Male Breast Huge Malignant Phyllodes

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Rezumat

Tumoră Phyllodes gigantă, malignă, la un bărbat


Discussii şi concluzii: Au fost puţine cazuri de tumori Phyllodes raportate la bărbaţi. Patologic, tumorile Phyllodes pot fi de trei tipuri: benigne, borderline şi maligne în funcţie de activitatea mitotică, natura marginilor de resecţie, creşterea stromală, celularitate şi atipie. Tumorile Phyllodes maligne tind să se extindă hematologic către plămâni, apoi către oase. Tumorile Phyllodes tind să recidiveze chiar după excizie completă, în special cele maligne, dar şi cele beneigne. Tratamentul de elecție pentru tumorile Phyllodes este excizia locală largă · cu sau fără radio·
terapie adjuvantă în leziunile maligne - fără rezultate dovedite ale chimioterapiei sau hormonoterapiei.

**Cuvinte cheie:** san, tumoră Phyllodes, leziune malignă

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**Abstract**

*Introduction:* Phyllodes tumors are rare breast tumors. The best treatment is wide local excision with 1 cm safety margin unless metastatic. The three pathological types (benign, borderline and malignant were reported in men.

*Case presentation:* A 73-year-old male with huge left breast swelling extending from the clavicle to the left hypochondriac region. Core needle biopsy suggested malignant phyllodes tumor. Post-contrast CT revealed a huge mass seen at the left anterolateral chest wall measuring about (22 x 25 x 26 cm). Simple mastectomy was performed en bloc with the tumor. The microscopic examination led to the diagnosis of high grade malignant phyllodes. IHC showed diffuse positive vimentin, CD10 and negative CK in the neoplastic cells. The patient lost follow up for three months. Then he was presented with fungating local recurrence with bilateral metastatic pulmonary. The patient underwent palliative excision. After the second surgery, he was prepared for palliative chemoradiotherapy but the patient died one month later at home.

*Discussion & conclusions:* Very few cases of phyllodes tumor were reported in men. Pathologically, phyllodes tumors are subdivided into three types: benign, borderline and malignant according to mitotic frequency, nature of margins, stromal growth, cellularity and atypia. Malignant phyllodes tumors tend to spread via hematological route mainly to the lung, then to the bone. Phyllodes tumors –even benign type- tend to recur even after complete excision with higher tendency for malignant cases. Wide local excision is the standard of care for phyllodes tumors – with or without adjuvant radiotherapy in malignant lesions- with no proved value for chemotherapy or hormonal therapy.

**Key words:** breast, phyllodes, malignant

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**Introduction**

Phyllodes tumors are rare breast tumors, encountering for about 1% of all breast neoplasms with very rare incidence in men and few cases reported in literature, commonly preceded by gynecomastia. Best treatment is wide local excision with 1 cm safety margin unless metastatic. The three pathological types (benign, borderline and malignant) were reported in men (1).

**Case report**

Seventy three year-old male, farmer, married with six offspring, with neither relevant medical nor surgical history. He was presented to our center in August 2015 complaining from huge left breast swelling. Physical examination revealed accepted general condition with huge left breast swelling extending from the clavicle to the left hypochondriac region with normal right breast & no suspicious axillary lymph nodes. Core needle biopsy revealed breast tissue showing infiltration by malignant spindle cell proliferation, necrotic areas were also seen. The primary diagnosis was malignant spindle cell proliferation most probably malignant phyllodes tumor. Post contrast CT (Fig. 1) revealed a huge heterogeneous mass
seen at the left anterolateral chest wall showing central necrosis with coarse calcifications inside measuring about (22 x 25 x 26 cm). The mass was seen abutting the axillary artery with no encasement. Simple mastectomy was performed en bloc with the tumor. The gross specimen was received as a huge soft tissue mass measuring about 40x30x15 cm covered by nipple, areola and skin ellipse measuring 10x7 cm. On dissection, it was firm in consistency and grayish white in colour with areas of extensive necrosis. Microscopic examination (Fig. 2) showed hypercellular malignant tumoral proliferation formed of

Figure 1. Post contrast CT axial & sagittal reformatted images (A & B): Huge heterogeneously enhancing left anterolateral chest wall soft tissue mass with extensive central necrosis & coarse calcification (arrow) is seen inseparable from left breast fibroglandular tissue as well as anterior chest wall muscles & extending down to the left 5th anterior rib with no bony destruction. The mass is seen abutting left axillary artery (arrowhead) with preserved fat plane in-between. No pathological axillary LNs could be detected.

Figure 2. Microscopic examination of the first excision. (A): Hypercellular malignant tumoral proliferation formed of fascicles of malignant spindle cells. H & E x200. (B): The spindle cells show high degree anaplasia and pleomorphism with frequent mitotic figures (> 10/10HPF). H & E x400. (C): Malignant spindle cells with wide area of necrosis. H & E x100.
fascicles and bundles of malignant spindle cells exhibiting high degree anaplasia, pleomorphism and frequent mitotic figures (> 10/10 HPF) and wide necrotic areas. Immunohistochemistry (IHC) was done (Fig. 3) and it showed diffuse positive vimentin, CD10 and negative CK in the neoplastic cells. Sections prepared from skin were free from tumor tissue. The final diagnosis was high grade malignant phyllodes tumor. The patient was discharged on the following day. He was referred to clinical oncology department for consultation about radiotherapy. Yet, the patient lost follow up for three months. Three months later, the patient was presented with fungating local recurrence. Chest CT (Fig. 4) revealed few heterogeneous- ly enhancing necrotic masses at the left anterolateral chest wall, the largest measured about 4.5 x 6.5 cm. Other enlarged masses were seen in the left axilla and the largest was 7.6 x 16 cm. Few similar masses were seen encasing the left axillary artery for more than 180° with no definite infiltration. The length of contacting segment measured about 3.7 cm. Few variable sized bilateral metastatic pulmonary nodules were seen: the largest was seen in left upper lung lobe measuring about 2.2 cm. The patient underwent palliative excision with microscopic examination and IHC showing recurrent high grade malignant phyllodes tumor. After the second surgery, he was prepared for palliative chemoradiotherapy with single agent Holoxane with 25% reduction and radiotherapy on palliative base 50 gy/25 sessions. The patient died one month later at home from a non-rivaled cause.

**Discussion**

Phyllodes tumors (previously named cystosarcoma phyllodes) are biphasic tumors

![Figure 3. Immunohistochemical study of the first excision. (A): Diffuse positive CD10 in the neoplastic cells (immunoperoxidase-DAB x100). (B): Diffuse positive vimentin in the neoplastic cells (immunoperoxidase-DAB x200). (C): Negative CK in the neoplastic cells (immunoperoxidase-DAB x100).]
composed of both epithelial and stromal elements. They are pathologically classified into: benign, borderline and malignant. This classification is based on variable features such as degree of cellular pleomorphism, mitotic count, stromal overgrowth and margin type. The incidence in women is less than 1% of all breast tumors and 2-3% of fibroepithelial tumors. Very few cases were reported in men due to infrequent lobular development. The reported cases are mostly linked with pre-existing gynecomastia. Phyllodes tumors were also reported in seminal vesicles & prostate (1–5).

The incidence link with gynecomastia led to a proposal that development of male phyllodes may be preceded with lobular development in male breast due to hormonal imbalance between estrogen & androgens (3,6) Our patient was not obese (BMI: 21) with no relevant medical history. He was moderate smoker for 35 years.

The previously reported cases had a mean age of 53 years old while the tumor diameter ranged from 1-45 centimeters (3,7). Our case can be considered one of the oldest cases as well as being one of the largest reported tumors.

On radiologic evaluation, phyllodes tumors appear in mammogram as well defined hyperdense oval polylobulated masses which may contain calcifications. At ultrasonography, they may present as a solid oval well circumscribed hypoechoic mass with parallel orientation, it may contain cystic areas. MRI may use cystic changes and wall irregularity as differentiating features between benign and malignant phyllodes. However, in general, radiological evaluation is not accurate for differentiating between benign phyllodes and fibroadenoma–however, the presence of a cystic component and strong lobulation favors the diagnosis of phyllodes tumor rather than fibroadenoma–as well as sharp delineation between the three subtypes of phyllodes tumors where imaging may be suggestive but not diagnostic of grading (8–12).

Pathologically, most phyllodes tumors show increased cellularity with increased stroma cells around ductal epithelium. FNAC is not sufficient to differentiate it from fibroadenoma. Core needle biopsy may be – but not always–
able to reach a diagnosis guided by the degree of cellularity as well as increased stromal mitotic activity and nuclear atypia. According to mitotic frequency, nature of margins, stromal growth, cellularity and atypia, phyllodes tumors are subdivided pathologically into three types: benign, borderline and malignant which composes about 20-50% of all phyllodes tumors. Malignant phyllodes tumors tend to spread via hematological route mainly to the lung then bone. Phyllodes tumors – even benign type – tend to recur even after complete excision with higher tendency for malignant cases. Recurrent tumors may present with a higher grade than the primary ones (2,4,13,14).

Wide local excision is the standard of care for benign, borderline and non-metastatic malignant phyllodes tumors – with or without adjuvant radiotherapy in malignant lesions with no proved value for chemotherapy or hormonal therapy. Axillary dissection is not recommended. Metastatic phyllodes tumor patients are candidate for palliative therapy with overall poor prognosis (3,4,8,12,15).

Authors’ Contributions

Conflict of interest: None declared.

References