

# For Moderate to Severe Scalp Dermatoses and Mild to Moderate Psoriasis on the Body

OLUX<sup>®</sup> (clobetasol propionate) Foam, 0.05%



(clobetasol propionate)  
Foam, 0.05%

Please see Important Safety Information on back page  
and accompanying Full Prescribing Information.

OLUX<sup>®</sup> (clobetasol propionate) Foam, 0.05% is a super-potent corticosteroid indicated for short-term topical treatment of the inflammatory and pruritic symptoms of moderate to severe corticosteroid-responsive dermatoses of the scalp, and for short-term topical treatment of mild to moderate plaque-type psoriasis on non-scalp regions excluding the face and intertriginous areas.

The foam is applied for no more than 2 consecutive weeks; patients should use no more than 50 grams per week.

Treatment beyond 2 consecutive weeks is not recommended and the total dosage should not exceed 50 grams per week because of the potential for the drug to suppress the hypothalamic-pituitary-adrenal (HPA) axis. In a controlled pharmacokinetic study, some subjects experienced reversible suppression of the adrenals following 14 days of Olux therapy.



## DISCOUNTS FOR YOUR PATIENTS

### PRESTIUM PRESCRIPTION SAVINGS PROGRAM

With Prestium Pharma's Prescription Savings Program, your patients can save up to \$45 off out-of-pocket costs each time their pharmacist fills or refills the prescription. **Ask for a supply of Prescription Savings Cards (see sample below) to provide your patients with their script.**

The cards are valid at participating pharmacies for one use per medicine per month. Not valid for patients under Medicaid, Medicare (including Medicare Part D), or similar state or federal programs. The card is not valid for residents of Massachusetts unless the patient is paying the full cost of the prescription.

**INSTANT SAVINGS CARD**  
Pay the initial \$15 and receive up to \$45 off your out-of-pocket expenses.

<b>DenaVir<sup>®</sup></b> (periclovir cream, 1%)	<b>ELIMITE<sup>™</sup> CREAM</b> (Pimecrolimus) 5%	<b>evoclin</b> (clindamycin phosphate) foam, 1%	<b>extina<sup>®</sup></b> (ketotifen) foam, 1%
<b>Luciq<sup>®</sup></b> (bimatoprost) foam, 0.12%	<b>Olux<sup>®</sup></b> (olopatadine hydrochloride) foam, 0.025%	<b>Olux-E</b> (olopatadine hydrochloride) foam, 0.025%	<b>Vision<sup>®</sup></b> (ketotifen) foam, 1% (0.25% w/v) (0.25% w/v) (0.25% w/v) (0.25% w/v)

**BIN: 610020    GROUP: 99992198    ID: XXXXXXXXXXXX**  
Please see back of card for details.



## How the Prestium Pharma Combo Savings Card Works for Your Patients:

- Pay the initial \$15, you could receive up to \$45 off your co-pay or out-of-pocket expenses.
- You may pay more than \$15 if your co-pay exceeds \$60; if your insurance does not cover Denavir, Elimite, Evoclin, Extina, Luxiq, Olux, Olux-E, or Vusion; or if you are a cash payer.
- Go to [www.PrestiumPharma.com](http://www.PrestiumPharma.com) for more information.
- For assistance with the Prestium Pharma Combo Savings Card, contact **1-855-820-3232**.
- After each transaction, keep this card for up to 12 total uses.

**APPLY FOR YOUR  
PRESCRIPTION CARD TODAY!**



## IMPORTANT SAFETY INFORMATION

Please see accompanying Full Prescribing Information

- OLUX<sup>®</sup> (clobetasol propionate) Foam, 0.05% is for topical use only. Avoid contact with eyes.
- OLUX Foam is a super-potent topical corticosteroid that has been shown to suppress the adrenals at 7.0 g of OLUX Foam per day. Lesser amounts of OLUX Foam were not studied.
- Systemic absorption of topical corticosteroids has caused reversible adrenal suppression with the potential for glucocorticosteroid insufficiency after withdrawal of treatment. Evaluate patients periodically for evidence of HPA axis suppression.
- Cushing's Syndrome, hyperglycemia, and glucosuria can also result from systemic absorption of topical corticosteroids. Use of OLUX Foam for longer than 2 weeks may suppress the immune system.
- Pediatric patients may be more susceptible to systemic toxicity from equivalent doses due to their larger skin surface to body mass ratios. Use in children under 12 years of age is not recommended.
- OLUX Foam should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.
- Caution should be exercised when OLUX Foam is administered to a nursing woman.
- In clinical trials, the most common adverse events associated with the use of OLUX Foam were burning, dryness, and other reactions at the application site.
- The propellant in this foam is flammable; patients should avoid fire, flame, and/or smoking during and immediately following application.



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