

ÚLCERA DA ASA DO NARIZ - UM CASO INVULGAR

Guida Santos¹, Alexandre João²

¹Interna do Internato Complementar de Dermatologia e Venereologia/ Resident of Dermatology and Venereology, Serviço de Dermatologia e Venereologia/Dermatology Department, Hospital Santo António dos Capuchos - Centro Hospitalar de Lisboa Central, Lisboa

²Assistente Hospitalar de Dermatologia e Venereologia/Consultant of Dermatology and Venereology, Serviço de Dermatologia e Venereologia/Dermatology Department, Hospital Santo António dos Capuchos - Centro Hospitalar de Lisboa Central, Lisboa, Portugal

RESUMO – A dermite *artefacta* é uma doença caracterizada por lesões cutâneas auto-induzidas, resultado ou manifestação de perturbações psiquiátricas. As manifestações clínicas são variáveis e compreendem erosões superficiais ou úlceras profundas em áreas acessíveis ao alcance das mãos. Pela sua raridade e polimorfismo das lesões, a dermatite *artefacta* é frequentemente um desafio para os médicos. Descreve-se o caso de uma mulher de 62 anos com úlcera do nariz com três anos de evolução causada por manipulação digital.

O reconhecimento precoce da dermatite *artefacta* é difícil mas permite evitar tratamentos desnecessários. É necessária uma abordagem pluridisciplinar desta entidade de modo a obter os melhores resultados.

PALAVRAS-CHAVE – Dermite *artefacta*; Úlcera; Depressão.

A NONHEALING ULCER ON THE NOSE - A CASE REPORT

ABSTRACT – *Dermatitis artefacta* is a disease characterized by self-inflicted skin lesions as the result or manifestation of psychiatric disorders or specific stress situations.

Clinical manifestations range from superficial erosions to deep wounds. Because of its rarity and the polymorphism of the lesions, *dermatitis artefacta* is often a challenge for the clinicians. This report presents the case of a 62-year-old woman who had an ulcer of the nose lasting for three years caused by digital manipulation. Early recognition of *dermatitis artefacta* is difficult but avoids unnecessary treatments. A multidisciplinary approach to this entity is necessary to obtain the best results.

KEY-WORDS – *Dermatitis*; *Ulcer*; *Nose diseases*; *Depression*.

Conflitos de interesse: Os autores declaram não possuir conflitos de interesse.

No conflicts of interest.

Suporte financeiro: O presente trabalho não foi suportado por nenhum subsídio ou bolsa.

No sponsorship or scholarship granted.

Direito à privacidade e consentimento escrito / Privacy policy and informed consent: Os autores declaram que pediram consentimento ao doente para usar as imagens no artigo.

The authors declare that the patient gave written informed consent for the use of its photos in this article.

Recebido/Received – Fevereiro/February 2013; Aceite/Accepted – Março/March 2013

Caso Clínico

Correspondência:

Dr.ª Guida Santos

Serviço de Dermatologia

Hospital Santo António dos Capuchos - Centro Hospitalar de Lisboa Central

Alameda Santo António dos Capuchos

1169-050 Lisboa, Portugal

Email: guidadossantos@gmail.com

INTRODUCTION

Dermatitis *artefacta* is a rare artifactual skin disease¹⁻³ with an etiopathology that is not completely understood^{1,3} and the prevalence may be higher than perceived^{1,2}. It is caused entirely by the actions of the fully aware patient on the skin, hair, nails or mucosa, with no rational motive for this behavior⁴. The condition is more common in women¹⁻³. The lesions are usually bilaterally symmetrical, within easy reach of the dominant hand, and may have bizarre shapes with sharp geometrical borders^{2,4}. Patients may induce lesions by rubbing, scratching, picking, cutting, punching, sucking, biting, injecting substances, applying dyes, heat or caustics or using some instruments¹⁻⁴. Reported associated conditions include obsessive-compulsive disorder, borderline personality disorder, depression and psychosis¹⁻⁵.

CASE REPORT

A sixty six-year-old caucasian woman was referred to our Department of Dermatology because of deep ulcerative lesion involving the left nasal ala associated to intense pruritus, without hypoesthesia, lasting for 2 years. The patient appeared remarkably unconcerned and only answered with difficulties to the questions of the anamnesis, explaining poorly her symptoms. During the first consultation, she related that the lesion has begun after the use of a nasal canula during a hospitalization for laterobulbar stroke 2 years before, and that failed to heal and gradually expanded. Her past medical history was significant for renal polycystic disease with right nephrectomy, leading to terminal renal insufficiency. Her father and her two sisters died of renal polycystic disease.

At presentation, the patient was noted to be missing approximately half of the left nasal ala. In its stead, there was erosion surrounded by hemorrhagic crusting and a margin of mild erythema and edema. The erosion was indurated but non tender. The nares appeared to have some mucosal inflammation (Fig. 1). In order to

exclude a cutaneous tumor and others dermatological disorders, a punch biopsy was performed and the patient was treated with fusidic acid cream. The histopathological examination showed an unspecific ulceration with dense inflammatory infiltrate and did not define neither malignant-cellular proliferation nor granulomatous infiltration (Fig. 2). A biopsy of the lesion revealed reactive hyperplasia of the dermis with neither malignant-cellular proliferation nor granulomatous infiltration. The ulcer continued to progress. Intralesional infiltration of corticosteroids was tried to diminish the itch and the inflammation with slight and temporary improvement. Because of the progression of the lesion, an incisional biopsy were performed and revealed a dense inflammatory infiltrate, consistent with unspecific ulceration. The patient's mood was even lower leading to be



Fig. 1 - Defect of the left nasal ala with slight inflammation.

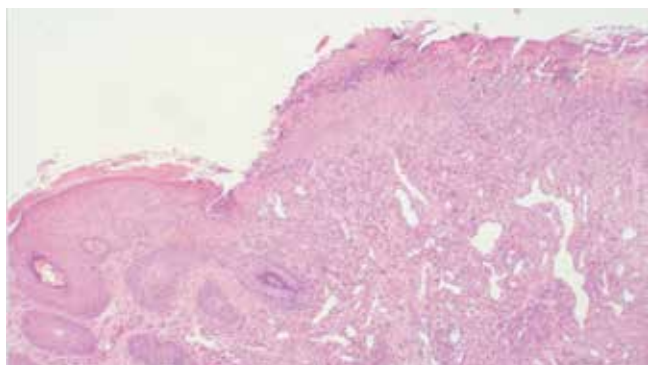


Fig 2 - Unspecific ulceration with dense inflammatory infiltrate on histopathological examination (H&E, 100x).

unable to be in public. After exclusion of others dermatological diseases, the suspicion of dermatitis *artefacta* was considered. When confronted in a non-accusational manner, the patient admitted scratching the nose but insisted that she did it because of the itch caused by the lesion. She mentioned that the whole problem began with the trauma of the nasal canula that causes intensive itch. She was treated with hydroxysine (25mg 2-3 times per day) and amitriptyline (25mg once a day). A local dressing was tried in order to disrupt the cycle of digital trauma with no significant results.

After several consultations, the approach that a psychiatrist would help her to diminish the itch was done and then posteriorly, we referred her to a psychiatrist who confirmed the diagnosis of dermatitis *artefacta* and make the diagnosis of a Major Depression with obsessive compulsive symptoms. The psychiatrist added agomelatine (25mg once a day) and suggested the nasal reconstruction for improving the patient's depression. During the follow-up, the patient admitted that she took previously antidepressants that she decided to stop. After several months of treatment with amitriptyline and agomelatine, she still scratched her nose and we obtained no significant results in stopping the progression of the lesion but improvement in the patient's mood were noticed. The patient was demanding for reconstructive surgery of her nasal defect. She went to a consultation of dermatological surgery and the reconstruction was planned.

DISCUSSION

Comorbidity of depression and dermatologic disorders is around 30%⁷. The skin is the most common site for self-damaging behaviors because it is easily

accessible and often the lesions are presented in an ostentatious manner⁷. The most common manifestations of self-destructive behavior are manipulations on the arms and legs, especially incisions on the forearms and are rarer in the nasal area⁸. The compulsive nose picking (rhinotillexomania) is a common benign habit in children and adults that may rarely become a serious affliction advancing to significant self-injury⁹.

Characteristically, the patient with dermatitis *artefacta* appears remarkably unconcerned in face of lesions that are morphologically bizarre, often geometric in outline, destructive, and reportedly of sudden, mysterious yet fully formed appearance, but are otherwise cooperative, allowing multiple diagnostic procedures or therapeutic measures^{1,5,7}. It is extremely difficult to comprehend why individuals intentionally inflict damage on themselves. Indirect diagnostic confirmation may be obtained by lesion healing after wound isolation with occlusive dressings.

Physicians may be hesitant to diagnose dermatitis *artefacta* because they fear that they might be missing organic disease or perhaps because they are unwilling to believe that the patients could be deceiving them⁵.

The differential diagnosis of a facial ulceration is broad and includes, besides dermatitis *artefacta*, trigeminal trophic syndrome that was not suggested in our case due to the absence of paresthesias and/or dysesthesias after trigeminal damage, and others etiologies as neoplastic (basal or squamous cell carcinoma), infective (herpetic reactivation, syphilis, leishmaniasis, mycobacterial infection, leprous trigeminal neuritis, paracoccidioidomycosis, blastomycosis), and inflammatory (Wegener's granulomatosis, pyoderma gangrenosum and sarcoidosis) etiologies were excluded by the biopsy.

Management must be delicate and a strong rapport with the patient is essential¹. Psychiatric referral should be carefully considered, as the patient may interpret it as rejection and intensify the self-induced cutaneous lesions. Patients should be referred for psychiatric evaluation only after an adequate patient-physician relationship has been established. Initially, direct confrontation with the patient and the diagnosis is discouraged^{3,5} as the patient will be in denial and may be lost of follow-up^{1,5}. When confronted it has to be done in a non-accusational manner. It is common for these patients to refuse psychiatric referrals¹. Although long-term studies are rare, the prognosis is considered poor^{1,6}.

This case illustrates a rare and exuberant cause of ulcer of the nose^{3,4}. The establishment of the diagnosis is often a time-consuming and complicated process that

Caso Clínico

precedes acceptance of the underlying pathology^{1,5}. Thus, recognizing and correctly diagnosing dermatitis artefacta is critical to avert unnecessary tests and treatments, allowing for more efficient management and better healing^{1,7,9}. This case shows typically the indifference of the patient. It also demonstrates that the strong rapport with the patient was essential in order to adhere to treatment and to begin evaluation with a psychiatrist.

REFERENCES

1. Gattu S, Rashid RM, Khachemoune A. Self-induced skin lesions: a review of dermatitis artefacta. *Cutis*. 2009; 84(5):247-51.
2. Potenza C, Bernardini N, Mambrin A, Skroza N. Dermatitis artefacta in a patient affected by impulse control disorder: case report. *Acta Dermatovenerol Croat*. 2011; 19(1):28-30.
3. George AE, Sarojini PA. Dermatitis artefacta - A case of 'munchausen's syndrome by proxy'? *Indian J Dermatol Venereol Leprol* 1994; 60:349-50.
4. Nielsen K, Jeppesen M, Simmelsgaard L, Rasmussen M, Thestrup-Pedersen K. Self-inflicted skin diseases. A retrospective analysis of 57 patients with dermatitis artefacta seen in a dermatology department. *Acta Derm Venereol*. 2005; 85(6):512-5.
5. Sun DK, Siegel DM. A nonhealing ulcer on the face. *Arch Fam Med*. 2000;9(9):787-9.
6. Verraes-Derancourt S, Derancourt C, Poot F, Heenen M, Bernard P. Dermatitis artefacta: retrospective study in 31 patients. *Ann Dermatol Venereol*. 2006; 133(3):235-8.
7. Baranska-Rybak W, Cubala WJ, Kozicka D, Sokołowska-Wojdyło M, Nowicki Roszkiewicz J. Dermatitis artefacta - a long way from the first clinical symptoms to diagnosis. *Psychiatr Danub*. 2011; 23(1):73-5.
8. Rudolph S, Schu U, Herrmann-Lingen C, Werner JA, Folz BJ. Nasal manifestations of self-destructive behaviour. *Rhinology*. 2007; 45(4):299-304.
9. Caruso RD, Sherry RG, Rosenbaum AE, Joy SE, Chang JK, Sanford DM. Self-induced ethmoidectomy from rhinotillexomania. *Am J Neuroradiol*. 1997; 18(10):1949-50.
10. Joe EK, Li VW, Magro CM, Arndt KA, Bowers KE. Diagnostic clues to dermatitis artefacta. *Cutis*. 1999; 63(4):209-14.