CASE REPORT

Cellular angiofibroma: A rare labia minor tumor

Patrícia Gomes Ferreira, Carolina Carneiro, Susana Saraiva, Vânia Ferreira, Horácio Scigliano, Isabel Monteiro

ABSTRACT

Cellular angiofibroma of the vulva is a rare benign mesenchymal tumor in middle-aged women, first reported in 1997 by Nucci et al. It is important to differentiate cellular angiofibroma from other tumors as these may be more aggressive and recurring. Cellular angiofibroma has a limited potential for local recurrence and is usually treated with complete local excision. A 45-year-old woman was referred to the gynecology appointment with a complaint of a discomfort mass in the right labium minus 1 year before, which has been progressively increasing in size for the past three months up to 6 cm. No change in vulvar skin color, local itching, or bilateral inguinal adenopathy. The tumor was excised, and the histopathological exam revealed a cellular angiofibroma. The patient recovered well and a good aesthetic result was achieved. This is the first case described of the cellular angiofibroma which arises from one of the labia minora. It is bigger in size and growing more rapidly than usual within three months. A simple excision was carried out and until now (12 months after) no recurrence signals.

Keywords: Cellular angiofibroma, Histopathology, Immunohistochemistry, Mesenchymal tumors, Vulva

INTRODUCTION

A fast-growing mass and local discomfort should be viewed as suspicious. The authors present the case of a woman with a large vulvar tumor arising from the right labium minus.

Cellular angiofibroma of the vulva is a rare benign mesenchymal tumor in middle-aged women, first reported in 1997 by Nucci et al. [1]. Mostly, it is presented in labia majora, being small-sized (<3 cm), painless, and well-circumscribed [1–4]. Histopathologically, it is characterized by a spindle cell component and abundant small-sized to medium-sized thick-walled vessels, and pathologically resembles angiomyofibroblastoma and aggressive angiomyxoma [1, 5, 6]. It is important to differentiate cellular angiofibroma from other tumors as these may be more aggressive and recurring. Cellular angiofibroma has a limited potential for local recurrence and is usually treated with complete local excision [1–4].

This report describes an atypical size and location of cellular angiofibroma of the vulva, which was successfully treated by excision.

CASE REPORT

A 45-year-old woman was referred to the gynecology appointment with a complaint of a tenderness mass in the right labium minus 1 year before, which has been progressively increasing for the past three months.
The patient’s menarche was at 16 years old, and she had 5 pregnancies and eutocic deliveries. She was submitted to bilateral tubal ligation at 40 years of age. She denied any personal or family history of breast or gynecological cancer. The patient had no history of trauma, chronic infection, or insect bite.

On examination, a peduncled tumor with 6 cm diameter was found, arising from the right labium minus (Figure 1), with no redness, and without signs of infection or abscess. At palpation, it was smooth and compressible. Vaginal examination was normal. No bilateral inguinal adenopathy. The patient was thoroughly investigated, and all her laboratory findings were within normal limits.

The tumor was excised with the peduncle and sent for histopathology (Figure 2). The surgical specimen was a soft grayish-brown mass measuring 6×4 cm. It was flat, well-delineated, with a disorganized fascicular pattern on section. Microscopic examination disclosed a well-circumscribed non-capsulated dermo-hypodermic benign neoplastic vascular tumor, predominantly of medium-large size. It presents a thick-walled hyalinized, arterial and venous vascular structures, with fibrous, rarely myxoid stroma, the latter showing small, monotonous, spindle cells, with bland monochromatic nuclei. Severe atypia, necrose, hemorrhage or mitosis figures were not identified (Figure 3).

Immunohistochemical study revealed positive stain for vimentin, CD34, estrogen, and progesterone receptors and absence of expression for epithelial membrane antigen (EMA) and smooth muscle actin (SMA). Overexpression of p16 nor p53 was not observed. Based on the above features, cellular angiofibroma was diagnosed (Figure 4).

Surgical excision with a free margin is the appropriate treatment for cellular angiofibroma. The postoperative period was uneventful (Figure 5). There was no recurrence in the follow-up until 12 months (Figure 6).

As there is just one case reported of recurrence and metastasis, there are no guidelines concerning the follow-up of patients with cellular angiofibroma.
DISCUSSION

Tumors primarily arising from the vulvovaginal area are relatively rare and they include soft tissue specific and non-specific tumors, as well as a spectrum of fibroepithelial tumors [7, 8]. Cellular angiofibroma is an uncommon benign mesenchymal neoplasm. It usually occurs on the vulva, particularly on the labia majora [9]. There are a few extra pelvic cases also reported [6].

Mandato et al. reviewed cases from 1997 to 2014 and found a total of 79 cases of female cellular angiofibroma [10]. Tumor sizes were between 0.6 and 12.3 cm, with a mean size of 3.6 cm. After this review, two vulvar cases with 20 cm were published [11, 12].

Histologically, cellular angiofibroma has two principal components: fusiform cells and prominent blood vessels. Fusiform cells form small fascicles in the middle of collagen bundles, frequent blood vessels of small to medium calibre, sometimes with a hyalinized wall, and a variable component of mature adipocytes. The tumor may present few mitotic activities and atypical cells in the stroma, but necrosis remains absent, although few instances of sarcomatous transformation and nuclear pleomorphism have also been reported [2, 4, 13–15]. The sarcomatous component can show variable features, such as: atypical lipomatous tumor, pleomorphic liposarcoma, and pleomorphic sarcoma. This phenomenon seems not to predispose to recurrence based on limited clinical follow-up available [14, 16].

All 79 cases summarized by Mandato et al. were treated with simple resection, and the surgical boundaries were not defined in 47 cases; the surgical boundary was positive in 18 cases and negative in 29 cases [10]. In the follow-up, five cases with positive surgical boundaries required reexcision [10]. In the present case, simple surgical resection was performed, and the surgical boundaries were reported to be histopathologically negative.

Of the 79 cases summarized by Mandato et al., 48 were followed up after 3–240 months; only a local recurrence of vulvar cellular angiofibroma is described after 6-month follow-up [10, 17]. Cellular angiofibroma has been initially excised with a rim of free-tumor tissue [10]. In another study of 12 patients, no recurrence or metastasis was reported at 14-month follow-up [14]. There are no reports of tumor metastasis, although the follow-up period in most of the studies is short [10, 14, 18].

CONCLUSION

This is the first case described of the cellular angiofibroma which arises from one of the labia minora. It is bigger sized (6 cm) than usual (<3 cm) and with rapid growth, within three months. A simple excision was carried out and until now (12 months after) no recurrence signals. In literature, a single case report of recurrence and no cases of metastasis, so there are no guidelines concerning the follow-up of patients with cellular angiofibroma.

REFERENCES


Author Contributions
Patricia Gomes Ferreira – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved
Carolina Carneiro – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved
Susana Saraiva – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved
Vânia Ferreira – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved
Horácio Scigliano – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved
Isabel Monteiro – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

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All relevant data are within the paper and its Supporting Information files.

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