

JOCK ITCH- clotrimazole cream
TARGET Corporation

Jock Itch Cream

Drug Facts

Active ingredient

Clotrimazole 1%

Purpose

Antifungal

Uses

- cures most jock itch
- relieves itching, burning, scaling, chafing and discomfort associated with jock itch

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 eyes

Stop use and ask a doctor if

- irritation occurs
- there is no improvement within 2 weeks

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 2 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
Target Corporation
Minneapolis,
MN 55403

PRINCIPAL DISPLAY PANEL - 14.2 g Tube Carton

up & up™

Compare to active ingredient in Lotrimin® AF*

jock itch cream

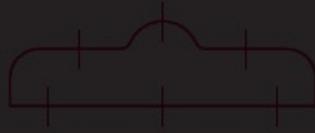
clotrimazole cream USP, 1% antifungal cream

cures most jock itch

relieves itching, burning, scaling and chafing

NET WT ½ OZ (14.2 g)

NDC 11673-062-15



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Drug Facts (continued)

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Minneapolis,
MN 55403
Made in Canada
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Target Brands, Inc.

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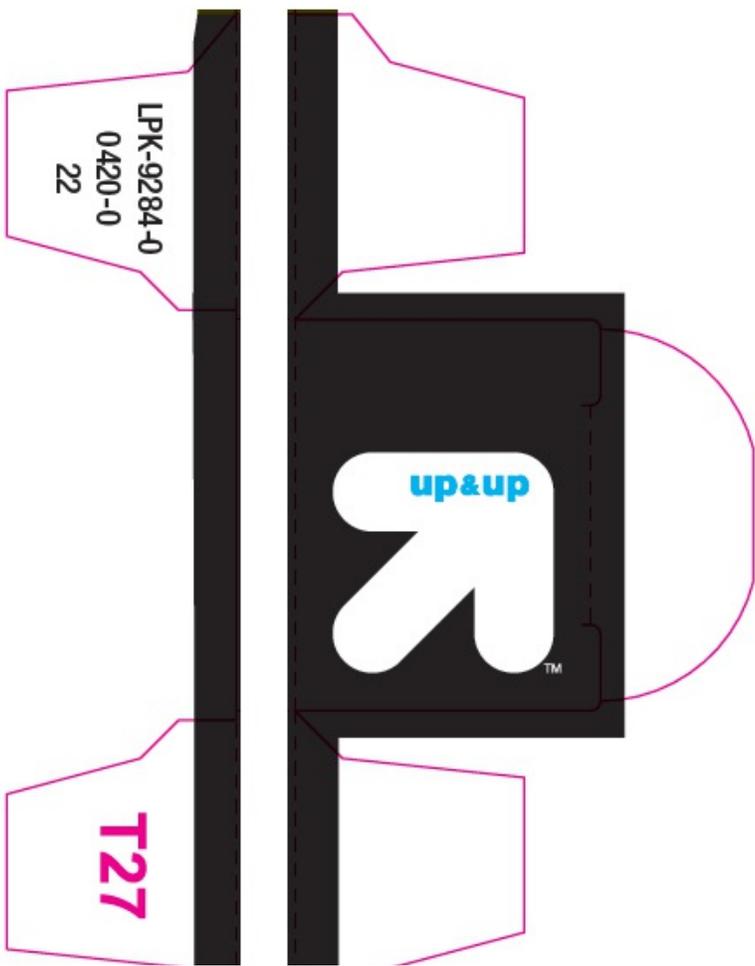
Directions

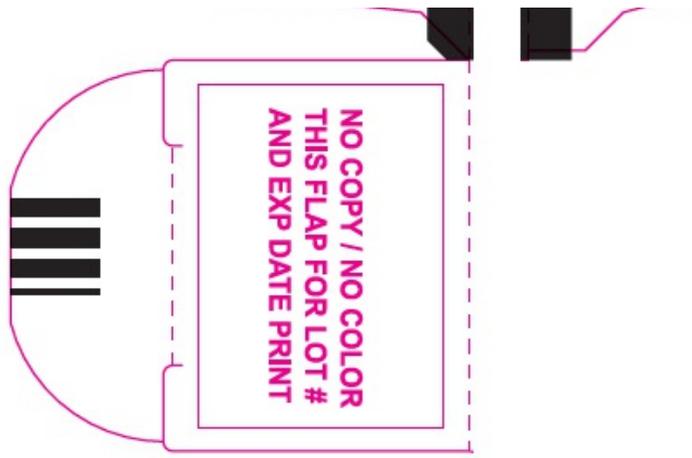
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245 07 0403 R00
C-001227-01-077
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Target Corporation





JOCK ITCH

clotrimazole cream

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:11673-062 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|--------------|
| CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65) | CLOTRIMAZOLE | 10 mg in 1 g |

Inactive Ingredients

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

Product Characteristics

| | | | |
|-----------------|-------|---------------------|--|
| Color | white | Score | |
| Shape | | Size | |
| Flavor | | Imprint Code | |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|-----------|---------------------|----------------------|--------------------|
| | | | | |

| | | | | |
|------------------------------|---|---|-----------------------------|---------------------------|
| 1 | NDC:11673-062-15 | 1 in 1 CARTON | 04/27/2020 | |
| 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 | | 04/27/2019 | |

Labeler - TARGET Corporation (006961700)

| | | | |
|------------------------|----------------|---------------|----------------------------|
| Establishment | | | |
| Name | Address | ID/FEI | Business Operations |
| Sun Pharma Canada Inc. | | 243339023 | manufacture(11673-062) |

Revised: 7/2025

TARGET Corporation