XIMONTH BEE VENOM PAIN CREAM- allantoin cream
Guangdong Ximonth Technology Co., Ltd.

Active Ingredient(s)

ARNICA CHAMISSONIS FLOWER EXTRACT□GLUCOSAMINE□SODIUM BUTYROYL/FORMOYL CHONDROITIN SULFATE□ MENAQUINONE-7□DIMETHYL SULFONE

Purpose

Absorbed through the skin, a variety of plant ingredients penetrate into the joints, dissolve uric acid crystals, soothe swelling, and reduce pain and discomfort.

Use

1. Clean and dry the affected area of skin.
2. Take an appropriate amount of cream and apply it evenly on the skin.
3. Massage gently with your hands in circular motions for absorption.
4. Use it 2-3 times a day for better results.

Warnings

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.

Do not use

Discontinue use if signs of irritation or rash occur.

STOP USE

Discontinue use if signs of irritation or rash occur.

Please keep out of reach of children. Do not swallow.

Avoid freezing and excessive heat above 40C (104F) □
Store in a cool and dry place.

ALLANTOIN

纸盒

规格尺寸：长4.3x4.3x4cm

XIMONTH BEE VENOM PAIN CREAM

allantoin cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84660-033
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

GLUCOSAMINE (UNII: N08U5BOQ1K) (GLUCOSAMINE - UNII:N08U5BOQ1K)	GLUCOSAMINE	4.5 g in 30 g
MENAQUINONE 7 (UNII: 8427BML8NY) (MENAQUINONE 7 - UNII:8427BML8NY)	MENAQUINONE 7	9 g in 30 g
ARNICA CHAMISSONIS FLOWER (UNII: 88WK5I8R3L) (ARNICA CHAMISSONIS FLOWER - UNII:88WK5I8R3L)	ARNICA CHAMISSONIS FLOWER	3 g in 30 g
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT) (DIMETHYL SULFONE - UNII:9H4PO4Z4FT)	DIMETHYL SULFONE	3 g in 30 g

Inactive Ingredients

Ingredient Name	Strength
ALLANTOIN (UNII: 344S277G0Z)	6 g in 30 g
SODIUM BUTYROYL/FORMOYL HYALURONATE (UNII: F63GOY24UQ)	4.5 g in 30 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84660-033-01	30 g in 1 BOX; Type 0: Not a Combination Product	09/26/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	09/26/2024	

Labeler - Guangdong Ximonth Technology Co., Ltd. (699436264)

Registrant - Guangdong Ximonth Technology Co., Ltd. (699436264)

Establishment

Name	Address	ID/FEI	Business Operations
Guangdong Ximonth Technology Co., Ltd.		699436264	manufacture(84660-033)

Revised: 9/2024 Guangdong Ximonth Technology Co., Ltd.