PUVIDA FUNGAL NAIL- tolnaftate 1% fungal nail patch Jiangxi Hemei Pharmaceutical Co., Ltd

84010-158 update pack

Active Ingredient

Tolnaftate 1%

Purpose

Anti-Funga

Use

Repairs fungal nail damage and improves nail appearance

Warnings

For external use only.

Do not use

Do not use if you are allergic to any of the ingredients Do not use if you are pregnant or breastfeeding

When Using

Avoid direct contact with eyes, mouth, mucous membrane, wounds

Stop Use

■if irritation occurs and discomfort persists

Ask Doctor

if you have diabetes or poor blood circulation

Keep Oot Of Reach Of Children

In case of contact with eyes, flush with water for at least 15 minutes. If swallowed, seek medical assistance immediately or contact a Poison Control Center.

Directions

Clean and thoroughly dry the affected area before application. ■Peel off the patch's protective film, starting with the outer side and then the inner side. ■Place the sticky side of the patch directly onto the infected area on the nail and remove the release paper. Gently wrap the sides of the toe or finger to ensure a snug fit. ■Fold the remaining adhesive part to cover the toe/fingertip. Leave the patch onfor 6-8 hours (It is recommended to use overnight for 3-6 months).

Other information

Store at room temperature and keep away from direct sunlight. Skin discoloration may occur during or after use.

Inactive ingredients

Water, Polyester Spandex, Acrylate Copolymer, Tackifying Resin, Glycerin, Silicone Oil Paper

PRINCIPAL DISPLAY PANEL



PUVIDA FUNGAL NAIL

tolnaftate 1% fungal nail patch

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:84010-158

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TOLNIATTATE (LINIII, OCKDOSOTION /TOLNIATTATE LINIII, OCKDOSOTION	TOLNIATTATE	1 - :- 100

TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV) TOLNAFTATE 1 g in 100

Inactive Ingredients

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Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DICLORAN (UNII: F0BE9UC5J7)	
ACRYLIC ACID (UNII: J94PBK7X8S)	
GLYCERIN (UNII: PDC6A3C0OX)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
POLYESTER-7 (UNII: 0841698D2F)	

Packaging

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	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:84010-158- 01	32 in 1 BOX; Type 0: Not a Combination Product	07/22/2025	

Marketing Information

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	07/22/2025	

Labeler - Jiangxi Hemei Pharmaceutical Co., Ltd (724892056)

Establishment				
Name	Address	ID/FEI	Business Operations	
liangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-158)	

Revised: 7/2025 Jiangxi Hemei Pharmaceutical Co., Ltd