PRESCRIBING INFORMATION

Pr PROPADERM®
(beclomethasone dipropionate)

Cream 0.025%

Topical Corticosteroid

Valeant Canada LP
2150 St-Elzear Blvd. West,
Laval, Quebec,
Canada H7L 4A8

Date of preparation:
October 23, 2014

Control No: 178645
PRESCRIBING INFORMATION

Pr PROPADERM®
(beclomethasone dipropionate)

Cream 0.025%

Topical Corticosteroid

ACTION AND CLINICAL PHARMACOLOGY

PROPADERM® (beclomethasone dipropionate) Cream is a potent anti-inflammatory steroid. In the vasoconstriction test on human skin, beclomethasone dipropionate is five thousand times as potent as hydrocortisone.

Clinical studies:

It has been shown in clinical trials that the optimal concentration of beclomethasone dipropionate cream is 0.025%.

In two hundred and sixty-seven patients, PROPADERM® Cream and fluocinolone acetonide were used simultaneously in a double-blind fashion. There was no significant difference in the action of the two substances. No side-effects or toxic reactions were reported.

One patient discontinued treatment due to the development of a pruritic eruption during medication with another formulation of PROPADERM® (previously marketed lotion).

INDICATIONS AND CLINICAL USES

PROPADERM® (beclomethasone dipropionate) Cream is indicated for all skin conditions where a topical anti-inflammatory steroid is indicated, including psoriasis, eczema, allergic dermatoses, neurodermatitis, seborrhea, intertrigo, lichen simplex, lichen planus, discoid lupus erythematosus and anogenital pruritus.
CONTRAINDICATIONS

Infected skin lesions if no anti-infective agent is used simultaneously; fungal and viral infections of the skin, including herpes simplex, vaccinia and varicella; pregnancy; hypersensitivity to any of the ingredients. Topical corticosteroids are also contraindicated in tuberculous lesions of the skin.

WARNINGS

PROPADERM® (beclomethasone dipropionate) Cream should not be used in the eye. When topical anti-inflammatory steroids are used under occlusive dressing, over extensive areas, it is possible that sufficient absorption may take place to give rise to systemic effects. Such effects have not been reported with PROPADERM®. Patients should be advised to inform subsequent physicians of the prior use of corticosteroids.

ADVERSE REACTIONS

Local burning, irritation, itching, skin atrophy, striae, hypertrichosis and adrenal suppression have been observed following topical corticosteroid therapy. Posterior subcapsular cataracts have been reported following the systemic use of corticosteroids.

SYMPTOMS AND TREATMENT OF OVERDOSAGE

Discontinuation of therapy, when the typical signs of hypercorticism appear.

DOSAGE AND ADMINISTRATION

PROPADERM® (beclomethasone dipropionate) Cream is applied thinly to cover the affected area and gently rubbed in. Application is usually one to three times daily or as indicated by the severity of the condition. In certain resistant dermatoses the effect of PROPADERM® can be enhanced if the treated area is covered with an occlusive dressing using impermeable material such as polyethylene film of cellophane.
PHARMACEUTICAL INFORMATION

DRUG SUBSTANCE

Common Name: beclomethasone dipropionate
Chemical Name: 9α-chloro-11β-hydroxy-16β-methyl-17α, 21-dipropionyloxy-pregna-1, 4-diene-3, 20-dione

Structural Formula:

HO

CH₃

C₁

C=0

CH₂OCOC₂H₅

Molecular Formula: C₂₈H₃₇ClO₇.
Molecular Weight: 521.04

STABILITY AND STORAGE RECOMMENDATION

PROPADERM® Cream: store between 15° and 25°C

AVAILABILITY OF DOSAGE FORMS

PROPADERM® (beclomethasone dipropionate) Cream: each gram contains beclomethasone dipropionate 0.025%. Available in tubes of 45 g.
REFERENCES


