

LANIMPRESS1%KETOCONAZOLEANTIDANDRUFFSHAMPOO-
lanimpres1%ketoconazoleantidandruffshampoo lotion
Shenzhen Zhumeng Times Technology Co., Ltd.

Active ingredients

Ketoconazole 1% Anti-dandruff shampoo

Purpose

controls flaking, scaling and itching associated with dandruff

Uses

wet hair thoroughly

apply shampoo, generously lather, rinse thoroughly. Repeat

use every 3-4 days for up to 8 weeks or as directed by a doctor. Then use only as needed to control dandruff

Warnings

For external use only

Do not use

on scalp that is broken or inflamed

if you are allergic to ingredients in this product

Dosage and administration

For external use only.

Do not use

Stop use and ask a doctor if rash occurs

When using section

rash appears

condition worsens or does not improve in 2-4 weeks.

stop use

Do not use

on scalp that is broken or inflamed
if you are allergic to ingredients in this product

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Inactive ingredients

benzyl alcohol, BHT, blue 1, citric acid, cocamide MEA, fragrance, glycol distearate, hydrochloric acid, hydroxypropyl methylcellulose, polyquaternium-7, sodium chloride, sodium cocoyl sarcosinate, sodium hydroxide, sodium lauryl sulfate, tetrasodium EDTA, water, CAPRYLATE, CARBOMER, METHYLPARABEN, TRIETHANOLAMINE, SODIUM HYALURONATE, DISODIUM EDTA, water



LANIMPRESS™

Anti-Dandruff
KETOCONAZOLE 1%
ANTI-DANDRUFF SHAMPOO



**Anti-Dandruff
Shampoo**

CLINICALLY PROVEN
to control flaking, scaling and
itching from dandruff

CONTROLS
fungus that can cause dandruff

CLEAN
fresh scent

4.22fl.oz(125mL)

LANIMPRESS™

1%KETOCONAZOLE
Anti-Dandruff Shampoo

Drug Facts

Active ingredient	Purpose
Ketoconazole 1%	Anti-dandruff shampoo

Uses controls flaking, scaling and itching associated with dandruff

Warnings

For external use only

Do not use

- on scalp that is broken or inflamed
- if you are allergic to ingredients in this product

When using this product

- avoid contact with eyes
- if product gets into eyes, rinse thoroughly with water

Stop use and ask a doctor if

- rash appears
- condition worsens or does not improve in 2-4 weeks

If pregnant or breast-feeding, ask a doctor before use.
Keep out of reach of children. If swallowed, get medical help or
contact a Poison Control Center right away.

Directions

adults and children 12 years and over	<ul style="list-style-type: none"> ■ wet hair thoroughly ■ apply shampoo, generously lather, rinse thoroughly. Repeat ■ use every 3-4 days for up to 8 weeks or as directed by a doctor. Then use only as needed to control dandruff
children under 12 years	<ul style="list-style-type: none"> ■ ask a doctor

Other information

- store at 20°C to 25°C (68°F-77°F)

Inactive ingredients benzyl alcohol, BHT, blue 1, citric acid, cocamide MEA, fragrance, glycol distearate, hydrochloric acid, hydroxypropyl methylcellulose, polyquaternium-7, sodium chloride, sodium cocoyl sarcosinate, sodium hydroxide, sodiumlaureth sulfate, tetrasodium EDTA, water

Questions or comments?

call 1-818-579-7288

DISTRIBUTED BY:

Shenzhen Zhumeng Times Technology Co., Ltd.

ADDRESS:

102, No. 11, Yunlin District, Yabian Community, Shajing Street, Baoan District, Shenzhen, guangdong 518100

MFG:2024/08/30

EXP:2027/08/29

Made in CHINA



LANIMPRESS1%KETOCONAZOLEANTIDANDRUFFSHAMPOO

lanimpres1%ketoconazoleantidandruffshampoo lotion

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:84372-037

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
KETOCONAZOLE (UNII: R9400W927I) (KETOCONAZOLE - UNII:R9400W927I)	KETOCONAZOLE	5 g in 125 mL

Inactive Ingredients

Ingredient Name	Strength
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600 KD) (UNII: 0L414VCS5Y)	
TROLAMINE (UNII: 9O3K93S3TK)	
EDETATE SODIUM (UNII: MP1J8420LU)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM COCOYL SARCOSINATE (UNII: 1R9DUY89CZ)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
WATER (UNII: 059QF0KO0R)	
IVERMECTIN (UNII: 8883YP2R6D)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
8-PENTADECANONE (UNII: QO7H8D8OSB)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ACONITIC ACID (UNII: 93371T1BXP)	
FRAGRANCE 13576 (UNII: 5EM498GW35)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84372-037-01	125 mL in 1 BOX; Type 0: Not a Combination Product	09/02/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	09/02/2024	

Labeler - Shenzhen Zhumeng Times Technology Co., Ltd. (631852731)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Zhumeng Times Technology Co., Ltd.		631852731	manufacture(84372-037)

Revised: 9/2024

Shenzhen Zhumeng Times Technology Co., Ltd.