

HOUKEA RAPID MOLE SPOT ELIMINATION GEL- glycerin gel
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](#).

AVOCADO OIL,
LAMINARIA JAPONICA,
CHONDRUS CRISPUS CARRAGEENAN,
CHAMOMILEIN

Clean and dry your skin, apply an appropriate amount of this product evenly on the affected area with skin problems such as spots, and gently massage until absorbed.

Dissolves skin acne; removes moles; lightens dark spots; reduces skin blemishes; makes skin smoother; restores skin health

Discontinue use if signs of irritation or rash occur.

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

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.

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WATER,
GLYCERIN,
CARBOMER HOMOPOLYMER TYPE A□
TROLAMINE□
ALLANTOIN

纸盒
规格尺寸：长5.2×宽5.2×高4.4cm



HOUKEA RAPID MOLE SPOT ELIMINATION GEL

glycerin gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84984-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LAMINARIA JAPONICA (UNII: WE98HW412B) (LAMINARIA JAPONICA -	LAMINARIA JAPONICA	0.01 mg

UNII:WE98HW412B)	LAMINARIA JAPONICA	in 50 mg
CHAMOMILE (UNII: FGL3685T2X) (CHAMOMILE - UNII:FGL3685T2X)	CHAMOMILE	0.01 mg in 50 mg
CHONDRUS CRISPUS CARRAGEENAN (UNII: UE856F2T78) (CHONDRUS CRISPUS CARRAGEENAN - UNII:UE856F2T78)	CHONDRUS CRISPUS CARRAGEENAN	0.01 mg in 50 mg
AVOCADO OIL (UNII: 6VNO72PFC1) (AVOCADO OIL - UNII:6VNO72PFC1)	AVOCADO OIL	0.01 mg in 50 mg

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER TYPE A (UNII: F68VH75CJC)	0.2 mg in 50 mg
ALLANTOIN (UNII: 344S277G0Z)	0.01 mg in 50 mg
WATER (UNII: 059QF0KO0R)	45.75 mg in 50 mg
GLYCERIN (UNII: PDC6A3C0OX)	4 mg in 50 mg
TROLAMINE (UNII: 9O3K93S3TK)	0.1 mg in 50 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84984-002-01	50 mg in 1 BOX; Type 0: Not a Combination Product	04/02/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/02/2025	

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(84984-002)

Revised: 4/2025

Guangzhou Houkea Biotechnology Co., Ltd.