

BEE VENOM PSORIASIS TREATMENT- apis mellifera venom cream
Guangzhou Houkea Biotechnology 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 Ingredients

APIS MELLIFERA VENOM,LICORICE,ALOE VERA LEAF

Purpose

Supports skin health
Long-lasting hydration
Fast itch relief

use

Long-lasting soothing and moisturizing for skin; effectively relieves skin redness and itching; eliminates tinea pedis and quickly restores skin health

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.

NONE

NONE

none

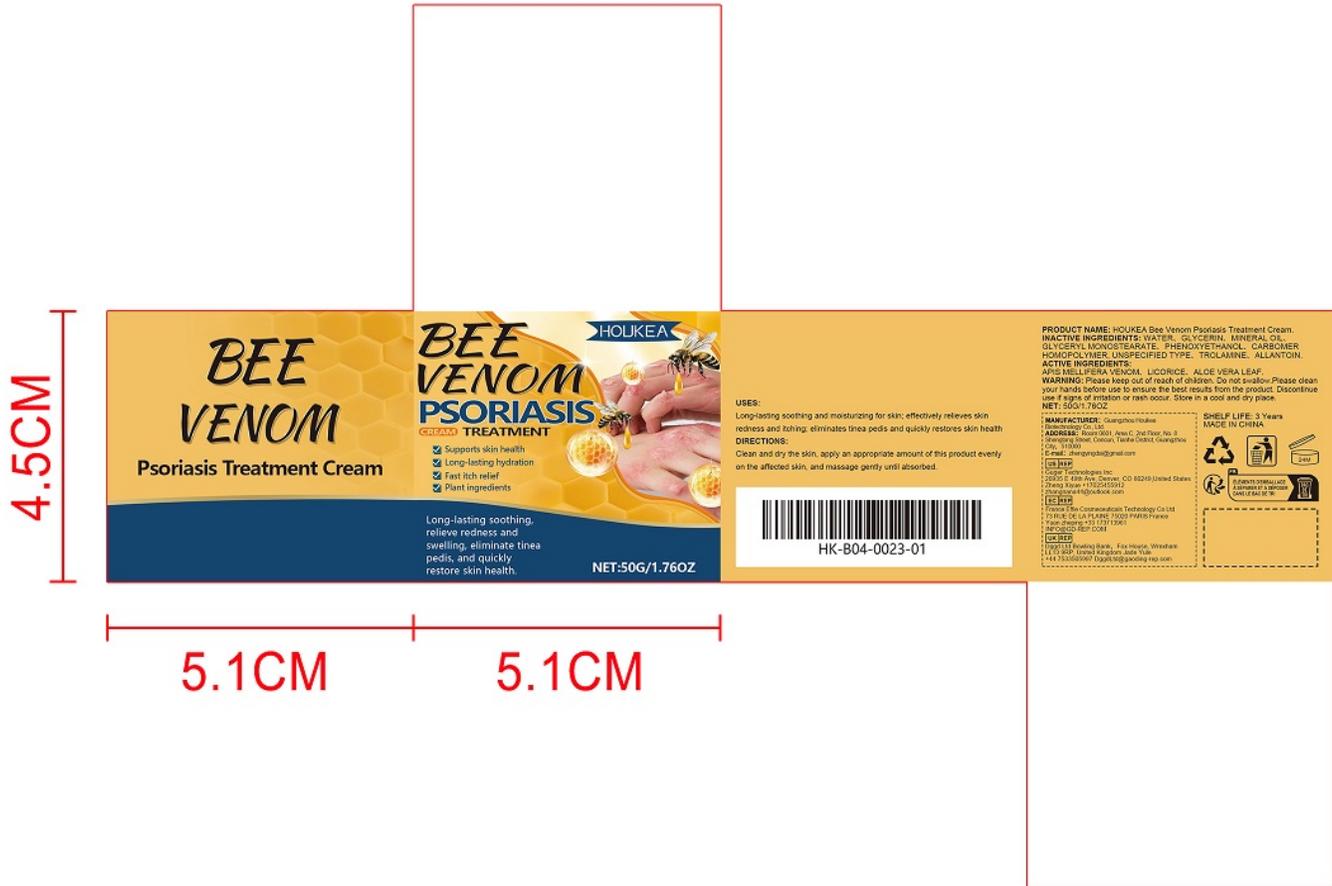
Please keep out of reach of children

Clean and dry the skin, apply an appropriate amount of this product evenly on the affected skin, and massage gently until absorbed.

WATER □ GLYCERIN □ MINERAL OIL □ GLYCERYL
MONOSTEARATE □ PHENOXYETHANOL □ CARBOMER HOMOPOLYMER, UNSPECIFIED
TYPE □ TROLAMINE □ ALLANTOIN

纸盒规格

尺寸:长5.1x宽5.1x高4.5cm



BEE VENOM PSORIASIS TREATMENT

apis mellifera venom cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84984-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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APIS MELLIFERA VENOM (UNII: 76013O881M) (APIS MELLIFERA VENOM - UNII:76013O881M)	APIS MELLIFERA VENOM	20 g in 100 g
LICORICE (UNII: 61ZBX54883) (LICORICE - UNII:61ZBX54883)	LICORICE	7 g in 100 g
ALOE VERA LEAF (UNII: ZY81Z83H0X) (ALOE VERA LEAF - UNII:ZY81Z83H0X)	ALOE VERA LEAF	3 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALLANTOIN (UNII: 344S277G0Z)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
MINERAL OIL (UNII: T5L8T28FGP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84984-001-01	1 in 1 BOX	12/02/2024	
1		50 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/02/2024	

Labeler - Guangzhou Houkea Biotechnology Co., Ltd. (456717258)

Registrant - Guangzhou Houkea Biotechnology Co., Ltd. (456717258)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Houkea Biotechnology Co., Ltd.		456717258	manufacture(84984-001)

Revised: 11/2025

Guangzhou Houkea Biotechnology Co., Ltd.