ARTHCAL SOOTHING PAIN RELIEF MASSAGE- soothing pain relief massage gel
Beijing JUNGE Technology Co., Ltd.

100g(3.53oz)

EMU OIL
MENTHOL

ARNICA MONTANA

TURMERIC

EUCALYPTUS OIL
CINNAMON OIL
GLYCERIN
ALCOHOL
ISOPROPYLAL COHOL
AQUA

Wash and dry the area to be applied.
Massage gently until absorbed into the skin.
Adult dosage: apply to affected area not more than 3 to 4 times daily.
For children under 14 years of age: Consult a doct

Topical Analgesic

Keep out of reach of children

For external use only
Do not apply to wounds or damaged skin
Do not bandage tightly .Avoid contact with eyes
Keep out of reach of children
If swallowed, get medical help or contact a Poison Control Center immediately.
lf condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within afew days, discontinue use of this product and consult a physician
lf pregnant or breast feeding, ask a health professional before use

Store in a clean, dry placeoulside of direct sunlight
Protect product fromexcessive moisture

ARTHCAL SOOTHING PAIN RELIEF MASSAGE

soothing pain relief massage gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85212-0039
Route of Administration	CUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
EMU OIL (UNII: 344821WD61) (EMU OIL - UNII:344821WD61)	EMU OIL	1 g in 100 g
TURMERIC (UNII: 856YO1Z64F) (TURMERIC - UNII:856YO1Z64F)	TURMERIC	1 g in 100 g
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	2 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	3 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
CINNAMON OIL (UNII: E5GY4I6YCZ)	0.5 g in 100 g

AQUA (UNII: 059QF0KO0R)	44.5 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX)	1 g in 100 g
ISOPROPYL ALCOHOL (UNII: ND2M416302)	16.2 g in 100 g
ALCOHOL (UNII: 3K9958V90M)	30 g in 100 g
EUCALYPTUS OIL (UNII: 2R04ONI662)	0.8 g in 100 g

Product Characteristics

Color		**Score**	
Shape	OVAL	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85212-0039-1	1 g in 1 BOX; Type 0: Not a Combination Product	01/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2025	

Labeler - Beijing JUNGE Technology Co., Ltd. (848718652)

Registrant - Beijing JUNGE Technology Co., Ltd. (848718652)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Ayan Biotech Co., Ltd.		544377996	manufacture(85212-0039)

Revised: 8/2025 Beijing JUNGE Technology Co., Ltd.