## Two UVA Sunscreen Filters Are Better Than One

BY BRUCE JANCIN

SAN FRANCISCO — An SPF 40 sunscreen containing two UVA filters—ecamsule and avobenzone—protected patients from flares of polymorphous light eruption significantly better than formulations containing only one UVA filter, in a large outdoor randomized trial.

La Roche-Posay's novel sunscreen, marketed over the counter by L'Oréal as Anthelios 40, contains ecamsule 3%, avobenzone 2%, octocrylene 10% for UVB protection, and titanium dioxide

Flares of polymorphous light eruption occurred later and with a higher cumulative UVA dose with the dual-UVA-filter sunscreen than with either of the single-filter products.

5% as a physical filter providing protection across the UV spectrum.

Ecamsule provides enhanced protection in the short-UVA range, where avobenzone is less effective. Ecamsule protects against UV in the 290- to 400-nm range, with peak protection at 345 nm. It is also more photostable than avobenzone, Dr. Vincent DeLeo explained at the annual meeting of the American Academy of Dermatology.

Dr. DeLeo of St. Luke's-Roosevelt

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were also found on the medial canthus (28%), upper eyelid (10%), and lateral canthus (6%).

Squamous cell tumors also were found most frequently on the lower eyelid (64 or 61%), followed by the medial canthus (17%), the upper eyelid (15%), and the lateral canthus (7%), Dr. Thiele reported.

Six of the 10 melanomas were also on the lower eyelid; 8 of the tumors were in females.

For BCCs, the pre- and postoperative sizes were smallest on the upper eyelids, while the largest tumors were found on the medial canthus. The mean number of Mohs layers needed for BCC clearance ranged from 1.33 in the lateral canthus to 1.42 in the medial canthus.

SCCs had larger pre-op and postop sizes, but the number of layers needed for clearance was lower. The mean number for SCC clearance was 1.5 in the medial canthus and 1.1 in the lateral canthus, Dr. Thiele said.

Although this study confirmed the results of some large Australian data-bases, the chart review found a two fold higher occurrence of SCCs on the upper eyelid than had been reported previously, he noted.

Better knowledge of high-risk histologies and locations of periocular skin cancers should assist surgeons, said Dr. Thiele, who reported no conflicts.

—Alicia Ault

Hospital in New York reported on 144 adult patients with polymorphous light eruption (PMLE) who participated in the randomized, double-blind clinical trial. They applied the dual-UVA-filter sunscreen daily on one side of the body and the same product minus either the ecamsule or avobenzone on the other side. Then they went outdoors for controlled doses of natural sunlight.

The primary study end point was a

composite efficacy measure consisting of delayed time to onset of PMLE or lower global flare severity, based on a 10point scale assessing itching, papules, vesicles, and erythema.

In paired comparisons, the success rate was 56% with the dual-UVA-filter sunscreen, vs. 11% for the ecamsule-deprived sunscreen, and 36% with the dual-UVA-filter sunscreen, compared with 16% for the avobenzone-deprived prod-

uct. Both differences were statistically significant.

Flares of PMLE occurred later and with a higher cumulative UVA dose with the dual-UVA-filter sunscreen than with either of the single-filter products.

L'Oréal, which funded the study, has exclusive patent rights to ecamsule (Mexoryl SX), approved by the Food and Drug Administration in July 2006 as the first new UVA filter in nearly 2 decades.



## myfinacea.com - a website for patient support and education

Finacea is indicated for topical treatment of inflammatory papules and pustules of mild to moderate rosacea. Although some reduction of erythema which was present in patients with papules and pustules of rosacea occurred in clinical studies, efficacy for treatment of erythema in rosacea in the absence of papules and pustules has not been evaluated.

Finacea is for dermatologic use only, and not for ophthalmic, oral, or intravaginal use. Finacea is contraindicated in individuals with a history of hypersensitivity to propylene glycol or any other component of the formulation. In clinical trials, sensations of burning/stinging/tingling occurred in 29% of patients, and itching in 11%, regardless of the relationship to therapy. Post-marketing safety—Skin: facial burning and irritation; Eyes: iridocyclitis on accidental exposure to the eye. There have been isolated reports of hypopigmentation after use of azelaic acid. Since azelaic acid has not been well studied in patients with dark complexion, these patients should be monitored for early signs of hypopigmentation.

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