TOLNAFTATE ANTIFUNGAL- tolnaftate powder Geri-Care Pharmaceuticals, Corp

GC TOLNAFTATE

Active ingredient

Tolnaftate 1%

PURPOSE

ANTIFUNGAL

USES

• for effective treatment of most athlete's foot (tinea pedis)

and ringworm (tinea corporis)

- for effective relief of itchy, scaly skin between the toes
- clears up most athlete's foot infection and with daily use helps keep it from coming back

WARNINGS

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 consult a doctor if

- irritation occurs
- 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.

DIRECTIONS

- shake well before use
- 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
- supervise children in the use of this product
- use daily for 4 weeks. If condition persists longer, consult a doctor.

For athlete's foot

- pay special attention to the spaces between the toes
- wear well-fitting ventilated shoes
- change shoes and socks at least once daily
- to prevent athlete's foot, clean and dry feet thoroughly.

Apply as above once or twice daily.

This product is not effective on the scalp or nails.

Other Information

store at room temperature

INACTIVE INGREDIENTS

corn starch and talc

PACKAGE LABEL



TOLNAFTATE ANTIFUNGAL

tolnaftate powder

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:57896-199

Route of Administration

TOPICAL

Active Ingredient/Active Moiety

-		
Ingredient Name	Basis of Strength	Strength
DESOTIVITY TO INVETATE LINIII-DEVENDESOTIVITY	TOLNATTATE	1 a in 100 a

TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)

OLNAFTATE 1 g in 100 g

Inactive Ingredients

5	
Ingredient Name	Strength
STARCH, CORN (UNII: 08232NY3SJ)	

TALC (UNII: 7SEV7J4R1U)

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:57896-199-45	45 g in 1 BOTTLE, DISPENSING; Type 7: Separate Products Requiring Cross Labeling	08/01/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		08/01/2011	5

Labeler - Geri-Care Pharmaceuticals, Corp (611196254)

Registrant - GCP Laboratories (965480861)

Establishment				
Name	Address	ID/FEI	Business Operations	
GCP Laboratories		965480861	manufacture(57896-199)	

Revised: 11/2023 Geri-Care Pharmaceuticals, Corp