

ATHLETE FOOT CREAM- terbinafine hydrochloride cream
Beautivity LLC

Update-85398-006-01

Terbinafine Hydrochloride 1%

Antifungal

- cures most athlete's foot (tinea pedis)
- cures most jock itch (tinea cruris) and ringworm (tinea corporis)
- relieves itching, burning, cracking and scaling which accompany these conditions

For external use only

- on nails or scalp
- in or near the mouth or eyes
- for vaginal yeast infections

do not get into eyes, if eye contact occurs, rinse thoroughly with water.

stop use and ask a doctor if too much irritation occurs or gets worse.

if swallowed, call a poison control centre or get medical help right away.

- adults and children 12 years and over
- use the tip of the cap to break the seal and open the tube
- wash the affected skin with soap and water and dry completely before applying
- for athlete's foot wear well-fitting, ventilated shoes. Change shoes and socks at least once daily.
- between the toes only: apply twice a day (morning and night) for 1 week or as directed by a doctor
- on the bottom or sides of the foot: apply twice a day (morning and night) for 2 weeks or as directed by a doctor
- for jock itch and ringworm: apply once a day (morning or night) for 1 week or as directed by a doctor
- wash hands after each use
- children under 12 years: ask a doctor
- Do not use if seal on tube is broken or is not visible.
- Store at 68–77°F (20–25°C)

Allantoin, Glycerin, Propylene Glycol, PCA Dimethicone, Ethylparaben, Glyceryl Stearate, Primary Alcohol Ethoxylate, Alumen, Sodium Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Isohexadecane, Polysorbate 80, Aloe Barbadensis Leaf Juice, Parfum

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ATHLETE FOOT CREAM

terbinafine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85398-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TERBINAFINE HYDROCHLORIDE (UNII: 012C11ZU6G) (TERBINAFINE - UNII:G7RIW8S0XP)	TERBINAFINE HYDROCHLORIDE	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
POTASSIUM ALUM (UNII: 1L24V9R23S)	
SODIUM ACRYLATE/SODIUM ACRYLOYLDIMETHYLTAURATE COPOLYMER (4000000 MW) (UNII: 1DXE3F3OZX)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
C12-15 PARETH-7 (UNII: 3XY03A79QH)	
ALLANTOIN (UNII: 344S277G0Z)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ETHYLPARABEN (UNII: 14255EXE39)	
ISOHEXADECANE (UNII: 918X1OUF1E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85398-006-01	60 g in 1 CARTON; Type 0: Not a Combination Product	08/18/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	08/18/2025	

Labeler - Beautivity LLC (096573788)

Revised: 3/2026

Beautivity LLC