CASE REPORT

Intense Pulsed Light Treatment of Persistent Facial Hypermelanosis Following Drug-Induced Toxic Epidermal Necrolysis

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BACKGROUND. Cutaneous hyperpigmentation is one of the most cosmetically disturbing sequel of drug-induced toxic epidermal necrolysis. Intense pulsed light is a promising tool for treating some melanocytic lesions.

OBJECTIVE. The objective was to assess the effect of intense pulsed light in treating post-toxic epidermal necrolysis facial hypermelanosis.

METHODS. Two Caucasian men aged 35 and 50 years presented with long-standing (32 and 39 years) severe hypermelanosis of the face after sulfonamide-induced toxic epidermal necrolysis. They were treated by intense pulsed light. Cutoff filters of 550, 590, and 615 nm were employed for five intense pulsed light sessions at 4-week intervals. The treatment was characterized by energy fluence of 25 to 32 J/cm², pulse width of 2.2 to 3.2 ms, and double- to triple-pulse mode respecting a 30-ms delay. Before intense pulsed light treatment, and 2 months after the fifth intense pulsed light session, clinical photographs and skin biopsies were performed in combination with quantitative narrow-band remittance spectrophotometry of melanin pigmentation. Patients were clinically followed-up for 8 months after the end of the treatment.

RESULTS. In both patients, clinical, histologic, and spectrophotometric assessments showed an average of 80% decrease in the hypermelanosis. No clinical recurrence of the hypermelanosis developed during the 8-month follow-up after intense pulsed light treatment. No major persistent side effects were experienced, especially hypopigmentation.

CONCLUSION. Intense pulsed light appears to be effective and safe for treating post-toxic epidermal necrolysis hypermelanosis in Caucasian patients.

PHILIPPE PAQUET, MD, PHD, AND GÉRALD E. PIÉRARD, MD, PHD HAVE INDICATED NO SIGNIFICANT INTEREST WITH COMMERCIAL SUPPORTERS.

DRUG-INDUCED TOXIC epidermal necrolysis is a rare, potentially lethal disease characterized by sudden epidermal necrosis. The only recognized cause of the disease is an adverse reaction to drugs, especially sulfonamides, phenytoin, nonsteroidal anti-inflammatory drugs, and allopurinol. The toxic epidermal necrolysis–induced death rate reaches approximately 30%. Long-term sequelae in the survivors remain frequent. Pulmonary, myocardial, osseous and gastrointestinal complications have also been described. The most common long-term consequences, however, affect the eyes, skin, oral and genital mucosae, and nails. Cutaneous sequelae include keloidal scars, alopecia, speckled eruptive melanocytic nevi, and hyper- or hypomelanosis.

We report two adult patients presenting with long-standing hypermelanosis of the face following toxic epidermal necrolysis that developed during childhood. The hyperpigmentation was treated successfully by intense pulsed light. To the best of our knowledge, these cases are the first descriptions of post-toxic epidermal necrolysis hyperpigmentation treated by intense pulsed light. We also briefly discuss the pathogenesis of post-toxic epidermal necrolysis hypermelanosis.

Case Report

Two phototype II Caucasian men, aged 35 and 50 years, suffered from toxic epidermal necrolysis at the age of 3 and 11 years, respectively. The clinical presentation was similar for the two patients, including epidermal sloughing over 50% of the body surface, and severe involvement of the face, eyes, and mucous membranes. Trimethoprim-sulfamethoxazole of the sulfonamide-class antibiotic was identified as the culprit drug in both cases. Cutaneous biopsies taken at the onset of the disease confirmed the diagnosis of toxic epidermal necrolysis. After the early recovery phase of the disease, facial skin showed a scar texture with a persistent diffuse and severe hypermelanosis, particularly prominent on the cheeks. By contrast, the
other parts of the body surface fully recovered without any pigmentary change.

An intense pulsed light device (Multilight\textsuperscript{T}, ESC Medical Systems Ltd, Yokneam, Israel) was used to treat the hypermelanosis in a five-session procedure with respective intervals of 4 to 6 weeks. Cutoff filters of 550, 590, and 615 nm were used. To deal with the superficial pigmentation, treatments started using the 550-nm cutoff filter, a single pulse mode at 25 J/cm\textsuperscript{2} fluence, and a pulse width of 3.6 ms. To tackle deeper pigmentation when previous attempts with the shorter cutoff filter failed to clear the melanoderma, the cutoff filter was switched to 590 or 615 nm. A double- or triple-pulse mode combined with a pulse width of 3.2 or 2.2 ms, a 30-ms delay, and a fluence of 25 to 30 and 26 to 32 J/cm\textsuperscript{2} were employed using 590- and 615-filters, respectively. The two cheeks were entirely treated with a 2.8 × 0.8-cm spot size, representing 25 to 35 pulses by session. The intense pulsed light spots overlapped by about 10%. Before treatment, skin cooling was obtained using cold packs for 1 min. In addition, a 2-mm-thick coating with a chilled colorless gel was applied to the treatment area to protect the epidermis from thermal injury and to help deliver the light uniformly at the skin surface. No local anesthesia was used. Patients were advised to apply a corticosteroid and antiseptic combination cream (Fucicort, Leo) twice daily for 7 days. In the period covering 2 months before treatment, the whole treatment phase, and 2 months after treatment, the patients were advised to avoid sun exposure and they regularly applied a sun-protective (SPF 60) product.

The treatment response was assessed by comparing pre- and posttreatment clinical photographs, cutaneous biopsies, and data from remittance spectrophotometry. Posttreatment evaluations were performed 2 and 8 months after the fifth intense pulsed light session. This period covered the summer season. Pre- and posttreatment biopsy specimens were taken close to each other. Sections were stained with Fontana-Masson silver stain to reveal melanin. A narrow-band spectrophotometer (Mexameter, C+K Electronic, Cologne, France) was used according to the EEMCO recommendations.\textsuperscript{5} The melanin index was measured on two different sites of the cheeks. Melanin index assessments were precisely performed each time on the same sites. To avoid any influence of skin color variation due to skin tanning, drug and alcohol intake, physical exercise, and environmental temperature, results were expressed as a differential melanin index (ΔM) between each of the melanin index values obtained at the target sites and another selected area of normal looking skin on the anterior part of the neck. M and ΔM values were expressed in arbitrary units. The posttreatment ΔM value was also calculated as a percentage relative to the pretreatment value.

Results

Deep purpuric spots developed rapidly after each intense pulsed light session. In some areas, especially at the beginning of the treatment, the patients developed blisters and crusts that resolved within 1 week leaving some discrete and transient depressed scars. After 3 months, the scars had cleared without leaving permanent marks.

The clinical assessment of treatment efficacy was supported by comparing the photographs. Overall, the two patients showed a marked clinical improvement of melanization after the fifth intense pulsed light session (Figure 1). The treatment did not induce any obvious hyper- or hypopigmentation. Eight months after the last intense pulsed light session, hypermelanosis did not recur even following summer.

The spectrophotometer assessments showed a sharp decrease in ΔM (range, −17% to −117%; mean, −84%) at completion of the treatment in both patients (Table 1). In each case, the initial histologic examination revealed a discrete superficial perivascular lymphoid infiltrate. The melanin overload in the deepest layers of the epidermis was accompanied by few melanophages in the superficial dermis. The Fontana-Masson stain was strongly positive. After intense pulsed light treatment, a marked decrease of the intraepidermal melanin load was evidenced, but the reduction in dermal melanophages was less obvious (Figure 2).

**Figure 1.** Post-toxic epidermal necrolysis facial hypermelanosis in Patient 2 before intense pulsed light treatment (A) and after five intense pulsed light sessions (B).
Discussion

Changes in skin pigmentation including hypo- and hypermelanosis are among the most common disturbing cutaneous sequelae of toxic epidermal necrolysis.\(^3,4\) The modifications in the melanocyte function are mainly seen on sun-exposed skin, although actinic exposure does not seem mandatory for their development.\(^4\)

The precise pathogenesis of post-toxic epidermal necrolysis hyperpigmentation is still obscure. In many inflammatory diseases, epidermal melanocytes increase in number, size, dendricity, and melanin production. Arachidonic acid metabolites (prostaglandin E2, thromboxane B2, leukotrienes C and D4) and histamine, which are increased in amount in inflammatory conditions, are thought to play a key role in the induction of postinflammatory hypermelanosis.\(^6,7\) The vasoconstriction peptide endothelin-1 is also a strong keratinocyte-derived mitogen and melanogen for human melanocytes.\(^8\) Endothelin-1 secretion is stimulated by tumor necrosis factor-\(\alpha\), which is one of the main cytokines involved in toxic epidermal necrolysis pathogenesis.\(^8\) Nitric oxide might also be involved in the process.\(^9\) Indeed, any nitric oxide excess induces keratinocyte apoptosis and boosts melanocyte activity.

A direct effect of drug metabolites on epidermal melanocytes may also be involved in the increased melanin production. Indeed, contact between unirradiated melanocytes in culture and DNA-damaging agents or thymine dinucleotides, the by-products of DNA excision repair, is sufficient to increase the melanin content and up regulate tyrosinase gene expression.\(^10\) In addition, tyrosinase expression may be up regulated by removal of a repressor gene given the tyrosinase is under negative control in melanocytes.\(^10\)

In our patients, the long-standing hypermelanosis resulted from toxic epidermal necrolysis developed during childhood and was exclusively located on the face. It should be noted that in children aged between 1 and 15 years, the number of active melanocytes is higher than in older subjects. The number of melanocytes also varies according to the region of the body, the cheeks exhibiting one of the highest densities.\(^11\) The persistence of hypermelanosis on sun-exposed skin in our patients suggests an ultraviolet-sustained pathomechanism. Moreover, the sulfonamides which are thought to be responsible for the presently reported toxic epidermal necrolysis cases are photosensitizing drugs.\(^12\) The mechanisms involved in ultraviolet-induced hypermelanosis include melanocyte DNA damage and/or repair, keratinocyte-derived hyperproduction of the melanogenic factor endothelin-1, or the destruction of tyrosinase gene repressor protein PRP (photolyase regulatory protein).\(^10\)

Dermal postinflammatory hypermelanosis results from melanin incontinence from the epidermis. Melanosomes are captured by dermal macrophages and dendrocytes.\(^12\) Among them, Factor XIIIa+ dendrocytes, whose population is particularly abundant in

**Table 1. Spectrophotometry Assessments of the Differential Melanin Index between the Hypermelanotic Cheeks and the Normal Looking Neck, Before and After Five Intense Pulsed Light Treatments**

<table>
<thead>
<tr>
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<th>Left Cheek</th>
<th>Right Cheek</th>
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<tbody>
<tr>
<td></td>
<td>Before Intense Pulsed Light</td>
<td>After Intense Pulsed Light</td>
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<tr>
<td><strong>Patient 1</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>External cheek</td>
<td>658</td>
<td>325</td>
</tr>
<tr>
<td>Internal cheek</td>
<td>506</td>
<td>422</td>
</tr>
<tr>
<td><strong>Patient 2</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>External cheek</td>
<td>715</td>
<td>–6</td>
</tr>
<tr>
<td>Internal cheek</td>
<td>1043</td>
<td>–146</td>
</tr>
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**Figure 2.** Fontana-Masson stain of cutaneous biopsies in Patient 2 before (a) and after five intense pulsed light sessions (b). Melanin deposits appear in black.
toxic epidermal necrolysis skin, have been shown to be able to store the released melanin products.\textsuperscript{13}

Different kinds of lasers (Q-switched ruby, alexandrite, or Nd-Yag)\textsuperscript{14–19} and intense pulsed light\textsuperscript{20–26} have been reported to be effective in treating some pigmented lesions, especially postinflammatory hyperpigmentation. There is, however, no information about the efficacy of intense pulsed light in treating post-toxic epidermal necrolysis hypermelanosis. The mechanism of action of these methods relies on the selective photothermolysis of pigmented cells. In contrast to lasers, intense pulsed light system offers the advantage of adjusting pulse width and wavelength according to the skin type and depth of pigment deposits in the skin.\textsuperscript{20–26} In our patients, the melanin overload was mostly located in the epidermis. Thus, short wavelengths (cutoff filter of 550 and 590 nm) were used preferentially. Clinical, histologic, and spectrophotometric assessments showed a prominent bleaching effect after five intense pulsed light sessions. This looks similar to the results obtained in treating postburn hypermelanosis especially on sun-exposed skin.\textsuperscript{20–26} In our patients, the melanin overload was mostly located in the epidermis. Thus, short wavelengths (cutoff filter of 550 and 590 nm) were used preferentially. Clinical, histologic, and spectrophotometric assessments showed a prominent bleaching effect after five intense pulsed light sessions. This looks similar to the results obtained in treating postburn hypermelanosis especially on sun-exposed skin.

In conclusion, toxic epidermal necrolysis can be followed by persistent postinflammatory hypermelanosis especially on sun-exposed skin. Intense pulsed light appears to be a promising treatment for post-toxic epidermal necrolysis epidermal hyperpigmentation, with the possibility to precisely adjust the beam depth penetration according to the site of melanin deposits in the skin.

References