Successful Treatment of Leiomyoma of the Nipple With Carbon Dioxide Laser

Buenos resultados tras tratamiento con láser CO₂ de leiomioama en el pezón

To the Editor:

Genital leiomyomas are the least common type of leiomyomas and those located on the nipple or areola are extremely rare.¹⁻⁴ To date, their treatment with carbon dioxide (CO₂) laser has not been reported in the literature.

We describe the case of a 33-year-old woman with no relevant past medical history who presented with a skin lesion on the left nipple; the lesion had been stable since its appearance in adolescence. The patient reported mild tenderness and great impact on personal appearance as the reasons for seeking treatment. The examination revealed elongation of the left nipple secondary to the presence of a well-defined, firm nodular lesion that was the same color as the nipple and about 15 mm in diameter; there were no signs of nipple retraction (Fig. 1). The rest of the left breast and the right breast revealed no significant findings.

Histopathology of the lesion revealed a hyperkeratotic epidermis with increased pigmentation of the basal layer and, in the dermis, the presence of a poorly defined, unencapsulated tumor composed of multiple interwoven layers of smooth muscle formed by cells with ovoid nuclei and eosinophilic cytoplasm (Fig. 2A). No atypical cells or mitoses were observed. Staining for smooth muscle actin was positive (Fig. 2B). In view of these clinical and histologic findings, the lesion was diagnosed as a genital leiomyoma. Under local anesthesia with mepivacaine 2%, the rest of the lesion was removed with a Sharplan CO₂ laser in unfocused mode with an irradiance of 55 W/cm² and a spot size of 3 mm. The lesion was treated in a single session with 5 to 8 passes; between each pass the coagulated tissue was removed until there was no visible sign of the leiomyoma. After 1 treatment session much of the lesion had been removed with good cosmetic and functional results. No recurrence was detected 20 months after treatment (Fig. 3).

Cutaneous leiomyomas are rare, benign tumors. They are classified into 3 categories according to the muscle fibers from which they originate: piloleiomyoma, originating from the smooth muscle fibers of the arrector pili muscle; angioleiomyoma, originating from the tunica media of the blood vessels; and dartos or genital leiomyoma, originating from the smooth muscles of the scrotum, vulva, nipple, or areola. Genital leiomyomas are the least common type and those located on the nipple or areola are extremely rare.¹⁻⁴ Leiomyomas of the nipple and areola, first described in 1854 by Virchow, present as small, solitary nodular lesions covered with healthy skin that can exceptionally cause nipple retraction and spread to the underlying breast tissue.⁴ The lesions are usually asymptomatic but in rare cases can cause pain spontaneously or in response to stimuli such as cold, pressure, or emotional stress.

Histologically, they are poorly defined, unencapsulated skin tumors composed of bundles of spindle cells with abundant eosinophilic cytoplasm and elongated nuclei with blunt ends, arranged in an interwoven pattern. The epidermis is usually unaffected because it is separated from the tumors by a free zone of dermis; cellular mitosis and atypia are

Figure 1 Skin lesion located on the patient’s left nipple.

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¹ Please cite this article as: López V, López I, Alcacer J, Ricart JM. Buenos resultados tras tratamiento con láser CO₂ de leiomioama en el pezón. Actas Dermosifiliogr. 2013;104:928–930.
Leiomyomas are positive for vimentin, desmin, and smooth muscle actin.

Differential diagnosis with the following diseases should be considered: Paget disease, fibroma, adenoma of the nipple, lymphadenosis benigna cutis, leiomyosarcoma of the areola, erosive adenomatosis of the nipple, and myoid hamartoma.²

Several treatment modalities are used. In cases involving pain it can be useful to use calcium channel blockers or α-blockers, which inhibit contraction of muscle fibers. Isolated cases have been treated with cryotherapy, nitroglycerin, hyoscine hydrobromide, lidocaine, and analgesics.³ In addition to medical treatment, surgical excision is also an option. However, surgery is sometimes contraindicated because of patient factors or because the lesions, as in our patient, are located in areas that require a good cosmetic result. CO₂ laser removal has been used to treat cutaneous leiomyoma with good results,³ but to date no cases of genital leiomyoma treated with this modality have been reported. CO₂ laser emits light in the infrared spectrum at a wavelength of 10 600 nm, which is absorbed by water. The laser vaporizes the skin and causes coagulative necrosis in the remaining dermis. CO₂ laser treatment is advantageous mainly because the procedure is rapid and enables precise selection of tissue. The technique is safe, as tissue damage is specific, with highly localized destruction, minimal damage to surrounding tissue, and minimal bleeding because it promotes coagulation. An alternative to the CO₂ laser is the erbium-doped yttrium aluminum garnet (Er:YAG) laser, which emits light in the infrared spectrum at a wavelength of 2900 nm and can vaporize the skin at a lower penetration depth than the CO₂ laser. The Er:YAG laser produces little thermal damage to the underlying tissue because of its high water absorption coefficient. However, this effective absorption limits the transmission of heat to the deep layers, where it may be insufficient to promote coagulation and prevent bleeding during surgery.

In conclusion, we have presented a case of a rare disease in which CO₂ laser treatment provided very satisfactory cosmetic results. We believe that this technique can be considered a good option in cases in which surgery is not possible or is contraindicated.

References


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Acute Generalized Exanthematous Pustulosis Induced by Galantamine

Pustulosis exantemática aguda generalizada inducida por galantamina

To the Editor:

At present, Alzheimer disease has no effective treatment due to its complex pathophysiology, which is not yet completely understood. However, abnormal cholinergic function has been shown to play an important role in the many types of brain neurotransmitter dysfunction found in Alzheimer patients. Accordingly, cholinesterase inhibitors that strengthen cholinergic neurotransmission, including galantamine, donepezil, and rivastigmine, are considered first-line drugs for the treatment of mild to moderate Alzheimer disease.

We present the first reported case of acute generalized exanthematous pustulosis (AGEP) following the administration of galantamine.

An 81-year-old woman with a history of hypertension, dyslipidemia, spondyloarthropathy, and recently diagnosed Alzheimer-type degenerative dementia was brought to our department for emergency examination. She presented with a generalized itchy skin rash that had developed 5 days earlier, associated with a sensation of poor temperature regulation and no measurable fever.

The physical examination showed large erythematous, edematous plaques on the trunk and limbs with multiple nonfollicular monomorphic pustules. The lesions on the limbs were more isolated and had a target morphology (Fig. 1). Small isolated areas of superficial epidermal detachment were also observed.

No involvement of the oral or genital mucosa or ocular discomfort was observed. Palpation did not reveal visceromegaly or enlarged lymph nodes.

The only relevant recent change in medication was the addition of galantamine, which had been prescribed by the neurologist in the preceding month to treat the patient’s dementia. Galantamine was being administered in increasing doses and the dosage of all other drugs had not changed.

Based on the suspected diagnosis of AGEP induced by galantamine, the patient was admitted to hospital for monitoring and treatment with the drug was discontinued. Her symptoms were treated with fluid therapy, oral antihistamines, and local dressings.

Histology revealed subcorneal pustules filled with neutrophils, edema in the papillary dermis, and a slight perivascular infiltrate of lymphocytes and eosinophils (Fig. 2).

Blood tests showed intense leukocytosis (38 × 109/L) with neutrophilia (34.6 × 109/L), normal blood cell morphology, and increased levels of acute phase reactants, including fibrinogen and C-reactive protein.

Figure 1 Erythematous, edematous plaques with multiple nonfollicular pustules.

Please cite this article as: Pérez-Garcia MP, Sánchez Motillas JM, Mateu-Puchades A, Díaz-Corpas T. Pustulosis exantemática aguda generalizada inducida por galantamina. Actas Dermosifiliogr. 2013;104:930-931.