

KOPA HAIKU ANTIFUNGAL CREAM- miconazole nitrate 2% cream

Trovina Co. Ltd

Miconazole Nitrate 2%

Antifungal

Anti-fungal for the effective treatment of Athlete's Foot, Jock Itch, Ringworm, Eczema, Tinea Versicolor, and fungal folliculitis.

For external use only

On children under 3 years of age.

Avoid contact with the eyes.

Irritation occurs or if there is no improvement within 4 weeks.

Keep out of reach of children. In case of ingestion, seek medical help or contact a Poison Control Center immediately.

1 Apply a small amount of cream to the affected area 2-3 times daily; 2 Ensure the skin in clean and dry before use; 3 For first-time users, perform a patch test on a small area of skin to check for sensitivity.

Aqua, Stemona Japonica Root Extract, Dictamnus Dasycarpus Root Bark Extract, Sophora Flavescens Root Extract, Aloe Barbadensis Leaf Extract, Cnidium Monnieri Fruit Extract, Tocopherol, Borneol, Menthol, Stearic Acid, Petrolatum, Paraffinum Liquidum, Glycerin, Triethanolamine

Store in a cool, dry place.

Email at: service@kopahaiku.co.uk



KOPA HAIKU ANTIFUNGAL CREAM

miconazole nitrate 2% cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87041-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)	
PARAFFINUM LIQUIDUM (UNII: T5L8T28FGP)	
DICTAMNUS DASYCARPUS ROOT BARK (UNII: LA97176ILS)	
TOCOPHEROL (UNII: R0ZB2556P8)	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K)	
GLYCERIN (UNII: PDC6A3C0OX)	
CNIDIUM MONNIERI FRUIT (UNII: V1A3S3CUS)	
BORNEOL (UNII: M89NIB437X)	
PETROLATUM (UNII: 4T6H12BN9U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIETHANOLAMINE (UNII: 9O3K93S3TK)	
STEMONA JAPONICA ROOT (UNII: FXG254HF10)	
MENTHOL (UNII: L7T10EIP3A)	
AQUA (UNII: 059QF0KOOR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87041-002-01	30 g in 1 CARTON; Type 0: Not a Combination Product	11/19/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	11/19/2025	

Labeler - Trovina Co. Ltd (233813758)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Zhongletang Pharmaceuticals Co., Ltd.		411295811	manufacture(87041-002)

Revised: 2/2026

Trovina Co. Ltd