JOINT BONE CARE- collagen cream
Guangdong Ximonth Technology Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Active Ingredient(s)

COLLAGEN □MENAQUINONE-7 □ARNICA MONTANA FLOWER EXTRACT□SODIUM CHONDROITIN SULFATE

Purpose

Relieve joint&bone pain

Use

1. Clean and dry the affected skin. 2. Take an appropriate amount of this product and apply it evenly on the skin of knees, shoulders, back joint pain areas or other bone and muscle pain areas. 3. Gently massage and absorb evenly with your hands in circular motions.

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

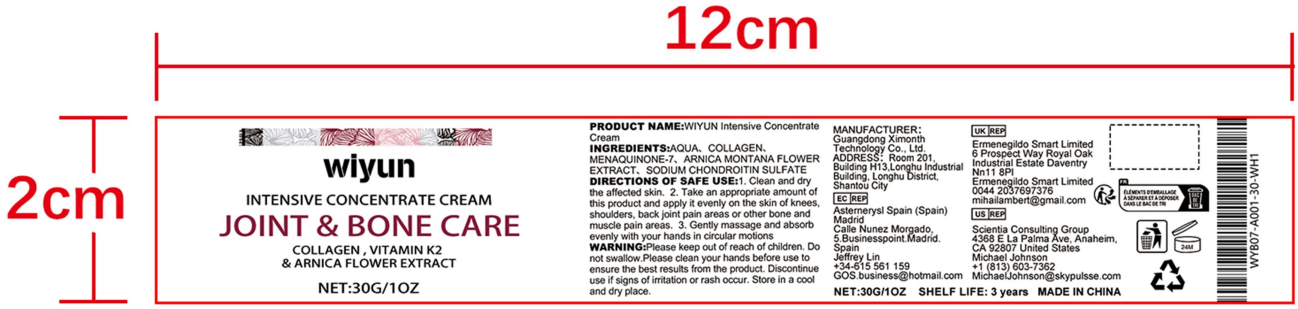
Discontinue use if signs of irritation or rash occur.

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.

AQUA



JOINT BONE CARE

collagen cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84660-001
Route of Administration	TOPICAL		

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

MENAQUINONE 7 (UNII: 8427BML8NY) (MENAQUINONE 7 - UNII:8427BML8NY)	MENAQUINONE 7	0.05 g in 100 g
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ) (ARNICA MONTANA FLOWER - UNII:OZ0E5Y15PZ)	ARNICA MONTANA FLOWER	1 g in 100 g
MARINE COLLAGEN, SOLUBLE (UNII: 8JC99XGU4W) (MARINE COLLAGEN, SOLUBLE - UNII:8JC99XGU4W)	MARINE COLLAGEN, SOLUBLE	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHONDROITIN SULFATE (PORCINE; 5500 MW) (UNII: H5BJH23Z9A)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84660-001-01	30 g in 1 BOTTLE; Type 0: Not a Combination Product	08/15/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/15/2024	

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(84660-001)

Revised: 8/2024 Guangdong Ximonth Technology Co., Ltd.