



# Surgical management of a giant malignant phyllodes tumour of the breast: a case report

Jenny Guevara-Martínez<sup>1^</sup>, Irene Osorio<sup>2</sup>, Juan Bernar<sup>2</sup>, Sergio Salido<sup>2</sup>, Cecilia Meliga<sup>2</sup>, Natascha Elsner<sup>3</sup>, Ricardo Pardo<sup>2</sup>

<sup>1</sup>General and Digestive Surgery, University Hospital La Paz (Hospital Universitario La Paz), Madrid, Spain; <sup>2</sup>General and Digestive Surgery, Breast and Endocrine Surgery Department, Fundación Jiménez Díaz, Madrid, Spain; <sup>3</sup>Department of Obstetrics and Gynecology, Hospital Universitario La Candelaria, Tenerife, Spain

*Correspondence to:* Jenny Guevara-Martínez. General and Digestive Surgery, University Hospital La Paz (Hospital Universitario La Paz), Paseo la Castellana 261, 28046, Madrid, Spain. Email: jennyguevaram@hotmail.com.

**Abstract:** Phyllodes tumour (PT) is a rare fibroepithelial neoplasm, being divided into benign, borderline or malignant, and usually presents as large masses with rapid growth. Breast tumours above 4 cm with these characteristics are highly suspicious of PT, and those above 10 cm are considered giant PTs, representing only 20% of these tumours. Prognosis relies on histological type and a mostly on a complete surgical resection with margins above 1 cm. Surgical management of giant PTs can be a technical challenge, and may require advanced breast reconstructive techniques. We present the case of a giant PT, completely resected with a mastectomy and nodal dissection. The patient was discharged without any complications and is currently on follow-up without recurrence. Adequate resection margins should always be the principal aim when providing an optimal surgical treatment of a PT. A multidisciplinary team evaluation by an experienced breast surgeon or a plastic reconstructive surgeon is recommended when planning a proper resection with further reconstruction. Axillary node metastases are rare, and dissection is limited to patients with pathological preoperative findings. Hematogenous dissemination may be present in malignant PTs. Classical adjuvant therapies like chemotherapy, hormone therapy or radiotherapy are not widely prescribed when treating PT. We strongly emphasize in providing a correct initial resection of the tumour.

**Keywords:** Giant phyllodes; malignant phyllodes; mastectomy; case report

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## Introduction

Phyllodes tumours (PT) of the breast are an uncommon type of fibroepithelial neoplasms, representing 0.5% to 1% of all fibroepithelial neoplasms (1-3). They are classified benign, borderline and malignant tumours, as accorded by the World Health Organization (1-4). They usually present as a large breast tumor with rapid growth above 4 cm, considering those over 10 cm as giant PT (2). Due to its rapid and considerable growth, their diagnosis is challenging with the usual techniques, and core biopsy is

necessary in most cases. Their management relies mostly on an adequate surgical resection, taking especial emphasis into achieving resection margins above 1 cm in order to avoid local recurrence. In cases of local recurrence, a new local excision or mastectomy is mandatory. Because of their rapid and accentuated growth, their surgical resection can be technically challenging, considering in large tumours the confection of flaps in order to maintain safe resection margins. Malignant lesions frequently show hematogenous dissemination, regardless of the presence or not of local

<sup>^</sup> ORCID: 0000-0001-9158-1610.

recurrence (1). Other treatments widely used in breast tumours, such as chemotherapy, radiotherapy or hormone therapy are not routinely used, making it fundamental to provide an adequate excision, with proper margins. We present the following case in accordance with the CARE reporting checklist (available at <https://abs.amegroups.com/article/view/10.21037/abs-20-150/rc>).

### Case presentation

We present the case of a female patient of 62 years old, without any personal or familiar history of breast cancer or any other malignancies, with a rapid growth of a breast mass, over the last month before consultation. She didn't have any trauma, pain, or infectious symptoms suggestive of other diagnosis. During the physical exam, the breast showed an ulcerated mass of 20 cm in diameter. Local curations of the ulcer were being performed by the patient. A core biopsy was performed under a probable diagnosis of a breast sarcoma. She underwent magnetic resonance and computed tomography (*Figure 1*), without evidence of any distant metastases, but with positive axillary nodes. After a multidisciplinary discussion in the breast tumour committee, we performed a mastectomy (*Figures 2-6*). Despite its large size, there was no need to make any advancements flaps, and the pectoral muscle fascia and axillary nodes were dissected to avoid local recurrence. Closed aspirate drains were left in the mastectomy and axillary dissection territories. The patient was discharged without any complications during the early postoperative period. The postoperative diagnosis was of a PT with sarcomatous degeneration, with negative axillary nodes. Our patient is currently on close follow up without local recurrence or distant metastases.

All procedures performed in this study were in accordance with the ethical standards of the institutional and/or national research committee(s) and with the Helsinki Declaration (as revised in 2013). Written informed consent was obtained from the patient.

### Discussion

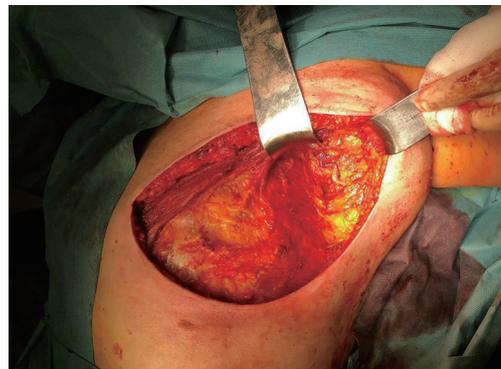
PT of the breast is an uncommon type of fibroepithelial neoplasms, with less than a 1% of incidence. They typically present an exaggerated intracanalicular pattern with leaf-like fronds protruding into dilated spaces, accompanied by hypercellularity (2). The definition of "phyllodes" originates from the Latin root "Phyllodium" meaning leaf, due to



**Figure 1** Preoperative CT-scan.



**Figure 2** Preoperative tumour.



**Figure 3** Mastectomy and axillary dissection.

its appearance on microscopy (2,3). They have a variety of behaviors, from benign fibroadenoma to malignant neoplasms with the ability to produce distant metastases.

Usually, the patients show a rapid onset of symptoms with a large, painless mass, as the one described in our case. Due to their large sizes, core needle biopsy or fine needle



**Figure 4** Axillary dissection with thoracodorsal pedicle.



**Figure 5** Immediate postoperative results.



**Figure 6** Mastectomy specimen.

aspiration may be insufficient for diagnosis. There may be strong similar histological findings, making it difficult to differentiate a benign PT from cellular fibroadenoma, requiring an evaluation of its borders after excisional biopsy (1,2). Both lesions have an increased stromal cellularity,

being differentiated by well-circumscribed borders in fibroadenomas and irregular borders in benign PT. On the other hand, malignant PTs must be distinguished from sarcomas and other types of carcinomas. Malignant PTs represent 1/4 of all PTs, having a strong stromal cellularity with cellular atypia, infiltrative borders, mitotic activity of at least 10/10 HPF and a stromal overgrowth with heterologous elements (2-5).

Mastectomy is not always necessary in small lesions, being only recommended when a margin of 1 cm cannot be achieved with a local excision. Positive surgical margins or less than 0.5 cm, recurrent tumours or a diameter over 10 cm—or 5 cm in malignant (6)—have been considered high risk for local recurrence (2,6,7). In patients with benign or borderline tumours with positive margins, is recommended a re-excision or close follow-up with physical examination and ultrasound every 6 months, and annual mammogram (7,8). In malignant lesions with positive margins, a re-excisional surgery would be necessary. Giant PTs represents only 20% of the total of PT and when planning their excision, different reconstructive techniques should be considered, including deep inferior epigastric artery perforator (DIEP), latissimus dorsi or rectus abdominis flaps (9,10). Their large sizes, and the need to achieve a complete excision with safe margins, may require a multidisciplinary approach in terms of reconstruction.

Although an axillary dissection is not routinely performed, 20% of patients with malignant PT have positive nodes (2). Positive nodes during preoperative radiology or physical exam, should orientate the need to complete an axillary dissection. In our case, an axillary dissection was done due to positive preoperative CT, although postoperative specimen of nodes was negative for metastases.

There is no final recommendation regarding radiotherapy. Although it might reduce the rate of local recurrence, a consensus has not been determined due to inconclusive results, leaving the use of postoperative radiotherapy only for determined patients, especially those considered as high risk of recurrence. In those tumours above 5 cm, postoperative radiotherapy should be considered because of the high risk of recurrence (6). After the resection of a recurrent lesion, adjuvant radiotherapy should be considered, following the same principles as soft tissue sarcoma (8).

Adjuvant or neoadjuvant chemotherapy is not routinely used, as there is no evidence of benefit, limiting its use only for compassionate indications. Despite the fact that hormone receptors might be positive in up to 75%, hormone therapy is not a recommended treatment in these neoplasms. In

cases of metastatic disease, the same recommendations as for soft tissue sarcoma should be considered (8).

A close follow-up is fundamental in these lesions due to its potential of local recurrence. Current NCCN guidelines recommend clinical follow-up for 3 years (8). If there is recurrence without metastasis, re-excision with tumorectomy or mastectomy can be performed without affecting the overall survival. Metastatic disease, mainly in the lung, abdomen and bone decreases the overall survival.

Despite the large size of the tumour we presented, a complete resection with adequate surgical margins was achieved with a mastectomy and without the need of further reconstructive techniques. When treating a PT, the aim to provide negative margins must be warranted, and multidisciplinary surgical strategies planned when needed. Since other adjuvant therapies are not demonstrated to avoid recurrence, all efforts must be towards an adequate resection.

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## Footnote

*Reporting Checklist:* The authors have completed the CARE reporting checklist. Available at <https://abs.amegroups.com/article/view/10.21037/abs-20-150/rc>

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*Ethical Statement:* The authors are 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. All procedures performed in this study were in accordance with the ethical standards of the institutional and/or national research committee(s) and with the Helsinki Declaration (as revised in 2013). Written informed consent was obtained from the patient for publication of this case report and accompanying images. A copy of the written consent is available for review by the editorial office of this journal.

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