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Neurofibroma Arising in Ectopic Breast Tissue: A Unique Case Report

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Authors' contributions

This work was carried out in collaboration between both authors. Author RC designed the study, wrote the first draft of the manuscript and managed the literature. Author AC supervised the entire work. Both authors read and approved the final manuscript.

Article Information

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Case Study

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ABSTRACT

Neurofibromas are benign nerve sheath tumours which occur in the dermis or subcutis with a variable distribution on the body surface. Accessory breast tissue results from incomplete involution of the embryologic mammary ridge. We report a case of a 63-year postmenopausal multiparous woman who presented to our department with a lump in the lower left abdominal quadrant near a supernumerary nipple. An abdominal ultrasound confirmed the presence of a 20mm mass and describes it as a hypo-anechoic neoformation below the accessory nipple in the context of likely breast parenchyma. A large local excision of the lesion was therefore planned. A final histologic diagnosis of neurofibroma arising in accessory breast tissue was made. This case report describes for the first time a case of a neurofibroma arising on ectopic breast tissue.

Keywords: Neurofibroma; breast; ectopic breast tissue.

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1. INTRODUCTION

Accessory breast tissue results from incomplete involution of the embryologic mammary ridge, also known as the milk line, wich extends from the axilla (most common location) to the groin. As a results, ectopic breast tissue can develop anywhere along this ridge. Neurofibromas are benign nerve sheath tumours and represent 5% of all benign soft tissue neoplasms. Nearly 10% of cases are found in associated with neurofibromatosis type 1 (NF-1), therefore, most of them occur in a solitary form. The NF-1 and solitary neurofibromas (SN) are regarded as two different disease processes, although present the same pathologic features, histologically. Most neurofibroma occur as solitary lesion; [1] solitary neurofibroma of the breast indipendent of neurofibromatosis is extremely rare and only a few case have been reported [2-6]. There is no evidence in the literature of neurofibroma cases arising in accessory breast tissue.

2. PRESENTATION OF CASE

A 63 year old postmenopausal multiparous woman presented to our department with a lump in the lower left abdominal quadrant near a supernumerary nipple that the patient reports to be present from the first pregnancy. On examination a 2 cm irregular mass, of likely belonging to the abdominal wall, was palpable. The abdominal lump was slowly increasing in size with no other associated symptoms. She had no medical comorbidities or family history of breast cancer. Systemic examination was unremarkable. Abdominal ultrasound was done. which confirmed the presence of a 20mm mass describes it as hypo-anechoic and а neoformation below the accessory nipple in the context of likely breast parenchyma. In relation to the dubious ultrasound characteristics a large local excision of the lesion was planned. Macroscopically, the sample consisted of a 3.8 x 1.8 cm skin lozenge containing a supernumerary nipple with underlying mammary tissue containing an irregular oval form of 1.6 cm of elastic consistency, a gray-white complexion and a cut collated appearance. Microscopically, benign neoplastic proliferation is appreciated, well delimited by the surrounding tissue, formed by cells with an elongated nucleus immersed in a fibrous stroma. No atypical mitoses were observed. The neoplastic cell population was positive for the reactions prepared with S100 anti-protein, anti-CD34, anti-BCL2 and antivimentin antibodies while it was negative for anti-CD117 antibody, smooth muscle anti-actin, antidesmin, anti CD45RB and anti-pancitocheratin. The determination of nuclear receptors for estrogen and progesterone hormones carried out using monoclonal antibodies gave a positive result (Er 75%, PgR 50%). The cell proliferation index evaluated with anti-Ki67 antibody gave positivity in 5% of neoplastic cells. A final histologic diagnosis of neurofibroma was made.

The patient did not relapse at 1 year of follow-up and the surgical scar is well re-epithelialized.



Fig. 1. Ultrasonic image showing position of lump

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Fig. 2. Lump in the lower left abdominal quadrant near a supernumerary nipple



Fig. 3. Morphology of the lump

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Fig. 4. Morphology of the lump showing its characteristcs



Fig. 5. Ultrasonic image

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Fig. 6. Side view of the lump



Fig. 7. Lateral view of the lump



Fig. 8. Lump after operative phase



Fig. 9. Ultrasonic image after operation

3. DISCUSSION

Breast development begins during the fifth or sixth week of gestation; mammary crests form from the anterior axillary fold to the inside of the inguinal folds. Physiologically, mammary ridges disappear shortly after their formation, with the exception of a small portion that persists in the pectoral region, where only one pair of mammary glands typically develop [7]. So if this regression fails, accessory nipples (polythelia) or accessory mammary glands (polymastia) can develop along the mammary ridges [8]. Accessory breast tissue, also known as ectopic breast tissue, is caused by the non-regression of primary breast tissue, except at the pectoral breast site. According Kayaka's classification. to supernumerary breast is defined, as in the case reported here, the presence of mammary gland associated with its components (nipple, areola, ducts) [9]. Multiple locations of accessory breast are reported, where the most frequent sites are the axilla and the infra-mammary region, but a location in the lower left abdominal guadrant is not described [10]. In the context of the accessory parenchyma, various types of lesions have been described such as fibroadenomas, [11] phyllodes tumors [12] and malignant tumors [10], but no cases of neurofibroma have been described at present. Neurofibromas are benign nerve sheath tumours which were first described bv Smith in 1849 and later by von Recklinghausen in 1882 [3]. These are relatively frequent tumours that can occur in both sexes and at any age. Most neurofibromas are solitary lesions (SN) which occur in the dermis or subcutis with a variable distribution on the body surface. [1] With regard to SN of the breast, Anglo-Saxon literature considers them as rare neoplasms. SN of the breast typically arise in the skin or near the nipple-areola complex [13]. In the context of the breast parenchyma, descriptions of such lesions are even rarer than typical sites. There is currently no work describing cases of neurofibroma originating in ectopic breast tissue.

4. CONCLUSION

To our knowledge this is the first description of case report of а neurofibroma а tissue. arising on ectopic breast The uniqueness of this finding underlines the importance of this description and lays the foundation for a thorough review of this tumor pathology.

CONSENT AND ETHICAL APPROVAL

As per university standard guideline participant consent and ethical approval has been collected and preserved by the authors.

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COMPETING INTERESTS

Authors have declared that no competing interests exist.

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