

## **QQE FUNGAL NAIL TREATMENT- tolnaftate 1% liquid**

### **Guangzhou Huixue Biotechnology Co., Ltd.**

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Tolnaftate 1%

Aloe Vera Extract, Helianthus Annuus (Sunflower) Seed Oil, Prunus Armeniaca (Apricot) Kernel Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Eugenia Caryophyllus (Clove) Bud Oil, Lavandula Angustifolia (Lavender) Oil, Hakka Yu, Tocopherol, Origanum Vulgare Oil, Simmondsia Chinensis (Jojoba) Seed Oil, Eucalyptol, Tocopheryl Acetate, Squalene, Propolis Extract, Citrus Bergamia (Bergamot) Fruit Oil.

#### Antifungal

1. Clean the affected area and dry thoroughly. Apply a thin layer of the product over affected area twice daily (morning and night) or as directed by a doctor. Gently file the nail to make it thinner and remove debris, being cautious not to harm the nail bed. For best results, file nails every 2-3 days. 2. Supervise children in the use of this product. For athlete's foot: Pay special attention to spaces between the toes; wear well-fitting, ventilated shoes, and change shoes and socks at least once daily. To prevent athlete's foot, apply once or twice daily (morning and/or night). For toe fungus, apply to skin around the nail and cuticle. Visible improvement may be seen in as little as 7 days. Continue regular use to help prevent fungus regrowth.

For external use only. Do not use on children under 2 years of age unless directed by a doctor. When using this product, avoid contact with the eyes. Stop use and ask a doctor if irritation occurs or if there is no improvement within 4 weeks. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Chest pain, rapid heart beat, faintness, or dizziness occurs sudden, unexplained weight gain occurs, your hands or feet swell.

Twice each day, 1ml each time. 15 days supply



## QQE FUNGAL NAIL TREATMENT

tolnaftate 1% liquid

### Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:87204-161

Route of Administration	TOPICAL
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**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>TOLNAFTATE</b> (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	0.15 mg in 15 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>EUCALYPTOL</b> (UNII: RV6J6604TK)	
<b>SQUALENE</b> (UNII: 7QWM220FJH)	
<b>TOCOPHEROL</b> (UNII: R0ZB2556P8)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
<b>1</b>	NDC:87204-161-01	15 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/30/2026	02/05/2027

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M029	01/30/2026	02/05/2027

**Labeler** - Guangzhou Huixue Biotechnology Co., Ltd. (418186127)

Revised: 1/2026

Guangzhou Huixue Biotechnology Co., Ltd.