



PRODUCT MONOGRAPH

Pr STIEPROX[®]

ciclopirox olamine (shampoo) 1.5% w/w

Topical Antifungal

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Pr **STIEPROX®**

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PART I: HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Route of Administration	Dosage Form / Strength	Clinically Relevant Nonmedicinal Ingredients
Topical	Shampoo 1.5% w/w	Coconut diethanolamide, Dipropylene glycol and Sodium lauryl ether sulphate <i>For a complete listing see DOSAGE FORMS, COMPOSITION AND PACKAGING section.</i>

INDICATIONS AND CLINICAL USE

STIEPROX® (ciclopirox olamine 1.5% w/w) shampoo is indicated for:

- Topical treatment and prophylaxis of dandruff or the treatment of seborrhoeic dermatitis in which the yeast *Malassezia furfur* is involved.

Geriatrics (≥ 65 years of age):

The safety and efficacy of STIEPROX® have not been established in geriatric patients.

Pediatrics (< 18 years of age):

The safety and efficacy of STIEPROX® have not been established in pediatric patients.

CONTRAINDICATIONS

- Patients who are hypersensitive to this drug or to any ingredient in the formulation or component of the container. For a complete listing, see the DOSAGE FORMS, COMPOSITION AND PACKAGING section.

WARNINGS AND PRECAUTIONS

General

STIEPROX[®] (ciclopirox olamine 1.5% w/w) shampoo is for external use only. Avoid contact with eyes. In case of contact with the eyes, rinse thoroughly with copious amounts of water. If irritation persists, a physician should be consulted.

Skin

If a reaction suggesting hypersensitivity or chemical irritation should occur with the use of STIEPROX[®] shampoo, treatment should be applied less often or should be discontinued and appropriate therapy instituted.

Hair Discolouration: A discolouration of the hair has been observed, mainly in patients with grey, white or chemically treated hair (for example, due to hair dye) (see ADVERSE REACTIONS).

Special Populations

Pregnant Women: The safety of STIEPROX[®] shampoo during pregnancy has not been established. There is evidence that ciclopirox olamine crosses the placental barrier in animals. Reproductive studies in mice, rats, rabbits and monkeys, at doses of ciclopirox olamine 10 times that of a topical human dose, have revealed no significant evidence of impaired fertility or harm to the fetus. STIEPROX[®] should only be used during pregnancy if the potential benefits to the mother justify the potential risks to the fetus.

No pregnant women were enrolled during clinical trials with STIEPROX[®].

Nursing Women: It is unknown if ciclopirox olamine is excreted in human milk. Because many drugs are excreted in human milk precaution should be exercised. Patients are advised to ensure that any residual product is fully washed off the breast prior to breast-feeding to avoid possible ingestion by nursing infant.

Pediatrics (< 18 years of age): The safety and effectiveness of STIEPROX[®] has not been established in pediatric patients. No data is available in children less than 12 years of age.

Geriatrics (≥ 65 years of age): The safety and effectiveness of STIEPROX[®] has not been established in a geriatric population.

ADVERSE REACTIONS

Adverse Drug Reaction Overview

STIEPROX[®] (ciclopirox olamine 1.5% w/w) shampoo is well tolerated with a low incidence of adverse reactions.

Clinical Trial Adverse Drug Reactions

Because clinical trials are conducted under very specific conditions the adverse reaction rates observed in the clinical trials may not reflect the rates observed in practice and should not be compared to the rates in the clinical trials of another drug. Adverse drug reaction information from clinical trials is useful for identifying drug-related adverse events and for approximating rates.

Table 1 provides the list of the Common Adverse Drug Reactions (>1%) observed with STIEPROX[®] (ciclopirox olamine 1.5% w/w) shampoo during the controlled clinical trials.

Table 1 Common Adverse Drug Reaction to STIEPROX[®] or its vehicle

	STIEPROX[®] Shampoo (1.5% ciclopirox olamine) (n= 258)	Active Comparator (n= 84)	Placebo (n= 164)
Skin and subcutaneous tissue disorders			
Pruritus*	3 (1.2%)	--	3 (1.8%)
Skin Irritation*	5 (1.9%)	--	3 (1.8%)
Scalp Irritation	0	2 (2.4%)	0

Less Common Clinical Trial Adverse Drug Reactions (<1%)

Administration site conditions: erythema^{*}, erosion of scalp, application site rash^{*}

Eye disorders: stinging, burning, watering, infection

Gastrointestinal disorders: nausea

Nervous system disorders: headache

* exacerbation of symptoms

Abnormal Hematologic and Clinical Chemistry Findings

Hematologic and clinical chemistry parameters were not evaluated during clinical trials with STIEPROX[®].

Post-Market Adverse Drug Reactions

In addition to adverse drug reactions identified from clinical trials, the following adverse reactions have been identified during post-approval use of STIEPROX[®]. Because these events have been reported voluntarily from a population of unknown size, estimates of frequency cannot be made.

Immune system disorder: application site hypersensitivity

Skin and subcutaneous tissue disorders: burning sensation^{*}, skin exfoliation^{*}, eczema, alopecia^{*}, hair colour changes and hair texture abnormal^{*}.

* exacerbation of symptoms

DRUG INTERACTIONS

Overview

No drug interactions have been reported with ciclopirox olamine. The possibility of interaction with alcohol has not been evaluated.

Drug-Drug Interactions

Interactions with other drugs have not been established.

Drug-Food Interactions

Interactions with food have not been established.

Drug-Herb Interactions

Interactions with herbal products have not been established.

Drug-Laboratory Interactions

Interactions with laboratory tests have not been established.

DOSAGE AND ADMINISTRATION

Dosing Considerations

Situations that may affect the dosing of STIEPROX[®] have not been established.

Recommended Dose and Dosage Adjustment

Dandruff: Use 2 or 3 times a week or as often as necessary.

Seborrhoeic dermatitis: Use 3 times a week or as often as necessary.

Missed Dose

Any missed application of the shampoo should be done the next day.

Administration

Note that this product is only for topical application to the scalp and adjacent areas.

The hair should be wet and sufficient STIEPROX[®] (ciclopirox olamine 1.5% w/w) shampoo should be applied to produce an abundant lather. The affected areas (scalp and/or its edges) should be vigorously massaged with the fingertips for two to three minutes. The hair should then be thoroughly rinsed and the procedure repeated.

OVERDOSAGE

For management of a suspected drug overdose, contact your regional Poison Control Centre.

There have been no clinical reports of acute overdosage with STIEPROX[®] (ciclopirox olamine 1.5% w/w) shampoo.

Symptoms

Oral ingestion is usually followed by nausea and vomiting due to the detergent.

Treatment

Management should be as clinically indicated. If ingested, treatment should be symptomatic. In the event of accidental ingestion, only supportive measures should be carried out. In order to avoid aspiration, neither emesis nor gastric lavage should be performed.

ACTION AND CLINICAL PHARMACOLOGY

Mechanism of Action

Ciclopirox olamine is a synthetic antifungal which is structurally unrelated to the common imidazoles or other antifungals. Unlike most antifungals, ciclopirox olamine does not affect sterol biosynthesis. It has been suggested that ciclopirox olamine interferes with the active uptake and accumulation of some essential substrates and/or polyvalent cations which results in cell death because of cellular depletion. Ciclopirox olamine may act through chelation of polyvalent cations such as Fe^{3+} or Al^{2+} .

Pharmacodynamics

Except for its specific fungicidal activity, ciclopirox olamine when formulated in a 1.5% w/w shampoo is not expected to exert any other pharmacodynamic effect when applied topically. No pharmacodynamic studies were conducted in humans with STIEPROX[®] (ciclopirox olamine 1.5% w/w) shampoo.

Pharmacokinetics

No pharmacokinetic studies were conducted in humans with STIEPROX[®] (ciclopirox olamine 1.5% w/w) shampoo.

Special Populations and Conditions

STIEPROX[®] was not tested in special populations.

STORAGE AND STABILITY

Store STIEPROX[®] between 4° and 30°C.

SPECIAL HANDLING INSTRUCTIONS

None.

DOSAGE FORMS, COMPOSITION AND PACKAGING

STIEPROX[®] contains ciclopirox olamine USP / EP 1.5% w/w in a mild aqueous shampoo base.

STIEPROX[®] contains the following excipients: purified water, sodium lauryl ether sulphate, cocamidopropyl betaine, disodium phosphate dodecahydrate, citric acid

monohydrate, coconut diethanolamide, hexylene glycol, oleyl alcohol, polysorbate 80, polyquaternium 10, fragrance (including dipropylene glycol), and sodium hydroxide.

STIEPROX[®] shampoo is a clear, yellow to light orange coloured, viscous liquid.

STIEPROX[®] shampoo is packaged into 150 mL HDPE bottles, with polypropylene screw caps and 10 mL, laminated, aluminum foil sample size sachets.

PART II: SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

Drug Substance

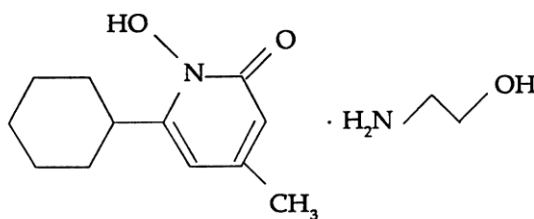
Proper name: Ciclopirox olamine

Chemical name: 2(1H)-Pyridinone, 6-cyclohexyl-1-hydroxy-4-methyl-,
compound with 2-aminoethanol (1:1)

Molecular formula and molecular mass: $C_{12}H_{17}NO_2 \bullet C_2H_7NO$

Structural formula: 268.35

Structural formula:



Physicochemical properties: White-to-cream white crystalline odourless powder. Very soluble (1 part in less than 1 part of solvent) in ethanol, methylene chloride and chloroform; slightly soluble (1 part in 100 to 1,000 parts of solvent) in ethyl acetate and water; practically insoluble (1 part in more than 10,000 parts of solvent) in cyclohexane and ether. Ciclopirox olamine has a melting point of superior to 100°C with decomposition and a pH of 8.0 to 9.0 (in a 1% water solution).

CLINICAL TRIALS

A) DANDRUFF

Study demographics and trial design

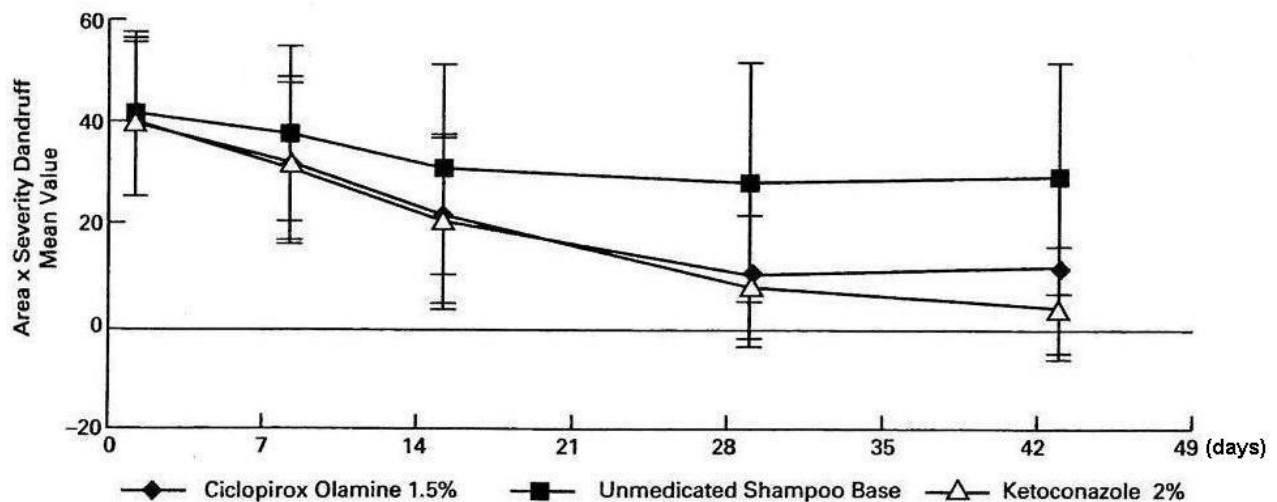
Table 2 Summary of patient demographics for clinical trials in specific indication

Study #	Trial design	Dosage, route of administration and duration	Study subjects (n=number)	Mean age (Range)	Gender
S177-GB-030[Ref 1]	Double blind, parallel groups, randomized single-centre	two times per week topical 29 days of treatment	n=163	45.2 years (18-70 years)	female: 35% male: 65%

All patients were healthy Caucasian men or women with severe dandruff and/or seborrhoeic dermatitis.

Study results

Figure 1 Whole-head study, showing the mean area x severity dandruff scores over time



Note: The mean reductions from baseline were all significantly greater than for the control for both the ciclopirox olamine ($P < 0.01$ at days 8 and 15, $P < 0.001$ at days 29 and 43) and ketoconazole ($P < 0.05$ at day 8, $P < 0.01$ at day 15, $P < 0.001$ at days 29 and 43) treatment groups.

B) SEBORRHOEIC DERMATITIS

Study demographics and trial design

Table 3 Summary of patient demographics for clinical trials in specific indication

Study #	Trial design	Dosage, route of administration and duration	Study subjects (n=number)	Mean age (Range)	Gender
SCI-2002-01	Double blind, parallel groups, randomized multi-centre	three times per week topical 42 days of treatment	n=103	43.8 years (18-84 years)	female: 36% male: 64%

All patients were healthy men or women with moderate to severe seborrhoeic dermatitis.
88% were Caucasians.

Table 4 Severity of Seborrhoeic Dermatitis

Treatment Week	Severity of Seborrhoeic Dermatitis				P value
	STIEPROX®		placebo		
	Enrolled / Completed	mean severity	Enrolled / Completed	mean severity	
Baseline	69/69	3.17	34/34	3.29	0.1636
Week 2	69/69	2.14	34/34	2.65	0.0094
Week 4	69/68	1.40	34/33	2.30	0.0003
Week 6	69/68	1.22	34/33	2.27	0.0004

Table 5 **Severity of Erythema**

Treatment Week	Severity of Erythema				P value
	STIEPROX®		placebo		
	Enrolled / Completed	mean severity	Enrolled / Completed	mean severity	
Baseline	69/69	3.17	34/34	3.29	0.1636
Week 2	69/69	1.80	34/34	2.47	0.0025
Week 4	69/68	1.07	34/33	1.94	0.0009
Week 6	69/68	1.06	34/33	1.79	0.0188

Table 6 **Severity of Scaling**

Treatment Week	Severity of Scaling				P value
	STIEPROX®		placebo		
	Enrolled / Completed	mean severity	Enrolled / Completed	mean severity	
Baseline	69/69	2.32	34/34	3.26	0.5950
Week 2	69/69	1.93	34/34	2.50	0.0154
Week 4	69/68	1.26	34/33	2.03	0.0047
Week 6	69/68	1.19	34/33	1.94	0.0164

Table 7 **Severity of Pruritus**

Treatment Week	Severity of Pruritus				P value
	STIEPROX®		placebo		
	Enrolled / Completed	mean severity	Enrolled / Completed	mean severity	
Baseline	69/69	2.32	34/34	2.44	0.7699
Week 2	69/69	1.45	34/34	2.06	0.0099
Week 4	69/68	0.97	34/33	1.73	0.0010
Week 6	69/68	1.09	34/33	1.61	0.0417

Table 8 **Surface Area of Target Lesion**

Treatment Week	Surface Area of Target Lesion				P value
	STIEPROX®		placebo		
	Enrolled / Completed	mean severity	Enrolled / Completed	mean severity	
Baseline	69/69	37.08	34/34	27.88	0.9860
Week 2	69/69	30.61	34/34	29.16	0.3837
Week 4	69/68	19.51	34/33	25.45	0.0261
Week 6	69/68	15.86	34/33	24.15	0.0335

Comparative Bioavailability Studies

No bioavailability studies were conducted with STIEPROX® shampoo.

DETAILED PHARMACOLOGY

In addition to its antifungal activity, ciclopirox olamine has anti-inflammatory activity as a result of its ability to inhibit the synthesis of prostaglandins and leukotrienes. It has been shown to significantly reduce arachidonic acid-induced ear-oedema ($p < 0.05$), as measured by percentage change from control-inflamed ears in mice. The anti-

inflammatory effect of antifungal preparations, including 1% ciclopirox olamine cream on the skins of healthy human volunteers was examined. Compared with oxiconazole, econazole, hydrocortisone and ketoconazole, ciclopirox olamine was significantly superior in inhibiting UVB-induced erythema. It was equipotent when compared with terbinafine or naftifine.

Although ciclopirox olamine has no characteristic pharmacological actions, it did reduce spontaneous behaviour, caused irregular respiration and lowered the body temperature of rats and mice. It did not affect the respiration, electrocardiogram or blood pressure of anaesthetised rabbits or cats. It had no effect upon isolated organ preparations, nor was it phototoxic or irritant after topical application.

When ciclopirox olamine was injected cutaneously, there was an irritant effect on the skin. Similarly, there was an irritant effect when it was instilled into the conjunctival sacs in rabbits.

Pharmacokinetics

Studies in animals and humans showed that dermal absorption of ciclopirox olamine after the topical application of a cream containing 1% ciclopirox olamine was relatively low. The percutaneous absorption in dogs was 5-15% of the given dose and 11-12% of the given dose in rats. The percutaneous absorption in humans was shown to be about 1.3% of the administered dose.

The *in vitro* penetration of ciclopirox olamine through the epidermis of animal and human skin has been examined. The results showed that in rats 20% to 40% of the applied ciclopirox olamine was absorbed over the first 24 hours, whereas, only 1.6% to 3.0% was absorbed through human skin over the same time period. For humans, this level of absorption was similar to that found by other investigators using cream formulations. However, the shampoo formulation appeared to provide a better vehicle than the cream formulation for penetration through rat skin. Ten percent (10%) dilutions of the concentrates in water showed very slow rates of absorption and the level of absorption was very much less than the level seen with the concentrates.

Penetration of the skin appears to be via the epidermis and the hair follicles.

Ciclopirox olamine has also been shown to penetrate through the surface of the fingernails. There was evidence from pregnant rat studies that the compound could enter the placenta. However, despite good absorption from the gastro-intestinal tract, placental transfer remained low.

Distribution

Following oral administration of ciclopirox olamine to humans, affinity of ciclopirox olamine to serum proteins was found to be $96\pm 2\%$ in the concentration range of 0.01 to 11.0 $\mu\text{g/mL}$.

After oral administration of radio labelled ciclopirox olamine to rats, radioactivity could be found in all organs, tissues and body fluid examined. These included spleen, adrenal glands, kidneys, gonads, liver, heart, lung, skeletal musculature, smooth muscle, retroperitoneal fatty tissue, brain, bone marrow, eyes, intestine, skin, blood and plasma. There was evidence that ciclopirox olamine crossed the placental barrier, but apparently only slightly. However, after dermal application, absorption was marginal and although radioactivity was found in a number of organs and tissues, only in the gall bladder, urinary bladder and large intestine were there any notable concentrations.

The metabolic patterns after oral and dermal application are similar. Glucuronidation of ciclopirox olamine appears to be the major form of its metabolism. In dogs, about 43% of the given dose was excreted via the kidneys within the first day, and of this, about three quarters was in the form of the glucuronide of ciclopirox olamine.

Excretion of ciclopirox olamine in dogs after topical application, was very slow and occurred primarily in the urine. Levels of excreted radioactivity in the urine and the feces were extremely low compared with the administered dose.

MICROBIOLOGY

Ciclopirox olamine (2(1H)-Pyridinone, 6-cyclohexyl-1-hydroxy-4-methyl-, compound with 2-aminoethanol (1:1)) is a broad spectrum antimycotic agent with some antibacterial activity. It is effective against pathogenic dermatophytes and yeasts. Important in its efficacy in the treatment of dandruff/seborrhoeic dermatitis and other mild dermatoses is that ciclopirox olamine is an effective antifungal against *Pityrosporum ovale* and *Pityrosporum orbiculare*. The latter are the yeast forms of *Malassezia furfur*, which have been implicated as the causative organisms in conditions such as dandruff. Ciclopirox olamine has a steep dose response line and a high penetration capacity, enabling it to reach deeper horny layers of the skin. These points allow for better elimination of the pathogens. Ciclopirox olamine exhibits antibacterial activity against a variety of Gram-positive and Gram-negative bacteria. Ciclopirox olamine inhibits the growth of dermatophytes, yeast, dimorphic fungi and actinomycetales at concentrations of 4 mg/L or less when these organisms were grown in Sabouraud dextrose media containing beef peptone free of certain minerals. The *in vitro* anti-fungal activity of ciclopirox olamine has been summarized in Table 9.

Table 9 *In-vitro* Susceptibility of 124 Dermatophyte Strains, 104 Yeasts and Yeast - Like Fungi and 29 Moulds of Various Species to Ciclopirox Olamine.

Test Organism	No. of Strains	Cumulative % of Strains Inhibited at MICs (mg/L)					
		0.49	0.98	1.95	3.9	7.8	<31
<i>Trichophyton rubrum</i>	37	2.7	40.5	94.6	100		
<i>Trichophyton mentagrophytes</i>	29	0	34.5	93.1	100		
Other <i>Trichophyton species</i>	22	0	27.3	95.5	100		
<i>Microsporum canis</i>	20	5	15	80	100		
Other <i>Microsporum species</i>	11	0	9.1	54.6	100		
<i>Epidermophyton floccosum</i>	5	0	40	100			
<i>Candida albicans</i>	37	0	56.8	86.5	100		
<i>Candida tropicalis</i>	12	0	50	100			
Other <i>Candida species</i>	36	0	8.3	63.9	100		
Other yeasts	19	0	21.1	89.5	100		
Moulds	29	0	3.5	34.5	69	79.3	100

In vivo studies in guinea pigs with experimental trichophytosis showed that application of ciclopirox olamine, 0.06% to 2.0% in isopropyl alcohol for 5 days after infection, resulted in a 20 to 85% suppression of the spread of mycosis and a decrease in the diameters of the areas of alopecia. Creams of ciclopirox olamine have also been shown to be effective in this experimental model.

Ciclopirox olamine affects the cytoplasmic membrane where it appears to impair active transport mechanisms, cell respiratory processes and membrane integrity. Ciclopirox also negatively influences the macromolecular synthesis of nucleic acids and proteins. Cell death occurs primarily due to cellular depletion.

TOXICOLOGY

Acute Toxicity

Acute toxicity studies with STIEPROX[®] (ciclopirox olamine 1.5%) shampoo have not been conducted. Ciclopirox olamine has a low order of oral toxicity with LD₅₀ values ranging between 2 and 4 g/kg in rats and mice.

Table 10 LD₅₀ (mg/kg) Values for Ciclopirox Olamine

Route of Administration	Species	
	Mice	Rats
Oral	1740 - >2500	3290
Intravenous	74 – 74	72 – 79
Intraperitoneal	83 – 88	146 – 172
Sub-cutaneous	>2500	>2500

Subacute Toxicity

In one-month and 3-month repeat oral dosing studies using rats, ciclopirox olamine at dose levels of 30 mg/kg and 10 mg/kg, respectively, were tolerated and were symptom-free.

Six-month dermal toxicity studies in dogs showed that the animals tolerated the daily application of 1% ciclopirox olamine. There was no sign of systemic effects at any of the doses applied (0.63-2.5 mg/kg body-weight) and reactions to treatment were only seen at the site of application.

Ciclopirox olamine 1% or 5% solutions in PEG-400, were applied twice weekly to the shaven dorsal skin of female mice (40 mice in each treatment group and 30 in the untreated control group), over a period of one year. There were no indications of any toxic or carcinogenic effects.

Local Toxicity

When ciclopirox olamine was injected intra cutaneously or applied to the conjunctival sac in rabbits, local irritative action was observed. However, when the compound was applied topically to the skin of rabbits, it was well tolerated. Ciclopirox olamine was not phototoxic in guinea pigs.

Ciclopirox olamine 1% solution in PEG-400 applied for 24 hours to the shaven skin of rabbits using gauze tape, did not produce any topical or systemic pathological changes. After application of 1, 2 or 4% solutions for 20 consecutive days, a correlation between

the severity of the skin reaction and the increasing concentration of ciclopirox olamine applied was noted. Histological investigations revealed inflammatory reactions in individual animals treated with the 1% and 2% solutions. Thickening of the germinative layer and hyperkeratosis, in addition to inflammatory changes, were noted in the animals treated with 4% solution. No systemic changes were found in any of the groups.

A one-month dermal toxicity study in rabbits showed that the animals tolerated the daily application of 1% ciclopirox olamine cream (0.5 g/60cm², 1.0 g/120cm², 2.0 g/240cm²) compared with 2.0 g of placebo cream applied to 240 cm² of skin. Redness and scabbing were seen in rabbits with either intact and abraded skin, to the same extent in the treatment and control groups. Four weeks after the discontinuation of the treatment, these changes were no longer evident. Matching results were seen in a similar study in guinea pigs which were given doses of 0.125 g/15cm², 0.25 g/30cm², 0.5 g/60cm² of 1% ciclopirox olamine cream compared with 0.5 g of placebo cream applied to 60 cm² of skin. Macroscopic examination revealed no skin reactions. Histologically, there was thickening of the epidermis in both study and control animals, which was reversible within four weeks.

Groups of 2 male/2 female beagle dogs were treated with 0.5 g of 1% ciclopirox olamine cream or 0.5 g of placebo cream applied to an area of 6 cm x 6 cm of skin for 30 consecutive days. Both formulations were well tolerated without any adverse reaction being recorded.

The results from the local irritation studies are somewhat contradictory. In the study in which ciclopirox olamine was in solution with PEG-400, there was evidence of histological changes and inflammatory reactions in the skin of rabbits after topical application over a period of 20 consecutive days. The severity of these changes was reported to be related to the concentration of ciclopirox olamine used, although the incidence and severity of the skin responses to 1% ciclopirox olamine solution were minimal. This study did not appear to have a placebo treated group. Therefore, it is difficult to relate the response of the skin to ciclopirox olamine to that caused by application of PEG-400 alone.

In two other studies, one involving rabbits and one involving dogs, a cream formulation of ciclopirox olamine applied for 30 consecutive days, did not cause any greater reaction in the skin, than that seen with the placebo control. From these observations, it must be concluded that the 1% cream formulation of ciclopirox olamine has a very low probability of causing any reaction to the skin of humans when used as instructed.

Teratology

No study to determine the teratologic potential of STIEPROX[®] was conducted. Studies in animals given oral or subcutaneous ciclopirox olamine failed to reveal any reproduction toxicity. There is evidence that ciclopirox olamine crosses the placental barrier in animals.

Mutagenicity

There is no evidence that ciclopirox olamine is mutagenic following topical, oral or subcutaneous administration to a number of animal species.

Carcinogenicity

There is no evidence from animal studies to suggest that ciclopirox olamine is carcinogenic when administered orally, topically or subcutaneously.

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PART III: CONSUMER INFORMATION

Pr **STIEPROX[®]**
ciclopirox olamine (shampoo) 1.5% w/w

This leaflet is part III of a three-part "Product Monograph" published when STIEPROX[®] was approved for sale in Canada and is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about STIEPROX[®]. Contact your doctor or pharmacist if you have any questions about the drug.

ABOUT THIS MEDICATION**What the medication is used for:**

STIEPROX[®] is a shampoo used to treat and prevent the return of dandruff or to treat seborrheic dermatitis in adults 18 years of age and older. It is important to read and follow these instructions for use.

It is not known if STIEPROX[®] is safe and effective in children less than 18 years old, or in adults more than 65 years old.

What it does:

Dandruff and seborrheic dermatitis can be caused by a fungus called *Malassezia furfur*. STIEPROX[®] contains the antifungal agent ciclopirox olamine which is a medication that, when applied to the skin of the scalp, will "kill" *M. furfur*.

When it should not be used:

STIEPROX[®] should not be used if you are hypersensitive to ciclopirox olamine or to any of the ingredients contained in this shampoo. (See **What the important nonmedicinal ingredients are**).

What the medicinal ingredient is:

Ciclopirox olamine.

What the important nonmedicinal ingredients are:

Purified water, sodium lauryl ether sulphate, cocamidopropyl betaine, disodium phosphate dodecahydrate, citric acid monohydrate, coconut diethanolamide, hexylene glycol, oleyl alcohol, polysorbate 80, polyquaternium 10, fragrance (including dipropylene glycol), and sodium hydroxide.

What dosage forms it comes in:

STIEPROX[®] is a shampoo with a ciclopirox olamine content of 1.5% w/w. It is available in a 150 mL bottle.

WARNINGS AND PRECAUTIONS

- STIEPROX[®] is for external use only.
- Like any other shampoo, care should be taken to keep STIEPROX[®] out of your eyes and off your eyelids. If contact occurs, flush eyes with copious amounts of water. If discomfort persists, consult your doctor.
- If skin irritation occurs and worsens or persists, consult your doctor.
- Consult your doctor prior to the use of STIEPROX[®] if you are pregnant, planning to become pregnant, nursing or planning to nurse.
- Nursing mothers are advised to ensure that any residual product is fully washed off the breast prior to breast-feeding to avoid possible ingestion by nursing infant.
- Do not use STIEPROX[®] if you have a reaction to ciclopirox olamine.
- Take special care with STIEPROX[®] if you have grey, white, or chemically damaged hair (for example, due to hair dye) as STIEPROX[®] may cause your hair to change colour.

INTERACTIONS WITH THIS MEDICATION

No drug interactions have been reported with ciclopirox olamine.

PROPER USE OF THIS MEDICATION

For use by adults 18 years and older on the scalp and areas next to the hairline only.

Directions for Use

1. Wet the scalp, then apply sufficient shampoo to produce lather.
2. The scalp and its edges should be vigorously massaged with the fingertips.
3. The shampoo should be left on the scalp for 2-3 minutes.
4. The hair should be thoroughly rinsed.
5. Apply a second time by repeating Steps 1 to 4.
6. **Dandruff** - Apply 2 or 3 times a week, or as directed by your doctor.

Seborrhoeic dermatitis - Apply 3 times a week, or as directed by your doctor.

Overdose:

In case of drug overdose, contact a health care practitioner, hospital emergency department or regional Poison Control Centre immediately, even if there are no symptoms.

Missed Dose:

If you forget to shampoo your hair with STIEPROX® on a given day, you should wash your hair with STIEPROX® the next day.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

Side effects may include:

Effects on your skin

Itching* (pruritus), skin and scalp irritation*, skin redness* (erythema), a skin burning sensation*, application site rash*, flaking of the skin*, sores on the scalp, eczema.

Effects on your eyes

Stinging, burning, watering, infection.

Other effects

Headache, hair loss*, hair discolouration, changes to hair texture and/or colour, nausea.

If any other side effects occur or existing side effects worsen, contact your doctor.

*Since these effects are also symptoms of the underlying disease, it is expected that these side effects would manifest as a worsening of symptoms.

HOW TO STORE IT

Store STIEPROX® between 4° and 30°C.

Keep out of reach and sight of children.

REPORTING SUSPECTED SIDE EFFECTS

You can report any suspected adverse reactions associated with the use of health products to the Canada Vigilance Program by one of the following 3 ways:

- Report online at www.healthcanada.gc.ca/medeffect
- Call toll-free at 1-866-234-2345
- Complete a Canada Vigilance Reporting Form and:
 - Fax toll-free to 1-866-678-6789, or
 - Mail to: Canada Vigilance Program
Health Canada
Postal Locator 0701E
Ottawa, ON K1A 0K9

Postage paid labels, Canada Vigilance Reporting Form and the adverse reaction reporting guidelines are available on the MedEffect™ Canada Web site at www.healthcanada.gc.ca/medeffect.

NOTE: Should you require information related to the management of side effects, contact your health professional. The Canada Vigilance Program does not provide medical advice.

MORE INFORMATION

This document plus the full product monograph, prepared for health professionals can be found at:

<http://www.stiefel.ca> or by contacting the sponsor,

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