Periarticular Thenar Erythema and Onycholysis Syndrome: A Manifestation of Taxane-Induced Cutaneous Toxicity

Síndrome de eritema tenar periarticular y onicolisis: una manifestación de toxicidad cutánea por taxanos

To the Editor:

In recent years, the widespread use of taxane-based chemotherapy regimens for the management of malignant tumors of various origins has evidenced the high frequency of cutaneous toxicity induced by this pharmacological group. In 2003, among its possible manifestations, Childress and Lokich defined periarticular thenar erythema and onycholysis (PATEO) syndrome as a rare subtype of hand-foot syndrome associated with nail changes in patients treated with paclitaxel and docetaxel. In contrast to typical palmar-plantar erythrodysesthesia, or classic hand-foot syndrome, skin involvement in this syndrome is characterized by an absence of involvement of the palms and soles and by the presence of alterations predominantly affecting the dorsum of the hands (particularly around the base of the thumb and fifth finger), the area of the Achilles tendon, and the perimaleolar skin.

We present 3 patients on chemotherapy with taxanes who developed clinical manifestations similar to those described in this syndrome. The 3 patients were women aged 35, 56, and 68 years, with infiltrating ductal adenocarcinoma of the breast on adjuvant treatment with docetaxel and cyclophosphamide (Table 1). They sought dermatologic evaluation for the appearance of painful erythematous-violaceous plaques with fine superficial desquamation on the dorsum of both hands (Fig. 1A). The lesions interfered with their activities of daily living. The lesions had arisen 3 to 12 days after the administration of chemotherapy, and showed a clearly progressive deterioration with each cycle, despite the use of topical corticosteroids. Associated nail involvement in the 3 cases varied from fine parallel Beau lines and subungual splinter hemorrhages to frank distal onycholysis of the nail plate (Fig. 1B). Physical examination revealed no lesions on the rest of the skin, and the patients presented an otherwise good general state of health. Histology of a 4-mm punch biopsy from the dorsum of the hand of patient number 2 revealed a hyperkeratotic epidermis with parakeratosis, necrotic keratinocytes, focal vascular degeneration of the basal layer, and a band-like perivascular lymphocytic inflammatory infiltrate with occasional eosinophils (Fig. 2, A and B). These findings were consistent with a clinical suspicion of taxane-induced toxicity. Dose adjustment of the chemotherapy, together with the application of topical corticosteroid therapy under occlusion controlled the lesions and significantly improved symptoms in all 3 patients. Only 1 patient required a single course of a tapered regimen of oral corticosteroids for a severe deterioration of the lesions on the dorsum of the hands that interfered with her activities of daily living.

PATEO syndrome is a manifestation of taxane-induced cutaneous toxicity predominantly affecting acral areas and with associated nail alterations. The frequency of this complication has been reported as 10% and 5%, respectively, in patients treated with paclitaxel and docetaxel, and its intensity appears to increase with the cumulative dose and with weekly regimens. Skin involvement is characterized by the presence of painful erythematous-violaceous plaques on the dorsum of the hands, on the feet, and in the area of the Achilles tendon; these lesions can subsequently develop vesicles and flaking. Severity is very variable, from asymptomatic forms to ones that have a marked impact on the basic activities of daily living. Some classifications, such as those of the World Health Organization and the National Cancer Institute, have established criteria of severity to evaluate acral skin involvement.

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Figure 1  Periarticular thenar erythema and onycholysis syndrome: Docetaxel-induced skin and nail changes. A, Patient 1. Erythematous-violaceous plaques on the dorsum of the hands and over the proximal phalanx. B, Nail changes in patient 3: parallel Beau lines and onycholysis of the distal third of the nail plate.

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Table 1  Epidemiologic, Clinical, and Therapeutic Details of 3 Cases of Periarticular Thenar Erythema and Onycholysis Syndrome.

<table>
<thead>
<tr>
<th>Sex</th>
<th>Age, y</th>
<th>Tumor</th>
<th>Adjuvant Chemotherapy</th>
<th>Latency\textsuperscript{a}</th>
<th>Cumulative Dose\textsuperscript{b}</th>
<th>NCI Grade\textsuperscript{c}</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Female</td>
<td>56</td>
<td>Infiltrating ductal adenocarcinaoma of the right breast, T2N1M0</td>
<td>Docetaxel + cyclophosphamide</td>
<td>12 d</td>
<td>Third cycle</td>
<td>2</td>
</tr>
<tr>
<td>2</td>
<td>Female</td>
<td>35</td>
<td>Infiltrating ductal adenocarcinaoma of the left breast, T2N0M0</td>
<td>Docetaxel + cyclophosphamide + G-CSF</td>
<td>3 d</td>
<td>Second cycle</td>
<td>3</td>
</tr>
<tr>
<td>3</td>
<td>Female</td>
<td>68</td>
<td>Infiltrating ductal adenocarcinaoma of the left breast, T2N0M0</td>
<td>Docetaxel + cyclophosphamide + G-CSF</td>
<td>8 d</td>
<td>Second cycle</td>
<td>2</td>
</tr>
</tbody>
</table>

Abbreviations: G-CSF, granulocyte colony stimulating factor; NCI, National Cancer Institute.
\textsuperscript{a} Time to onset of symptoms after the most recent administration of chemotherapy.
\textsuperscript{b} Chemotherapy cycle being administered when symptoms first arose.
\textsuperscript{c} National Cancer Institute classification of the severity of acral erythema.\textsuperscript{4,7}

Nail changes in these patients are due to the cytotoxic effect of taxanes and their presence is characteristic of this condition, in contrast to other hand-foot syndromes secondary to anthracyclines or multikinase inhibitors.\textsuperscript{5} The cytotoxicity of these agents is due to their inhibition of microtubule function in cell division, and is most intense in those epithelia with high rates of cell proliferation, such as the nail matrix. The most common changes described after the administration of these drugs are Beau lines, onychomadesis, onycholysis, subungual splinter hemorrhages, subungual hematomas, leukonychia, paronychia, and nail hyperpigmentation.\textsuperscript{4,5} Although the clinical manifestations are not usually severe, some manifestations, such as subungual hematomas, abscesses, and distal onycholysis can cause significant morbidity in oncologic patients and must be taken into consideration.\textsuperscript{6}

Histology of the skin lesions reveals changes similar to those observed in typical palmar-plantar erythrodysesthesia induced by other chemotherapeutic agents. It is characterized by the presence of a hyperkeratotic and acanthotic epidermis, with necrotic keratinocytes, focal vacuolar degeneration of the basal layer, blood-vessel dilatation, and a lichenoid inflammatory infiltrate with few lymphocytes, neutrophils, or eosinophils.\textsuperscript{3,7}

In clinical practice, the impact of PATEO syndrome on the quality of life of oncology patients is a frequent cause of dose reduction of the chemotherapy regimen, as occurred in our patient number 2. In the most severe and disabling cases, discontinuation of treatment may even be indicated.\textsuperscript{8} Preventive measures are an essential strategy to reduce the morbidity associated with these treatments in debilitated patients. Local hypothermia using socks or gloves during infusions of the drug can be effective, though this cannot be extended to other chemotherapy regimens with oral administration.\textsuperscript{9,10}

Given the high frequency of dermatologic consultations related to the side effects of chemotherapy in oncology patients, we consider it important to be aware of this

Figure 2  Histology of the lesion on the dorsum of the hand of patient 2. A, Hyperkeratotic epidermis with parakeratosis, foci of vacuolar degeneration of the basal layer, and a band-like perivascular infiltrate with occasional eosinophils. Hematoxylin and eosin, original magnification ×10. B, At higher power, evidence of necrotic keratinocytes in the epidermis. Hematoxylin and eosin, original magnification ×40.
atypical form of acral erythema and of its management as, in disabling cases, it can require dose reduction or a temporary interruption of the chemotherapy treatment. It is notable that all 3 of our patients were on combined treatment with cyclophosphamide and docetaxel for adenocarcinoma of the breast, and the possibility that the combination may have increased the cutaneous morbidity compared with the administration of docetaxel in monotherapy cannot therefore be excluded.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

References


E. Rodríguez-Lomba,* I. Molina-López, R. Suárez-Fernández, O. Baniandrés-Rodríguez

Servicio de Dermatología, Hospital General Universitario Gregorio Marañón, Madrid, Spain

* Corresponding author.
E-mail address: enriquerlomba@outlook.com (E. Rodríguez-Lomba).
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Oral Oxybutynin for Local and Multifocal Hyperhidrosis: A Multicenter Study

Estudio multicéntrico sobre el uso de oxibutinina oral en hiperhidrosis local y multifocal

To the Editor,

Although the psychosocial consequences of hyperhidrosis are well known, the impact of this disease has traditionally been underestimated by the medical community. Oxybutynin chloride is an effective, safe, and well-tolerated treatment for hyperhidrosis that has been increasingly used since 2006. Most authors recommend a starting dose of 2.5 mg/d and a maximum dose of 15 mg/d. The safety of oxybutynin has been demonstrated in patients with hyperhidrosis.1 However, despite its effectiveness, favorable adverse effect profile, and affordable price, this drug is not as widely used as would be expected.

We present a series of 56 patients with primary hyperhidrosis treated with oxybutynin (5-mg tablets) at 5 Spanish hospitals between May 2013 and February 2016. The patients began by taking half a tablet at breakfast and another half at lunchtime for a week. When this dosage did not achieve control of the sweating, the daily dose was increased by 2.5 mg and maintained for a week. This increase was repeated weekly until a maximum dose of 15 mg/d was reached.

The following variables were studied: sex, age, hyperhidrosis sites (palms and axillae, soles and axillae, palms and soles), starting dose (5 mg in all cases), maintenance dose, adverse effects, and, when reported, the adverse effect that caused the greatest discomfort. Patients aged over 14 years with hyperhidrosis at one or more sites that had not been treated with anything other than topical agents were included. Exclusion criteria were failure to meet the previous criteria, a contraindication for treatment with oral anticholinergics, and previous use of iontophoresis, botulinum toxin, or systemic drugs to treat their condition. All the patients or their legal representatives signed an informed consent form agreeing to the off-label use of oxybutynin.

The patients were assessed with the Hyperhidrosis Disease Severity Scale (HDSS) at the start of treatment and

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