



Surgical & Cosmetic Dermatology

ISSN: 1984-5510

ISSN: 1984-8773

Sociedade Brasileira de Dermatologia

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Surgical & Cosmetic Dermatology, vol. 12, no. 1, Supl., 2020, October-December, pp. 43-45  
Sociedade Brasileira de Dermatologia

DOI: <https://doi.org/10.5935/scd1984-8773.20201241291>

Available in: <https://www.redalyc.org/articulo.oa?id=265568336009>

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# Nodular fasciitis in the forehead: a rare presentation

*Fasciite nodular na fronte: uma rara apresentação*

DOI: <https://www.dx.doi.org/10.5935/scd1984-8773.20201241291>

## ABSTRACT

Nodular fasciitis is a benign tumor resulting from the reactive proliferation of fibroblastic or myofibroblastic cells presenting rapid growth and rich cellularity. In adults, the extremities' involvement is more frequent; however, other regions can be affected. In this report, we present the case of a 40-year-old woman with a nodular lesion on the forehead. The histopathology confirmed the diagnosis of nodular fasciitis. The reported case highlights its rare location and alerts the clinical dermatologist in its inclusion among the differential diagnoses of tumor lesions on the face.

**Keywords:** Facial Dermatoses; Fasciitis; Forehead

## RESUMO

*A fasciite nodular é um tumor benigno, decorrente da proliferação reativa de células fibroblásticas ou miofibroblásticas de rápido crescimento e rica celularidade. Em adultos, o acometimento das extremidades é mais frequente; entretanto, outras regiões podem ser acometidas. Neste relato, é apresentada paciente feminina de 40 anos, com lesão nodular na fronte, com diagnóstico de fasciite nodular confirmado à histopatologia. O caso relatado procura destacar sua rara localização e alertar o dermatologista clínico para a sua inclusão entre os diagnósticos diferenciais das lesões tumorais na face.*

**Palavras-chave:** Fasciite; Face; Testa

## INTRODUCTION

Nodular fasciitis (FN) is a benign tumor derived from the proliferation of fibroblasts and myofibroblasts that can affect hypodermis, muscle, and fascia. In 1955, Konwaler described the FN as pseudosarcomatous fasciitis or infiltrative fasciitis, because of its strong resemblance to sarcoma.<sup>1</sup> The etiology of the FN is still uncertain, but there are reports of previous trauma in about 10 to 15% of cases.<sup>2</sup> Clinically, the lesion presents as a solitary, fast-growing nodule measuring 1 to 5 cm and more frequently affecting the extremities of the upper limbs. Some presentations can be confused with sarcoma due to its rapid growth, rich cellularity, and high mitotic activity, being of fundamental importance the distinction of the FN with malignant neoplastic diseases.

## Case report

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Received on: 30/11/2018

Approved on: 18/08/2020

Financial support: None.

Conflict of interest: None.

Study conducted at the Lauro de Souza Lima Institute, Bauru (SP), Brazil.

### Acknowledgment:

We would like to thank Dr. Cleverson Teixeira Soares, pathologist at the Lauro de Souza Lima Institute, for providing the PAAF photographic documentation and the anatomopathological examination.



## CASE REPORT

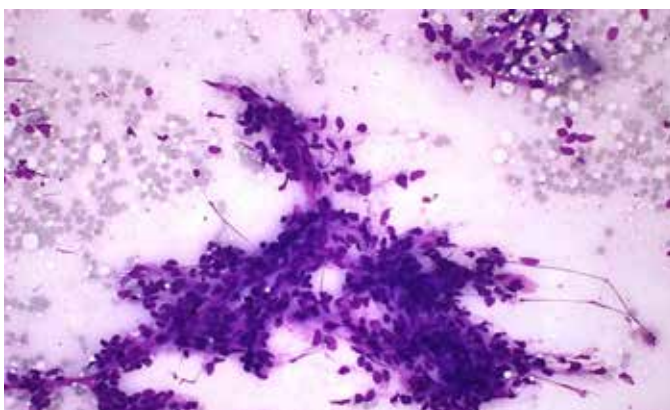
A 40-year-old hypertensive female patient with a history of onset of tumor in the forehead for 4 years, with progressive increase and mild local pain. On physical examination, the tumor of the patient had a 2 cm diameter, mobile, fibroelastic consistency, well-delimited, and not adhered to deep planes on the left forehead (Figure 1). With the diagnostic hypotheses of schwannoma, lipoma, and epidermoid cyst, and due to unavailability of ultrasonography exam (USG), fine needle aspiration (FNA) was performed, which showed fusocellular mesenchymal proliferative lesion suggestive of FN (Figure 2). We chose to perform surgical excision, with the removal of a 2cm-diameter tumor, with a light yellow color, well-delimited and parenchymal consistency (Figure 3). In the histopathological examination, the presence of mesenchymal proliferation with discrete atypias was noted (Figure 4), which, associated with FNAB and clinical presentation, confirmed the diagnosis of FN in the absence of immunohistochemistry.

## DISCUSSION

NF is a benign fibroproliferative disease of unknown etiology that affects men and women between the ages of 20 and 40. Reports of spontaneous remission and frequent location on bony prominences suggest the etiological hypothesis of previous trauma at the site of lesion.<sup>2</sup> In adults, the extremity involvement



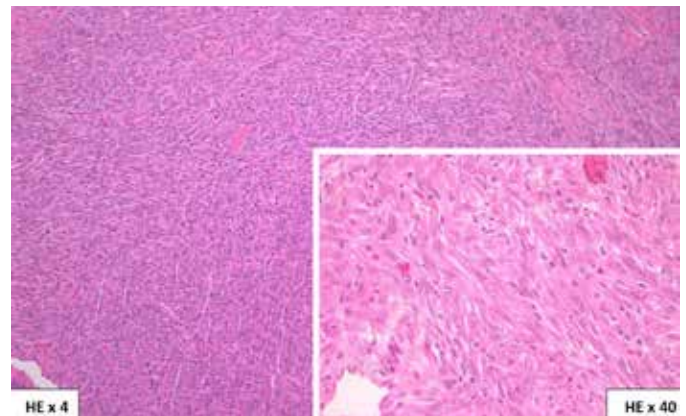
**FIGURE 1:** Movable nodule, about 2 cm in the left forehead, without involvement of the suprajacent skin



**FIGURE 2:** PAAF: presence of agglomerated fusel cell mesenchymal cells



**FIGURE 3:** Complete excision of the lesion, with removal of a pale yellow nodule, about 1.5 cm and parenchymal appearance on palpation



**Figure 4:** Histopathology: presence of mesenchymal proliferation with discrete atypia

is more frequent in the upper extremities (43%), followed by the trunk (25%) and lower extremities (22%), while only 10% of the FN occurs in the face and neck.<sup>3,4</sup> Most cases of NF in the face and neck occur in children.<sup>5</sup>

Clinically, FN manifests as a tumor lesion, well delimited, about 2 to 5 cm in size, with subcutaneous nodular growth fast and self-limited, and may present painful sensitivity at the site. The main differential diagnoses include pyogenic granuloma, cysts, lipoma, dermatofibroma, neurofibroma, and sarcoma.<sup>6,7</sup> Because it is infrequent, it is usually a neglected entity in the evaluation of benign tumor lesions, with other diagnostic hypotheses raised prior to FN. Many cases are usually confirmed by histopathological examination.<sup>8</sup>

USG can be performed and may show well-defined, hypoechoic, dermal nodular lesions, with or without the presence of a heterogeneous hyperechoic center, and may therefore make a differential diagnosis with malignant nodular lesions. Consequently, anatomopathological examination is necessary in these cases.<sup>9</sup>

Histopathology demonstrates a well-circumscribed, fascial or intramuscular subcutaneous nodule with a star-like appearance. A proliferation of rounded and fusiform fibroblast

and oval nucleus myofibroblasts with thin chromatin and prominent nucleolus is seen. In newly-onset lesions, the cells are loosely arranged in an edematous and myxomatous stroma, while the older lesions show hyalinized collagen bundles.<sup>10</sup> Fibroblasts and myofibroblasts react positively to vimentin and specific muscle actin and some cells are CD68 positive.<sup>11</sup> The histological differential diagnosis includes fibrosarcoma and malignant fibrous histiocytoma.

The treatment consists of complete surgical excision of the lesion, with a variable recurrence rate according to the literature, probably due to the incomplete excision of the lesion.

Other types of treatment may be considered conforming to the location of the lesion, such as CO2 ablative laser and intralesional infiltration with triamcinolone.<sup>12</sup>

The main objective of this study was to highlight the unusual location of the case and alert the dermatologist for the diagnosis and inclusion of NF as a differential diagnosis of other facial tumors, in order to define the best therapeutic approach of the lesion, avoiding diagnostic errors and possible local recurrences. ●

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