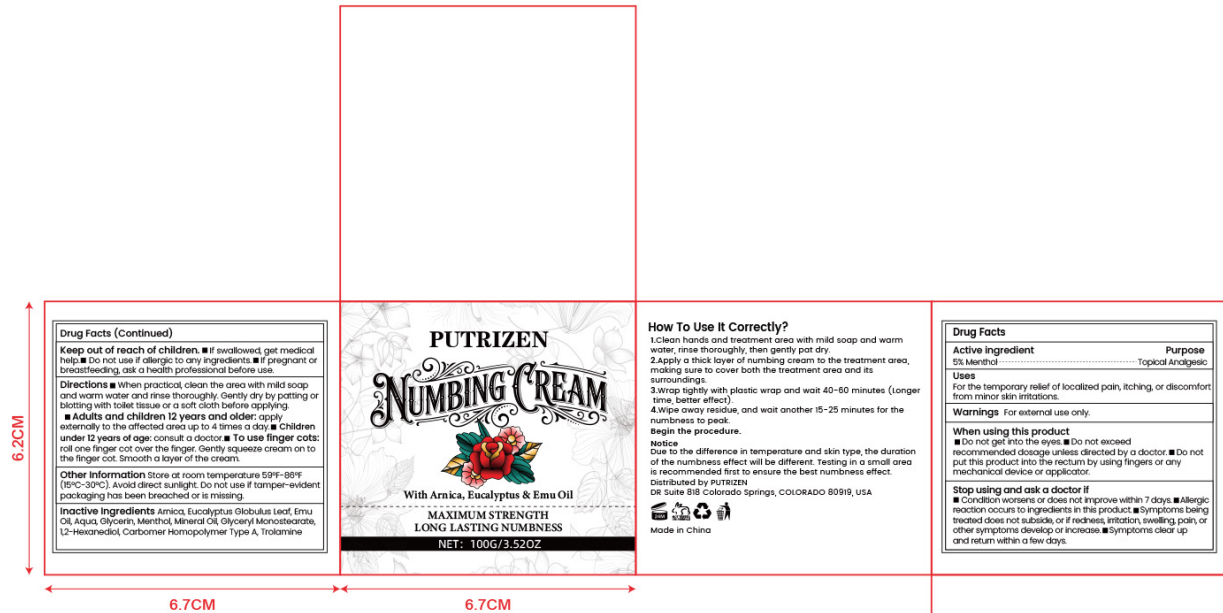
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纸盒 (哑膜)

规格尺寸：长6.7x宽6.7x高6.2cm



PUTRIZEN NUMBING CREAM

glycerin cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:84983-027

Route of Administration	TOPICAL
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Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ) (ARNICA MONTANA FLOWER - UNII:OZ0E5Y15PZ)	ARNICA MONTANA FLOWER	0.01 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	1 mg in 100 mg
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	0.01 mg in 100 mg
EMU OIL (UNII: 344821WD61)	0.01 mg in 100 mg
GLYCERIN (UNII: PDC6A3C0OX)	8 mg in 100 mg
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	0.1 mg in 100 mg
MENTHOL (UNII: L7T10EIP3A)	5 mg in 100 mg
MINERAL OIL (UNII: T5L8T28FGP)	2 mg in 100 mg
CARBOMER HOMOPOLYMER TYPE A (UNII: F68VH75CJC)	0.1 mg in 100 mg
AQUA (UNII: 059QF0KO0R)	83.67 mg in 100 mg
TROLAMINE (UNII: 9O3K93S3TK)	0.1 mg in 100 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84983-027-01	100 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/09/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	04/09/2026	

Labeler - Shantou South Moon Biotechnology Co., Ltd. (457126192)

Establishment

Name	Address	ID/FEI	Business Operations
Shantou South Moon Biotechnology Co., Ltd.		457126192	manufacture(84983-027)

Revised: 4/2026

Shantou South Moon Biotechnology Co., Ltd.