TOENAIL FUNGUS TREATMENT MAXIMUM STRENGTH LIQUID- tolnaftate liquid
Shenzhen Ctrip Technology Co., Ltd.

84614-002 TOENAIL FUNGUS TREATMENT MAXIMUM STRENGTH Liquid

TOENAIL FUNGUS TREATMENT MAXIMUM STRENGTH Liquid

Tolnaftate 1%

Anti-fungal and Nail Renewal

Antifungal use on fingemails, toenails, and the immediately adjacent skin.
Fungal nails, Athlete's Foot, Tinea Manuum, Ringwomcaused byfungus and bacterial infection.
For all caused by nail fungus, nail discoloration, nail thickening,nailsplitting, nail crumbling

Keep away from fire and flame.
Keep out of reach of children.
For use on nails and the immediately adjacent skin only.

If a reaction suggesting sensitivity or iritation occurs.

Not for ophthalmic, oral, or intravaginal use

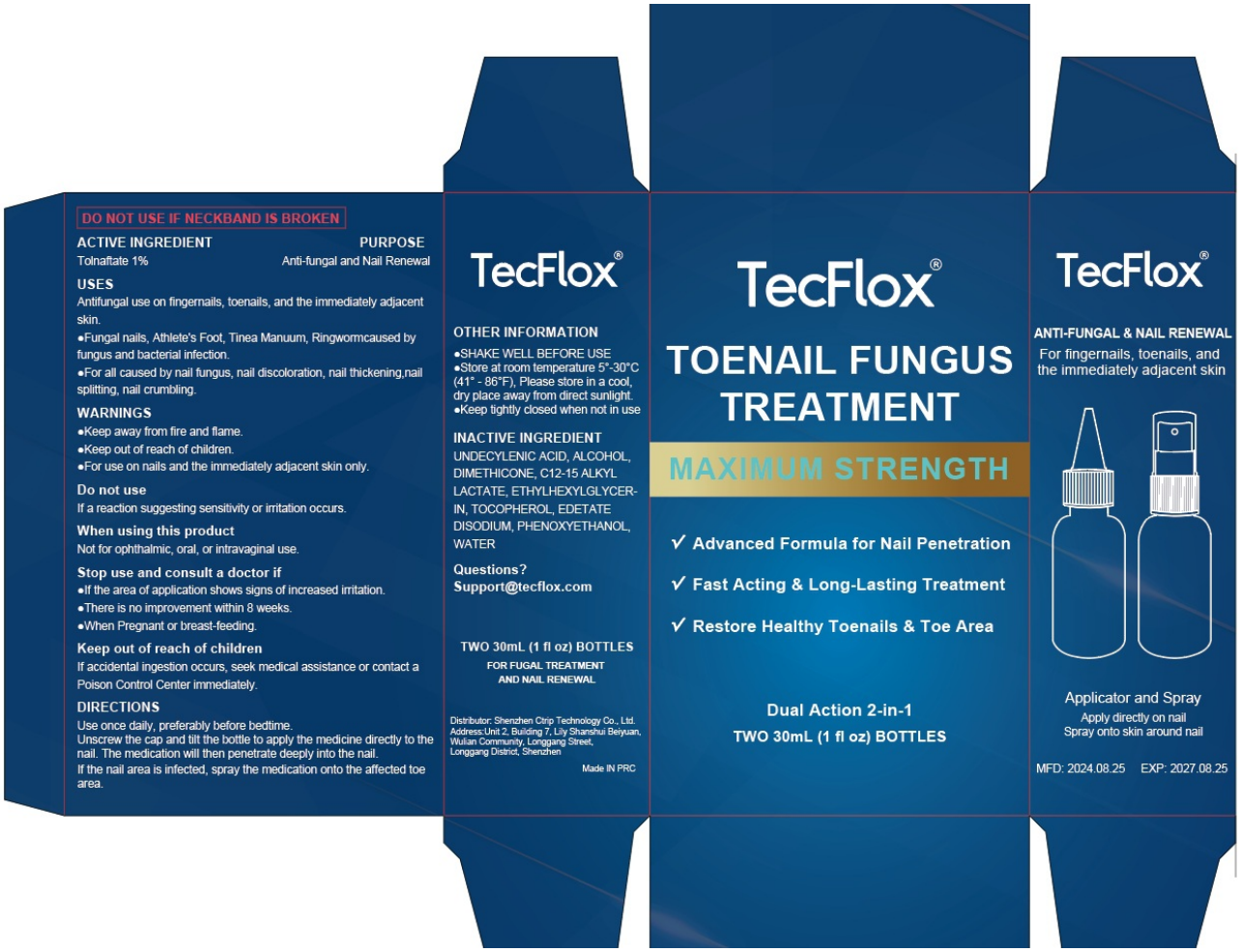
If the area of application shows signs of increased iritation.
There is no improvement within 8 weeks.
When Pregnant or breast-feeding

If accidental ingestion occurs, seek medical assistance or contact a Poison Control Center immediately.

Use once daily, preferably before bedtimeUnscrew the cap and tilt the bottle to apply the medicine directly to thenail. The medication will then penetrate deeply into the nail.If the nail area is infected, spray the medication onto the affected toearea.

SHAKE WELL BEFORE USE
Store at room temperature 5--30°C(41°-86°F), Please store in a cool,dry place away from direct sunlight.
Keep tightly closed when not in use

UNDECYLENIC ACID
ALCOHOL
DIMETHICONE
C12-15 ALKYL LACTATE
ETHYLHEXYLGLYCERIN
TOCOPHEROL
EDETATE DISODIUM
PHENOXYETHANOL
WATER



TOENAIL FUNGUS TREATMENT MAXIMUM STRENGTH LIQUID

tolnaftate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84614-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	1 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
UNDECYLENIC ACID (UNII: K3D86KJ24N)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
TOCOPHEROL (UNII: R0ZB2556P8)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

C12-15 ALKYL LACTATE (UNII: GC844VRD7E)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84614-002-01	2 in 1 PACKAGE	08/23/2024	
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	08/23/2024	

Labeler - Shenzhen Ctrip Technology Co., Ltd. (403113060)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Ctrip Technology Co., Ltd.		403113060	manufacture(84614-002)

Revised: 1/2026 Shenzhen Ctrip Technology Co., Ltd.