

For Scalp Dermatoses, A Medium-Potency Steroid

LUXIQ® (betamethasone valerate) Foam, 0.12%



(betamethasone
valerate) Foam, 0.12%

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

With its foam delivery system, LUXIQ® (betamethasone valerate) Foam, 0.12% provides a quickly penetrating topical steroid for hair-bearing and hairless areas of the scalp. The foam leaves minimal residue, does not stain, hydrate, or dry the skin, and is free of fragrance, preservatives, and parabens. Indicated for relief of the inflammatory and pruritic symptoms of corticosteroid-responsive dermatoses of the scalp, LUXIQ Foam is prescribed for adult patients.

This medium-potency foam, which disappears rapidly at skin temperature, is applied twice daily for up to 2 weeks.

- If irritation develops, LUXIQ Foam should be discontinued and appropriate therapy instituted. Allergic contact dermatitis with corticosteroids is usually diagnosed by observing a failure to heal rather than noting a clinical exacerbation, as with most topical products not containing corticosteroids.
- Concomitant skin infections should be treated with an appropriate antimicrobial agent. If the infection persists, LUXIQ Foam should be discontinued until the infection has been adequately treated.
- Local adverse reactions that have been reported with topical steroids include: irritation; dryness; folliculitis; acneiform eruptions; hypopigmentation; perioral dermatitis; allergic contact dermatitis; secondary infection; skin atrophy; striae; and malaria.



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.

Denavir (periclovir ointment, 1%)	ELIMITE CREAM (Pimecrolimus) 5%	evoclin (Erdafitinib) foam, 1%	extina (Aflibercept) foam, 1%
Luxiq (Bimatoprost) foam, 0.12%	Olux (Oxycodone hydrochloride) foam, 0.05%	Olux-E (Ezetimibe/ezetimibe) foam, 0.05%	Vision (Timolol maleate) foam, 0.25% off product / 1% on value (0.25% off product) / 1% on value

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 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

- LUXIQ® (betamethasone valerate) Foam, 0.12% is for topical use only.
- Systemic absorption of topical corticosteroids has caused reversible hypothalamic-pituitary-adrenal (HPA) axis suppression with the potential for glucocorticosteroid insufficiency. This may occur during treatment or after withdrawal of the topical corticosteroid.
- Manifestations of Cushing's Syndrome, hyperglycemia, and glucosuria can also be produced in some patients by systemic absorption of topical corticosteroids.
- Conditions that augment systemic absorption include the application of the more potent steroids, use over large surface areas, prolonged use, and the addition of occlusive dressings. Evaluate patients periodically for evidence of HPA axis suppression.
- If irritation develops, LUXIQ Foam should be discontinued and appropriate therapy instituted. Allergic contact dermatitis with corticosteroids is usually diagnosed by observing a failure to heal rather than noting a clinical exacerbation, as with most topical products not containing corticosteroids.
- Concomitant skin infections should be treated with an appropriate antimicrobial agent. If the infection persists, LUXIQ Foam should be discontinued until the infection has been adequately treated.
- LUXIQ Foam should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.
- Caution should be exercised when LUXIQ Foam is administered to a nursing woman.
- Safety and efficacy in pediatric patients have not been established. Pediatric patients may be more susceptible to systemic toxicity from equivalent doses due to their larger skin to surface body mass ratios. Chronic corticosteroid therapy may interfere with the growth and development of children.
- In clinical studies, the most common adverse events associated with the use of LUXIQ Foam were burning, stinging, or itching at the application site.
- Local adverse reactions that have been reported with topical steroids include: irritation; dryness; folliculitis; acneiform eruptions; hypopigmentation; perioral dermatitis; allergic contact dermatitis; secondary infection; skin atrophy; striae; and malaria.
- Because LUXIQ Foam is flammable, you should counsel patients to avoid fire, flame, or smoking during and immediately following application.



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