

RITE AID ANTIFUNGAL RINGWORM- clotrimazole cream
Rite Aid Corporation

Rite Aid®
antifungal ringworm

Drug Facts

Active ingredient

Clotrimazole 1%

Purpose

Antifungal

Uses

- cures most ringworm
- relieves itching, redness, irritation and discomfort which accompany this condition

Warnings

For external use only

When using this product avoid contact with eyes

Stop use and ask a doctor if

- irritation occurs
- there is no improvement within 4 weeks

Do not use on children under 2 years of age unless directed by a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- wash affected area and dry thoroughly
- apply a thin layer over affected area twice daily (morning and night)
- supervise children in the use of this product
- use daily for 4 weeks
- if condition persists longer, ask a doctor
- this product is not effective on the scalp or nails

Other information

- To open: unscrew cap, use the pointed end of cap to puncture seal.
- store between 20°- 25°C (68° - 77°F)

- see carton or tube crimp for lot number and expiration date

Inactive ingredients

benzyl alcohol (1%), cetostearyl alcohol, cetyl esters wax, 2-octyldodecanol, polysorbate 60, purified water, sorbitan monostearate

Questions?

Call **1-866-923-4914**

DISTRIBUTED BY:

RITE AID, 30 HUNTER LANE,
CAMP HILL, PA 17011

PRINCIPAL DISPLAY PANEL - 14.2 g Tube Carton





**RITE
AID®
PHARMACY**

Ringworm Cream

Clotrimazole Cream USP, 1%

ANTIFUNGAL

NET WT 0.5 OZ (14.2 g)

RITE AID ANTIFUNGAL RINGWORM			
clotrimazole cream			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-1110
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
OCTYLDODECANOL (UNII: 461N1O614Y)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
WATER (UNII: 059QF0KO0R)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ESTERS WAX (UNII: D072FFP9GU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-1110-1	1 in 1 CARTON	06/01/1995	
1		14.2 g in 1 TUBE; 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	06/01/1995	

Labeler - Rite Aid Corporation (014578892)**Establishment**

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(11822-1110)

Revised: 7/2025

Rite Aid Corporation