KETOCONAZOLE-shampoo
Patriot Pharmaceuticals, LLC
--------

KETOCONAZOLE 2% SHAMPOO

DESCRIPTION

Ketoconazole 2% shampoo is a red-orange liquid for topical application, containing the broad spectrum
synthetic antifungal agent ketoconazole in a concentration of 2% in an aqueous suspension. It also
contains: coconut fatty acid diethanolamide, disodium monolauryl ether sulfosuccinate, F.D.&C. Red
No. 40, hydrochloric acid, imidurea, laurdimonium hydrolyzed animal collagen, macrogol 120 methyl
glucose dioleate, perfume bouquet, sodium chloride, sodium hydroxide, sodium lauryl ether sulfate,
and purified water.

Ketoconazole is \( \text{cis-1-acetyl-4-[4-[[2-(2,4-dichlorophenyl)-2-(1H-imidazol-1-ylmethyl)-1,3-dioxolan-4-yl]methoxy]phenyl]piperazine} \) and has the following structural formula:

![Ketoconazole Structural Formula]

CLINICAL PHARMACOLOGY

Tinea (pityriasis) versicolor is a non-contagious infection of the skin caused by \( \text{Pityrosporum orbiculare} (\text{Malassezia furfur}) \). This commensal organism is part of the normal skin flora. In susceptible individuals the condition is often recurrent and may give rise to hyperpigmented or hypopigmented patches on the trunk which may extend to the neck, arms and upper thighs. Treatment of the infection may not immediately result in restoration of pigment to the affected sites. Normalization of pigment following successful therapy is variable and may take months, depending on individual skin type and incidental skin exposure. The rate of recurrence of infection is variable.

Ketoconazole 2% was not detected in plasma in 39 patients who shampooed 4–10 times per week for 6
months, or in 33 patients who shampooed 2–3 times per week for 3–26 months (mean: 16 months).

An exaggerated use washing test on the sensitive antecubital skin of 10 subjects twice daily for five
consecutive days showed that the irritancy potential of ketoconazole 2% shampoo was significantly less
than that of 2.5% selenium sulfide shampoo.

A human sensitization test, a phototoxicity study, and a photoallergy study conducted in 38 male and 22
female volunteers showed no contact sensitization of the delayed hypersensitivity type, no phototoxicity
and no photoallergenic potential due to ketoconazole 2% shampoo.

Mode of Action: Interpretations of \( \text{in vivo} \) studies suggest that ketoconazole impairs the synthesis of
ergosterol, which is a vital component of fungal cell membranes. It is postulated, but not proven, that the
therapeutic effect of ketoconazole in tinea (pityriasis) versicolor is due to the reduction of
\( \text{Pityrosporum orbiculare} (\text{Malassezia furfur}) \) and that the therapeutic effect in dandruff is due to
the reduction of \( \text{Pityrosporum ovale} \). Support for the therapeutic effect in tinea versicolor comes from a
three-arm, parallel, double-blind, placebo-controlled study in patients who had moderately severe tinea
(pityriasis) versicolor. Successful response rates in the primary efficacy population for each of both
three-day and single-day regimens of ketoconazole 2% shampoo were statistically significantly greater
(73% and 69%, respectively) than a placebo regimen (5%). There had been mycological confirmation of fungal disease in all cases at baseline. Mycological clearing rates were 84% and 78%, respectively, for the three-day and one-day regimens of the 2% shampoo and 11% in the placebo regimen. While the differences in the rates of successful response between either of the two active treatments and placebo were statistically significant, the difference between the two active regimens was not.

**Microbiology:** Ketoconazole 2% is a broad spectrum synthetic antifungal agent which inhibits the growth of the following common dermatophytes and yeasts by altering the permeability of the cell membrane: dermatophytes: *Trichophyton rubrum*, *T. mentagrophytes*, *T. tonsurans*, *Microsporum canis*, *M. audouini*, *M. gypseum* and *Epidermophyton floccosum*; yeasts: *Candida albicans*, *C. tropicalis*, *Pityrosporum ovale* (*Malassezia ovale*) and *Pityrosporum orbiculare* (*M. furfur*). Development of resistance by these microorganisms to ketoconazole has not been reported.

**INDICATIONS AND USAGE**

Ketoconazole 2% shampoo is indicated for the treatment of tinea (pityriasis) versicolor caused by or presumed to be caused by *Pityrosporum orbiculare* (also known as *Malassezia furfur* or *M. orbiculare*).

Note: Tinea (pityriasis) versicolor may give rise to hyperpigmented or hypopigmented patches on the trunk which may extend to the neck, arms and upper thighs. Treatment of the infection may not immediately result in normalization of pigment to the affected sites. Normalization of pigment following successful therapy is variable and may take months, depending on individual skin type and incidental sun exposure. Although tinea versicolor is not contagious, it may recur because the organism that causes the disease is part of the normal skin flora.

**CONTRAINDICATIONS**

Ketoconazole 2% shampoo is contraindicated in persons who have known hypersensitivity to the active ingredient or excipients of this formulation.

**PRECAUTIONS**

Severe hypersensitivity reactions, including anaphylaxis, have been reported during post-marketing use of ketoconazole shampoo. If a reaction suggesting sensitivity or chemical irritation should occur, use of the medication should be discontinued.

**Information for Patients:**

Patients should be advised of the following:
- Ketoconazole 2% shampoo may be irritating to mucous membranes of the eyes and contact with this area should be avoided.
- The following have been reported with the use of ketoconazole 2% shampoo: hair discoloration and abnormal hair texture, removal of the curl from permanently waved hair, itching, skin burning sensation and contact dermatitis, hypersensitivity, angioedema, alopecia, rash, urticaria, skin irritation, dry skin, and application site reactions.
- Patients who develop allergic reactions, such as generalized rash, skin reactions, severe swelling, angioedema, or shortness of breath should discontinue ketoconazole and contact their physician immediately.

**Carcinogenesis, Mutagenesis, Impairment of Fertility:**

Long-term studies to assess the carcinogenic potential of ketoconazole 2% shampoo have not been conducted. A long-term feeding study of ketoconazole in Swiss Albino mice and in Wistar rats showed no evidence of oncogenic activity. The dominant lethal mutation test in male and female mice revealed that single oral doses of ketoconazole as high as 80 mg/kg were not genotoxic. The Ames Salmonella microsomal activator assay was also negative.
**Pregnancy: Teratogenic effects:**

There are no adequate and well-controlled studies in pregnant women. Ketoconazole should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. In humans, ketoconazole is not detected in plasma after chronic shampooing on the scalp.

Ketoconazole has been shown to be teratogenic (syndactylyia and oligodactylia) in the rat when given orally in the diet at 80 mg/kg/day (a dose 10 times the maximum recommended human oral dose). However, these effects may be related to maternal toxicity, which was seen at this and higher dose levels.

**Nursing mothers:** There are no adequate and well-controlled studies in nursing women. Ketoconazole is not detected in plasma after chronic shampooing on the scalp. Caution should be exercised when ketoconazole 2% shampoo is administered to a nursing woman.

**Pediatric Use:** Safety and effectiveness in children have not been established.

**ADVERSE REACTIONS**

**Clinical Trials Experience**

In 11 double-blind trials in 264 patients using ketoconazole 2% shampoo for the treatment of dandruff or seborrheic dermatitis, an increase in normal hair loss and irritation occurred in less than 1% of patients. In three open-label safety trials in which 41 patients shampooed 4–10 times weekly for six months, the following adverse experiences each occurred once: abnormal hair texture, scalp pustules, mild dryness of the skin, and itching. As with other shampoos, oiliness and dryness of hair and scalp have been reported. In a double-blind, placebo-controlled trial in which patients with tinea versicolor were treated with either a single application of ketoconazole 2% shampoo (n=106), a daily application for three consecutive days (n=107), or placebo (n=105), drug-related adverse events occurred in 5 (5%), 7 (7%) and 4 (4%) of patients, respectively. The only events that occurred in more than one patient in any of the three treatment groups were pruritus, application site reaction, and dry skin; none of these events occurred in more than 3% of the patients in any one of the three groups.

**Post-marketing Experience**

Because these reactions are reported voluntarily from a population of uncertain size, it is not possible to reliably estimate their frequency. The following adverse drug reactions have been identified during post-marketing experience with ketoconazole 2% shampoo: there have been reports of hair discoloration and abnormal hair texture, itching, skin burning sensation, contact dermatitis, hypersensitivity, angioedema, alopecia, rash, urticaria, skin irritation, dry skin, and application site reactions.

**OVERDOSAGE**

Ketoconazole 2% shampoo is intended for external use only. In the event of accidental ingestion, supportive and symptomatic measures should be employed. Induced emesis and gastric lavage should not be performed to avoid aspiration.

**DOSAGE AND ADMINISTRATION**

Apply the shampoo to the damp skin of the affected area and a wide margin surrounding this area. Lather, leave in place for 5 minutes, and then rinse off with water.

One application of the shampoo should be sufficient.
HOW SUPPLIED

Ketoconazole 2% shampoo is a red-orange liquid supplied in a 4-fluid ounce (120 mL) nonbreakable plastic bottle (NDC 10147-0750-4).

Ketoconazole 2% shampoo is a red-orange liquid supplied in a 4-fluid ounce (120 mL) nonbreakable plastic bottle in a carton of one (NDC 10147-0750-5).

Storage conditions: Store at a temperature not above 25°C (77°F). Protect from light.

Keep out of reach of children

For more information call 1-800-667-8570

Product of Belgium

Manufactured by:
Janssen Pharmaceutica NV
Beerse, Belgium

Manufactured for:
Patriot Pharmaceuticals, LLC
Horsham, PA 19044

©2005 Patriot

Edition: December 2017

PRINCIPAL DISPLAY PANEL - 120 mL Bottle Box

NDC 10147-0750-5

KETOCONAZOLE 2%
Shampoo

for topical application only.
Rx only.
4 fl. oz. (120 mL)

PATRIOT
PHARMACEUTICALS LLC
### Active Ingredient/Active Moiety

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>ketoconazole</td>
<td>ketoconazole</td>
<td>20 mg in 1 mL</td>
</tr>
</tbody>
</table>

### Inactive Ingredients

<table>
<thead>
<tr>
<th>Ingredient Name</th>
<th>Strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>coco diethanolamide</td>
<td></td>
</tr>
<tr>
<td>disodium laureth sulfosuccinate</td>
<td></td>
</tr>
<tr>
<td>FD&amp;C RED NO. 40</td>
<td></td>
</tr>
<tr>
<td>HYDROCHLORIC ACID</td>
<td></td>
</tr>
<tr>
<td>IMIDUREA</td>
<td></td>
</tr>
<tr>
<td>PEG-120 METHYL GLUCOSE DIOLEATE</td>
<td></td>
</tr>
<tr>
<td>SODIUM LAURETH-3 SULFATE</td>
<td></td>
</tr>
<tr>
<td>WATER</td>
<td></td>
</tr>
<tr>
<td>SODIUM CHLORIDE</td>
<td></td>
</tr>
<tr>
<td>SODIUM HYDROXIDE</td>
<td></td>
</tr>
</tbody>
</table>

### Product Characteristics

<table>
<thead>
<tr>
<th>Color</th>
<th>RED (red-orange)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Shape</td>
<td></td>
</tr>
<tr>
<td>Size</td>
<td></td>
</tr>
<tr>
<td>Flavor</td>
<td></td>
</tr>
<tr>
<td>Imprint Code</td>
<td></td>
</tr>
<tr>
<td>Contains</td>
<td></td>
</tr>
</tbody>
</table>

### Packaging

<table>
<thead>
<tr>
<th>#</th>
<th>Item Code</th>
<th>Package Description</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>NDC:10147-0750-4</td>
<td>120 mL in 1 BOTTLE; Type 0: Not a Combination Product</td>
<td>07/01/2005</td>
<td>08/31/2019</td>
</tr>
<tr>
<td>2</td>
<td>NDC:10147-0750-5</td>
<td>1 in 1 BOX</td>
<td>04/01/2017</td>
<td>09/30/2020</td>
</tr>
<tr>
<td>2</td>
<td></td>
<td>120 mL in 1 BOTTLE; Type 0: Not a Combination Product</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Marketing Information

<table>
<thead>
<tr>
<th>Marketing Category</th>
<th>Application Number or Monograph Citation</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>NDA authorized generic</td>
<td>NDA019927</td>
<td>07/01/2005</td>
<td>09/30/2020</td>
</tr>
</tbody>
</table>

### Labeler

- Patriot Pharmaceuticals, LLC (178220385)

### Establishment

<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Janssen Pharmaceutica NV</td>
<td></td>
<td>370005019</td>
<td>ANALYSIS(10147-0750), MANUFACTURE(10147-0750)</td>
</tr>
</tbody>
</table>
### Establishment

<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Noramco, Inc.</td>
<td>057234486</td>
<td>API MANUFACTURE(10147-0750)</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
<th>Business Operations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Janssen Pharmaceutical Sciences Unlimited Company</td>
<td>985639841</td>
<td>API MANUFACTURE(10147-0750)</td>
<td></td>
</tr>
</tbody>
</table>

Revised: 10/2019

Patriot Pharmaceuticals, LLC