

**SOUTH MOON MELITTIN JOINT SOOTHING GEL- south moon melittin joint soothing gel gel**  
**Dongguan Haiyi Technology Co.,Ltd.**

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**Active ingredient**

BEE VENOM

GLUCOSAMINE

**Inactive ingredient**

AQUA,CURCUMIN,EICOSAPENTAENOIC ACID,CHOLE-CALCIFEROL

**Purpose section**

- 1.Solve various joint problems
- 2.passed the USA FDAapproved
- 3.Effectively relieves arthritis pain
- 4.Does not contain anyharmful ingredients.

**Warning**

Please keep out of reach of children. Do not swallow. Please clean your hands before use to ensure the best results from the product. Discontinue use if signs of irritation or rash occur. Store in a cool and dry place.

**stop use**

This product may cause allergic reactions in a small number of people. If you experience any discomfort, please stop using it immediately.

**not use**

Not suitable for use by children, pregnant or breast feeding people.

**OUT OF CHILDREN**

KEEP THE PRODUCT OUT OF REACH OF CHILDREN to

**HOW TO USE**

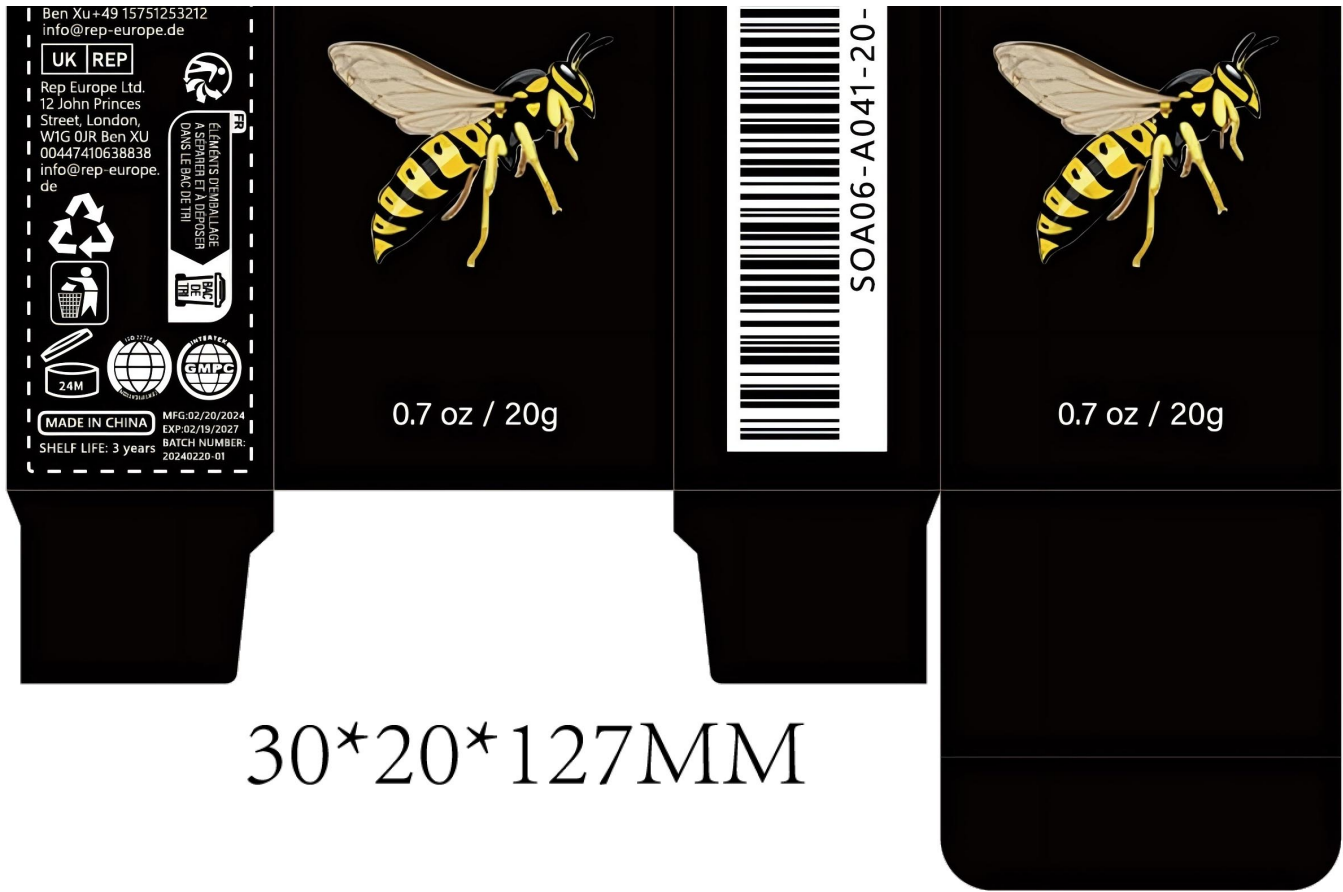
1. Clean and dry the skin on the joints.

2. Take an appropriate amount of gel and apply it evenly on the joints.
3. Gently massage with hands in circular motion until absorbed.
4. Daily use can get better results.

## Dosage

take an appropriate amount, Use 2-3 times a week





## SOUTH MOON MELLITTIN JOINT SOOTHING GEL

south moon melittin joint soothing gel gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:84732-055
<b>Route of Administration</b>	CUTANEOUS		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>APIS MELLIFERA VENOM</b> (UNII: 76013O881M) (APIS MELLIFERA VENOM - UNII:76013O881M)	APIS MELLIFERA VENOM	1 mg in 20 g
<b>GLUCOSAMINE</b> (UNII: N08U5BOQ1K) (GLUCOSAMINE - UNII:N08U5BOQ1K)	GLUCOSAMINE	1 mg in 20 g

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CURCUMIN</b> (UNII: IT942ZTH98)	
<b>CHOLECALCIFEROL</b> (UNII: 1C6V77QF41)	
<b>ICOSAPENT</b> (UNII: AAN7QOV9EA)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84732-055-01	20 g in 1 TUBE; Type 0: Not a Combination Product	10/21/2024	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	10/21/2024	

**Labeler** - Dongguan Haiyi Technology Co.,Ltd. (722030807)

## Establishment

Name	Address	ID/FEI	Business Operations
Dongguan Haiyi Technology Co.,Ltd.		722030807	manufacture(84732-055)

Revised: 10/2024

Dongguan Haiyi Technology Co.,Ltd.