

World Journal of *Clinical Cases*

World J Clin Cases 2020 August 26; 8(16): 3377-3620



OPINION REVIEW

- 3377 Novel computerized psychometric tests as primary screening tools for the diagnosis of minimal hepatic encephalopathy
Luo M, Mu R, Liu JF, Bai FH

REVIEW

- 3390 Management of cancer patients during COVID-19 pandemic at developing countries
González-Montero J, Valenzuela G, Ahumada M, Barajas O, Villanueva L

MINIREVIEWS

- 3405 Liver in the limelight in the corona (COVID-19) time
Chela HK, Pasha SB, Basar O, Daglilar E, Tahan V
- 3411 Steroid-responsive pancreatitis
Pelaez-Luna M, Soriano-Rios A, Lira-Treviño AC, Uscanga-Domínguez L

ORIGINAL ARTICLE**Clinical and Translational Research**

- 3431 Application of molybdenum target X-ray photography in imaging analysis of caudal intervertebral disc degeneration in rats
Su QH, Zhang Y, Shen B, Li YC, Tan J
- 3440 Accuracy study of a binocular-stereo-vision-based navigation robot for minimally invasive interventional procedures
Wang R, Han Y, Luo MZ, Wang NK, Sun WW, Wang SC, Zhang HD, Lu LJ

Retrospective Study

- 3450 Value of virtual bronchoscopic navigation and transbronchial ultrasound-guided sheath-guided exploration in diagnosis of peripheral lung cancer
Liu Y, Wang F, Zhang QC, Tong ZH
- 3458 Significance of serum fibroblast growth factor-23 and miR-208b in pathogenesis of atrial fibrillation and their relationship with prognosis
Chen JM, Zhong YT, Tu C, Lan J
- 3465 Home quarantine compliance is low in children with fever during COVID-19 epidemic
Lou Q, Su DQ, Wang SQ, Gao E, Li LQ, Zhuo ZQ

- 3474 Combination of endoscopic submucosal dissection and laparoscopic sentinel lymph node dissection in early mucinous gastric cancer: Role of lymph node metastasis

Li H, Zhao LL, Zhang XC, Liu DX, Wang GY, Huo ZB, Chen SB

- 3483 Factors affecting failed trial of labor and countermeasures: A retrospective analysis

Wang JG, Sun JL, Shen J

- 3493 Value of miR-1271 and glypican-3 in evaluating the prognosis of patients with hepatocellular carcinoma after transcatheter arterial chemoembolization

Guo Z, Wang J, Li L, Liu R, Fang J, Tie B

Observational Study

- 3503 Follow-up study on symptom distress in esophageal cancer patients undergoing repeated dilation

Liu L, Liu QW, Wu XD, Liu SY, Cao HJ, Hong YT, Qin HY

- 3515 Long-term medical treatment of patients with severe burns at exposed sites

Du Y, Lv GZ, Yu S, Wang D, Tan Q

CASE REPORT

- 3527 Laparoscopic management of a giant mucinous benign ovarian mass weighing 10150 grams: A case report

Sanna E, Madeddu C, Melis L, Nemolato S, Macciò A

- 3534 Concurrent hepatocellular carcinoma metastasis to stomach, colon, and brain: A case report

Kim R, Song J, Kim SB

- 3542 Disseminated osteomyelitis after urinary tract infection in immunocompetent adult: A case report

Kim YJ, Lee JH

- 3548 Pelvic lipomatosis and renal transplantation: A case report

Zhao J, Fu YX, Feng G, Mo CB

- 3553 Intestinal obstruction in pregnancy with reverse rotation of the midgut: A case report

Zhao XY, Wang X, Li CQ, Zhang Q, He AQ, Liu G

- 3560 Clinical laboratory investigation of a patient with an extremely high D-dimer level: A case report

Sun HX, Ge H, Xu ZQ, Sheng HM

- 3567 Recovery from a biliary stricture of a common bile duct ligature injury: A case report

Fan Z, Pan JY, Zhang YW

- 3573 Spontaneous pneumomediastinum in an elderly COVID-19 patient: A case report

Kong N, Gao C, Xu MS, Xie YL, Zhou CY

- 3578 Acute generalized exanthematous pustulosis with airway mucosa involvement: A case report

Li LL, Lu YQ, Li T

- 3583** Multifocal neuroendocrine cell hyperplasia accompanied by tumorlet formation and pulmonary sclerosing pneumocytoma: A case report
Han XY, Wang YY, Wei HQ, Yang GZ, Wang J, Jia YZ, Ao WQ
- 3591** Giant benign phyllodes breast tumour with pulmonary nodule mimicking malignancy: A case report
Zhang T, Feng L, Lian J, Ren WL
- 3601** Spontaneous multivessel coronary artery spasm diagnosed with intravascular ultrasound imaging: A case report
Wu HY, Cao YW, Chang FJ, Liang L
- 3608** Delayed perforation after endoscopic resection of a colonic laterally spreading tumor: A case report and literature review
Zhou GYJ, Hu JL, Wang S, Ge N, Liu X, Wang GX, Sun SY, Guo JT
- 3616** First branchial cleft cyst accompanied by external auditory canal atresia and middle ear malformation: A case report
Zhang CL, Li CL, Chen HQ, Sun Q, Liu ZH

ABOUT COVER

Editorial board member of *World Journal of Clinical Cases*, Dr. Kvolik is a Professor in the School of Medicine, Osijek University, Croatia. She obtained her MD degree, with specialization in the field of anesthesiology, resuscitation and intensive care from the Zagreb Medical School, Croatia. Afterwards, she undertook postgraduate training in Clinical Pharmacology at the same institution, defending both a Master's thesis and PhD thesis. In 2006, she joined the Osijek University Medical Faculty as a lecturer and was promoted to Professor in 2009. In 2012, she was elected Head of the Department of Anesthesiology, Resuscitation, Intensive Care and Pain Therapy, a position she occupies to this day. She is also the current Head of the Intensive Care Unit at the Osijek University Hospital, Croatia. (L-Editor: Filipodia)

AIMS AND SCOPE

The primary aim of *World Journal of Clinical Cases* (*WJCC*, *World J Clin Cases*) is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

WJCC mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

INDEXING/ABSTRACTING

The *WJCC* is now indexed in Science Citation Index Expanded (also known as SciSearch®), Journal Citation Reports/Science Edition, PubMed, and PubMed Central. The 2020 Edition of Journal Citation Reports® cites the 2019 impact factor (IF) for *WJCC* as 1.013; IF without journal self cites: 0.991; Ranking: 120 among 165 journals in medicine, general and internal; and Quartile category: Q3.

RESPONSIBLE EDITORS FOR THIS ISSUE

Production Editor: *Ji-Hong Liu*; Production Department Director: *Xiang Li*; Editorial Office Director: *Jin-Lai Wang*.

NAME OF JOURNAL

World Journal of Clinical Cases

ISSN

ISSN 2307-8960 (online)

LAUNCH DATE

April 16, 2013

FREQUENCY

Semimonthly

EDITORS-IN-CHIEF

Dennis A Bloomfield, Sandro Vento, Bao-Gan Peng

EDITORIAL BOARD MEMBERS

<https://www.wjgnet.com/2307-8960/editorialboard.htm>

PUBLICATION DATE

August 26, 2020

COPYRIGHT

© 2020 Baishideng Publishing Group Inc

INSTRUCTIONS TO AUTHORS

<https://www.wjgnet.com/bpg/gerinfo/204>

GUIDELINES FOR ETHICS DOCUMENTS

<https://www.wjgnet.com/bpg/GerInfo/287>

GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH

<https://www.wjgnet.com/bpg/gerinfo/240>

PUBLICATION ETHICS

<https://www.wjgnet.com/bpg/GerInfo/288>

PUBLICATION MISCONDUCT

<https://www.wjgnet.com/bpg/gerinfo/208>

ARTICLE PROCESSING CHARGE

<https://www.wjgnet.com/bpg/gerinfo/242>

STEPS FOR SUBMITTING MANUSCRIPTS

<https://www.wjgnet.com/bpg/GerInfo/239>

ONLINE SUBMISSION

<https://www.f6publishing.com>

Giant benign phyllodes breast tumour with pulmonary nodule mimicking malignancy: A case report

Ting Zhang, Liang Feng, Jie Lian, Wei-Li Ren

ORCID number: Ting Zhang 0000-0001-5026-8353; Liang Feng 0000-0002-8378-7157; Jie Lian 0000-0003-2822-7197; Wei-Li Ren 0000-0003-1858-4350.

Author contributions: Zhang T was the patient's oncologist, reviewed the literature and contributed to manuscript drafting; Feng L collected the patient's clinical data; Lian J performed the pathologic analyses and reviewed the histological sections of this case; Ren WL reviewed and revised the manuscript.

Informed consent statement:

Informed written consent was obtained from the patient for publication of this report and any accompanying images.

Conflict-of-interest statement: The authors declare that they have no conflict of interest.

CARE Checklist (2016) statement:

The authors have read the CARE Checklist (2016), and the manuscript was prepared and revised according to the CARE Checklist (2016).

Open-Access: This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative

Ting Zhang, Liang Feng, Wei-Li Ren, Department of Breast Disease Center, People's Hospital of Shangyu, Shaoxing 312300, Zhejiang Province, China

Jie Lian, Department of Pathology, People's Hospital of Shangyu, Shaoxing 312300, Zhejiang Province, China

Corresponding author: Wei-Li Ren, MBBS, Surgical Oncologist, Department of Breast Disease Center, People's Hospital of Shangyu, No. 517 Baiguan Street, Shangyu District, Shaoxing 312300, Zhejiang Province, China. yar201810@163.com

Abstract

BACKGROUND

Phyllodes tumours (PTs) are fibroepithelial breast tumours, which can be classified as benign, borderline or malignant, according to their histological characteristics. While various huge borderline or malignant PTs have been previously described, a benign PT with a pulmonary nodule mimicking malignancy has not yet been reported. In order that doctors may have a comprehensive understanding of super-giant benign PTs (≥ 20 cm), we also performed a literature review to summarize the clinical features, differential diagnosis, and treatment of this disease.

CASE SUMMARY

A 42-year-old woman with severe anaemia presented with a rapidly enlarging right breast mass, measuring approximately 30 cm \times 25 cm \times 20 cm that was first noticed 1 year previously. A region of skin ulceration and necrosis (20 cm \times 15 cm) was observed on the lateral side of the mass. Computed tomography (CT) of the chest revealed a pulmonary nodule, which initially suggested a diagnosis of metastasis. CT showed that the boundaries between the pectoralis major and the mass were blurred, which was presumed to be due to tumour invasion. However, two core needle biopsies of the mass showed no evidence of malignancy. Following these results, the tumour was removed by mastectomy of the right breast. Interestingly, postoperative pathology finally proved the diagnosis of a benign PT. After 1 year of follow-up, wedge resection of the small pulmonary nodule was performed, and it was confirmed that the lung nodule was actually adenocarcinoma rather than metastatic breast cancer. The patient recovered very well without any postoperative treatment.

CONCLUSION

This case is unique in that the giant breast mass initially mimicking a malignant

Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

Manuscript source: Unsolicited manuscript

Received: April 9, 2020

Peer-review started: April 9, 2020

First decision: April 29, 2020

Revised: May 11, 2020

Accepted: July 16, 2020

Article in press: July 16, 2020

Published online: August 26, 2020

P-Reviewer: Rosenberger LH

S-Editor: Zhang L

L-Editor: Webster JR

P-Editor: Liu JH



clinical presentation was eventually pathologically confirmed to be a benign PT, which misled the diagnosis and complemented the atypical features of benign PTs. The pathological and immunohistochemical results were important in the differential diagnosis. In addition, total mastectomy should be recommended due to difficulty in the precise diagnosis of PTs, especially in large breast masses. In the literature, almost one-half of super-giant benign cases were thought to be malignant tumours before surgery. This finding is a reminder to consider all conditions in order to make an accurate diagnosis and avoid excessive treatment.

Key words: Phyllodes tumour; Pulmonary neoplasms; Diagnosis; Treatment; Recurrence; Case report

©The Author(s) 2020. Published by Baishideng Publishing Group Inc. All rights reserved.

Core tip: Phyllodes tumours (PTs) are fibroepithelial breast tumours. We report the unique case of a female patient who presented with a rapidly expanding breast PT. This case shows that a giant benign PT may reveal malignant features. The clinical manifestations and imaging examinations led us to misdiagnose this mass as a malignant tumour. However, pathological diagnosis of the tumour after complete excision confirmed the tumour to be a benign PT. The lung nodule was found to be adenocarcinoma rather than metastatic tumour. We also summarized and analyzed 12 cases and the results demonstrated that we should not be fooled by appearances. All conditions should be considered to make an accurate diagnosis, in order that patients are given the appropriate treatment and avoid excessive treatment.

Citation: Zhang T, Feng L, Lian J, Ren WL. Giant benign phyllodes breast tumour with pulmonary nodule mimicking malignancy: A case report. *World J Clin Cases* 2020; 8(16): 3591-3600

URL: <https://www.wjgnet.com/2307-8960/full/v8/i16/3591.htm>

DOI: <https://dx.doi.org/10.12998/wjcc.v8.i16.3591>

INTRODUCTION

Phyllodes tumours (PTs) are rare fibroepithelial lesions with the proliferation of stromal and epithelial elements. The other two fibroepithelial lesions are fibroadenomas and hamartomas^[1]. The World Health Organization formally referred to this disease using the term “phyllodes tumour” in 2003. Phyllodes tumours exhibit a growth pattern of excessive proliferation of leaf-like stroma into dilated clefts^[2]. The incidence of PTs is low at only 2.5% of fibroepithelial lesions and 0.3%–1% of all primary breast tumours^[3]. According to their histologic grade, PTs can be classified into benign (60%-75%), borderline (15%-20%) and malignant (10%-20%)^[4-6]. Most cases occur in women aged 40-50 years, and cases in males have been reported occasionally^[7,8]. The preferred treatment for PT is surgical removal, and as lymph node metastasis is rare, axillary lymph node dissection is not routine^[2].

Many reports have described various large borderline or malignant PTs. However, a benign PT with lung nodule mimicking malignancy has not yet been reported. Here, we present the unique case of a female patient who developed a rapidly expanding PT mimicking malignancy, which misled the diagnosis. However, pathological diagnosis of the tumour after complete excision eventually confirmed a benign PT, and the accompanying lung nodule proved to be adenocarcinoma. The atypical clinical symptoms and confusing images of this benign PT make this case special. We searched PubMed from inception to September 2019 for “large OR huge OR massive OR giant OR big”, “phyllodes tumour” and “benign” as key words. We summarized and analyzed 12 cases of super-giant benign PTs (including our case) in order that doctors have a comprehensive understanding of this disease.

CASE PRESENTATION

Chief complaints

A 42-year-old Asian female presented with a rapidly enlarging right breast mass, which was originally noticed 1 year previously.

History of present illness

Over the past month, the breast mass had rapidly enlarged with symptoms of ulceration, bleeding and fever.

History of past illness

The patient had an unremarkable previous medical history.

Physical examination

On physical examination, the patient had a large right breast mass measuring 30 cm × 25 cm × 20 cm. It involved the whole right breast with a 20 cm × 15 cm area of ulceration complicated by necrosis located in the upper outer quadrant, which had a cauliflower-like neoplasm inside and was accompanied by an overpowering rotten stench. The skin of the right breast was stretched thin with superficial varicose veins, and the nipple was obviously enlarged (Figure 1A and B).

Laboratory examinations

The basic condition of the patient was poor with severe anaemia and hypoalbuminaemia at the time of admission. Blood analysis revealed red blood cells of $2.2 \times 10^{12}/L$, hemoglobin of 59 g/L, and albumin of 19.9 g/L, with a normal platelet count.

Imaging examinations

Due to ulceration and the sheer size of the breast mass, the patient was unable to undergo mammography or mammary magnetic resonance imaging. Breast ultrasound revealed an enormous mass with solid components occupying the entire breast. A contrast-enhanced chest computed tomography (CT) scan displayed the giant breast mass (Figure 2A and B), and a pulmonary nodule of 8 mm × 6 mm in the left lung, which was initially considered metastatic (Figure 2C). The right pectoralis major was coarse locally, and the boundaries between the mass and the pectoralis major were unclear, which was thought to be due to tumour invasion (Figure 2D). There were no suspicious findings in the left breast or axillary nodes, and other examinations were normal.

Further diagnostic work-up

The patient was further evaluated with a core needle biopsy. The first core needle biopsy of the massive tumour suggested mammary adenosis. However, this did not rule out the possibility of malignancy. Dillon *et al*^[5] reported that approximately 39% of breast diseases may give false negative results. In addition, as the mass was large, the core needle biopsy was unable to cover the entire area. A second biopsy was then performed, which consequently revealed lymphadenitis with neoplastic cells. Both biopsies found no evidence of malignant tumour. Subsequently, a right mastectomy without resection of the axillary lymph nodes was recommended^[5].

FINAL DIAGNOSIS

According to postoperative histological examination, the patient was diagnosed with benign PT.

TREATMENT

The patient refused the suggested reconstructive breast surgery, due to its high cost. A right mastectomy without resection of the axillary lymph nodes was performed. The minimum surgical margin of the mass was 1 cm. Accordingly, making use of the superior and inferior skin flaps (even the skin directly overlying the mass which was normal) (Figure 3A and B), the skin closure was approximated after excision of the

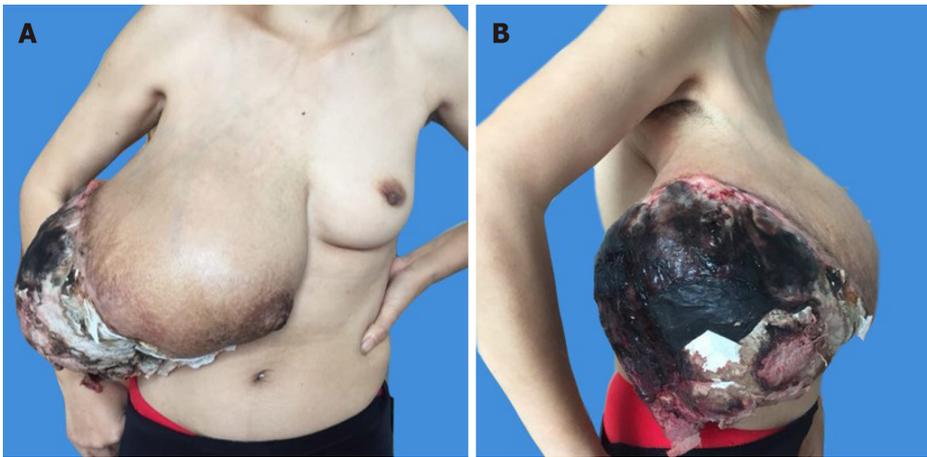


Figure 1 A giant phyllodes tumor of the right breast in a 42-year-old woman: A: Front image; and B: Lateral image.

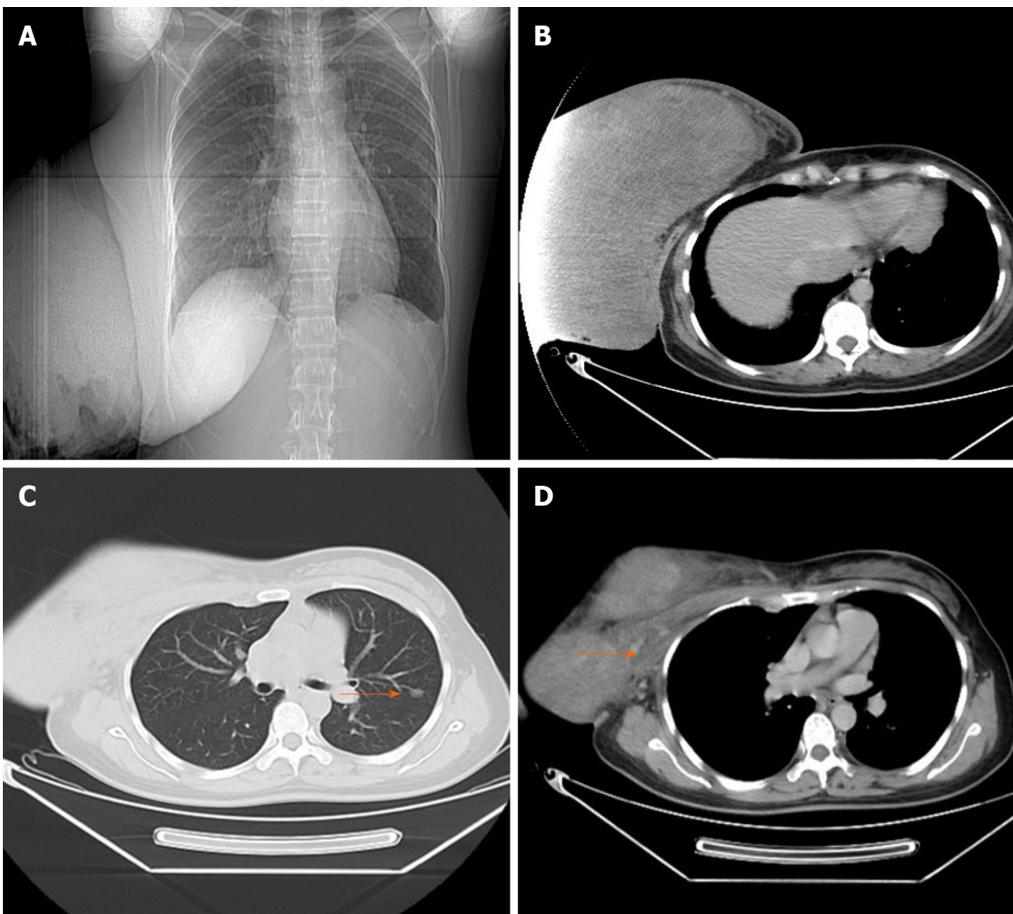


Figure 2 Preoperative radiologic evaluation. A and B: Chest computed tomography showed a huge mass; C: A pulmonary nodule was seen in the left lung (arrowheads); and D: Preoperative computed tomography showed that there was no clearance between the mass and pectoralis major (arrowheads).

giant mass. Dissection revealed that the tumour was partly adhered to the pectoralis major muscle, rather than invading the muscle.

The excised breast mass was 25 cm × 20 cm × 16 cm in size, and weighed 3.5 kg. Due to severe ulceration of the tumour before surgery, the patient underwent debridement and dressing changes every day. The weight of the tumour was reduced compared with that on admission. Cystic and solid lobulated changes could be seen after incision of the tumour (Figure 4A). Postoperative histological examination was consistent with benign PT with negative margins (Figure 4B). Histologic sections revealed a circumscribed lesion with a variable leaf-like growth pattern (Figure 4C), low-to-moderate stromal cellularity, minimal stromal cell atypia, and absent stromal



Figure 3 Making use of the superior and inferior skin flaps and even the skin directly overlying the mass which was normal. A: Preoperative photograph of the design to allow skin approximation and closure after removal of the large tumour; and B: Postoperative photograph after tumour resection and skin closed with placement of two drains under the flaps.

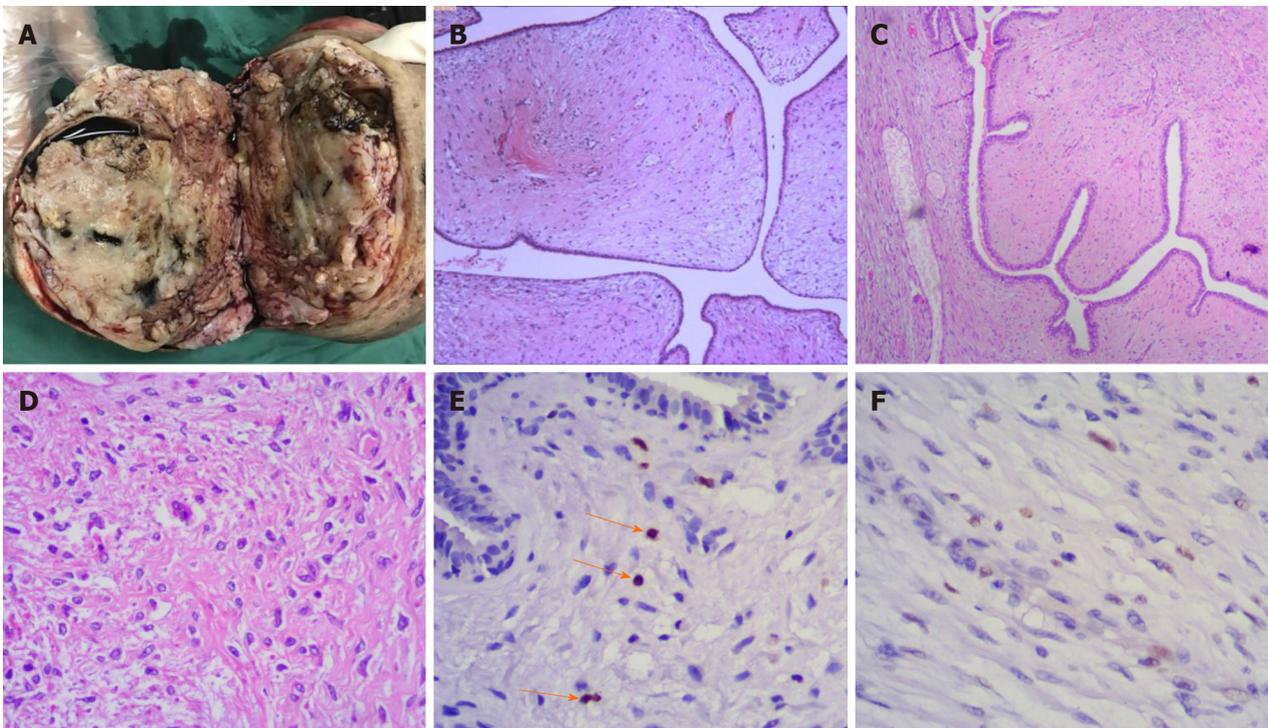


Figure 4 The tissue section showing benign phyllodes tumor. A: Cystic components after incision of the tumour; B: (10 ×) Well-circumscribed fibroepithelial neoplasm; C: (40 ×) Prominent leaf-like architecture and areas of hypocellular stroma; D: (400 ×) Bland stromal spindle cells without mitoses or nuclear atypia; E: Ki-67 proliferation index of the tumour was 1 for the stromal component; and F: The P53 index of the stromal component was focally positive.

overgrowth and mitoses (Figure 4D). The Ki-67 proliferation index of the tumour was 1% for the stromal component (Figure 4E). The P53 index for the stromal component was focally positive and consistent with benign stromal proliferation (Figure 4F). All of these analyses showed no evidence of malignancy.

OUTCOME AND FOLLOW-UP

The patient has recovered very well without any postoperative treatment. No recurrence or metastasis was observed 12 mo after her breast operation, and wedge resection of the small lung nodule was performed after 1 year of follow-up, which confirmed the pulmonary nodule to be adenocarcinoma, rather than metastatic breast

cancer (Figure 5A and B). Immunohistochemical results showed that thyroid transcription factor-1 and napsin-A were positive (Figure 5C and D) and GCDPF-15 was negative (Figure 5E). Ki-67 was focally positive (Figure 5F). The patient provided written informed consent for publication of the case details.

DISCUSSION

This case is interesting in that the patient had a giant benign PT with a pulmonary nodule mimicking malignancy, the patient also had severe anaemia, hypoalbuminaemia and infection. It was confusing and extremely difficult to diagnose whether the tumour was benign or malignant at first due to the following features: (1) The mass which was more than 30 cm with skin ulcers and necrosis increased rapidly in size in a short time. The patient had significant anaemia and malnutrition, which was more like cachexia. (2) CT showed there were no clear boundaries between the mass and pectoralis major, which was considered tumour invasion. (3) The pulmonary nodule was thought to be metastasis preoperatively, according to the chest CT and her breast disease. These findings suggested a diagnosis of malignant PT until postoperative pathology proved the breast mass was a benign PT. The reasons for this confusion were as follows: (1) Abundant vessels supported the tumour, resulting in the patient's poor overall condition and cachexia. (2) Intraoperative findings revealed there was no infiltration between the mass and pectoralis major, only local adhesion. (3) The pulmonary nodule, which was resected by video-assisted thoracic surgery, was a primary lung adenocarcinoma, rather than breast metastasis.

The average size of PTs are usually 4-7 cm^[9]. Giant PTs usually have a diameter of more than 10 cm^[10]. It was reported that the largest benign PT even exceeded 50 cm^[4]. We searched PubMed from inception to September 2019 for "large OR huge OR massive OR giant OR big", "phyllodes tumour" and "benign" as key words. We summarized and analyzed 12 cases of super-giant benign PTs (including our case) that exceeded 20 cm. These cases as well as the one presented are shown in Table 1^[4]. The majority of cases presented with rapid growth. The clinical characteristics of seven cases, including our case, were ulceration, bleeding or infection^[1,2,6,7,10-12]. The huge masses in four cases were adherent to the pectoral muscle^[3,10,11,12]. Our case is unique in that it is the first report of a patient with a super-giant PT presenting atypical clinical manifestations with a co-diagnosis of lung adenocarcinoma. In the literature, only one other patient presented atypical clinical manifestations^[11], and in one other case the PT was combined with another malignant tumour^[8].

An interesting fact about this case is our patient's manifestations and imaging examinations led us to misdiagnose the mass as a malignant tumour. However pathological diagnosis of the tumour after complete excision confirmed it was a benign PT. Of the 12 cases of super-giant benign PTs diagnosed according to the postoperative pathology results, five cases were thought to be malignant tumours before surgery, accounting for almost one-half^[1,3,6,8,12]. This finding shows that super-giant PTs may be more likely to reveal "malignant features". Due to oversensitivity to "malignant features", excessive treatments were performed in these cases, including pectoral muscle excision^[3,7,8,10], peripheral muscle excision^[4], and lymph node sampling^[5,7,8]. Most of the patients were stable without recurrence during the follow-up period. Only one patient died after 6 mo due to malignant pleural effusion; however, the patient had no prior history of lung disease and her breast tumour was a benign PT^[7].

To avoid confusion regarding the diagnosis, and being misled by the atypical clinical presentation of the mass, and precisely differentiating benign from malignant PTs, it is more important to pay attention to the pathology results. Histologically, the lobulated structure is typical, and the mesenchymal cells in the lower epithelium undergo significant proliferation, which is helpful in the diagnosis of PT. According to World Health Organization recommendations, PTs can be classified into benign, borderline and malignant tumours depending on their histological features, including stromal hypercellularity, stromal atypia, mitosis, stromal overgrowth, and tumour margins (Table 2). Sometimes, the distinction between benign, borderline and malignant tumours may be particularly difficult from core biopsies, Dillon *et al*^[5] reported that approximately 39% of false negative results were obtained. In addition, the mass was too large for the core needle biopsy to cover the entire area in our patient. Thus, immunohistochemistry can be helpful. One study by Kleer *et al*^[6] showed that the Ki-67 labelling index was notably higher in high-grade malignant tumours compared to low-grade malignant tumours, and the Ki-67 labelling index in the low-grade malignant PT group was notably higher than that in the benign PT

Table 1 Case reports of giant benign phyllodes tumours with a diameter ≥ 20 cm

Ref.	Age	Disease duration	Size/weight	Clinical characteristics	Pectoral muscle adherence	Pre-op diagnosis/core biopsy	Surgery/margins	Recurrences
Miyaguni <i>et al</i> ^[20]	43	5 mo	20 cm/unknown	Ulceration, bleeding	No	Malignant/Unknown	Mastectomy/unknown	Unknown
Udapudi <i>et al</i> ^[21]	21	3 mo	45 cm/6.5 kg	Ulceration	No	Benign PT/Benign PT	Mastectomy/unknown	None for 2 yr
Liang <i>et al</i> ^[22]	64	2 yr	36 cm/unknown	No	Yes, suggested invasion	Malignancy not excluded/Highly atypical cells	Mastectomy, partial PME /< 1 cm	None for 7 yr
Zhao <i>et al</i> ^[23]	63	2 yr	45 cm/11 kg	No	No	Unknown/Unknown	Mastectomy, peripheral muscle excision /unknown	Unknown
Likhitmaskul <i>et al</i> ^[24]	35	5 mo	20 cm/unknown	Gestation	No	Benign PT/Benign PT	Mastectomy, LNs removed/unknown	Unknown
Sbeih <i>et al</i> ^[25]	41	7 yr	25 cm/unknown	Ulceration	No	Malignancy not excluded/Pseudoangiomatous or PT	Mastectomy, skin graft/negative margins	Unknown
Islam <i>et al</i> ^[4]	44	1 yr	50 cm/unknown	Ulceration, fungating, anaemia, malnourished	No	Benign PT/Benign PT	Mastectomy, partial PME, LN samples, LD flap closure/unknown	Died of MPE after 6 mo
Yan <i>et al</i> ^[26]	54	6 mo	20 cm/unknown	Non-myoeptithelial tumour of the parotid	No	Malignancy not excluded/Fibroepithelial lesion	Mastectomy, PME, LN samples/adequate margins	None for 3 mo
Kallam <i>et al</i> ^[27]	32	8 mo	20 cm/unknown	Gestation	No	Benign PT/Benign PT	Mastectomy/unknown	None for 4 wk
Rathore <i>et al</i> ^[28]	25	6 wk	30 cm/5 kg	Fungating, anaemia	Yes	Benign PT/Benign PT	Mastectomy, PME/> 2 cm	None for 10 mo
Benoit <i>et al</i> ^[29]	40	1 mo	29 cm/4 kg	Ulceration, bleeding, infection	Yes	Benign PT/Adenofibroma or benign PT	Mastectomy/< 1 mm	Unknown
Our case	42	2 mo	30 cm/3.5 kg	Ulceration, bleeding, infection, Lung adenocarcinoma	Yes, suggested invasion	Malignancy not excluded/Benign PT	Mastectomy/< 1 mm	None for 12 mo

PT: Phyllodes tumour; PME: Pectoral muscle excision; LN: Lymph node; Pre-op: Preoperative; LD: Latissimus dorsi breast reconstruction; MPE: Malignant pleural effusion.

Table 2 Histologic features of benign, borderline and malignant phyllodes tumors (adopted from the World Health Organization classification 2012)

Histologic features	Benign	Borderline	Malignant
Stromal hypercellularity	Mild	Moderate	Marked
Stromal mitotic activity	0-4/10 HPF	5-9/10 HPF	$\geq 10/10$ HPF
Stromal cell atypia	Mild	Moderate	Marked
Stromal overgrowth	Absent	Absent or focal	Often present
Tumour borders	Well-defined	Well-defined focally infiltrative	Infiltrative
Malignant heterologous elements	Absent	Absent	May be present

group ($P = 0.012$). P53 has also been used for distinction; however, both benign and malignant PTs show focally positive p53 occasionally. Bode *et al*^[7] reported that p63, p40 and cytokeratin were only labelled in malignant tumours. Fibroadenomas, benign PTs, and borderline PTs are not labelled in this way. A study by Chia *et al*^[8] revealed

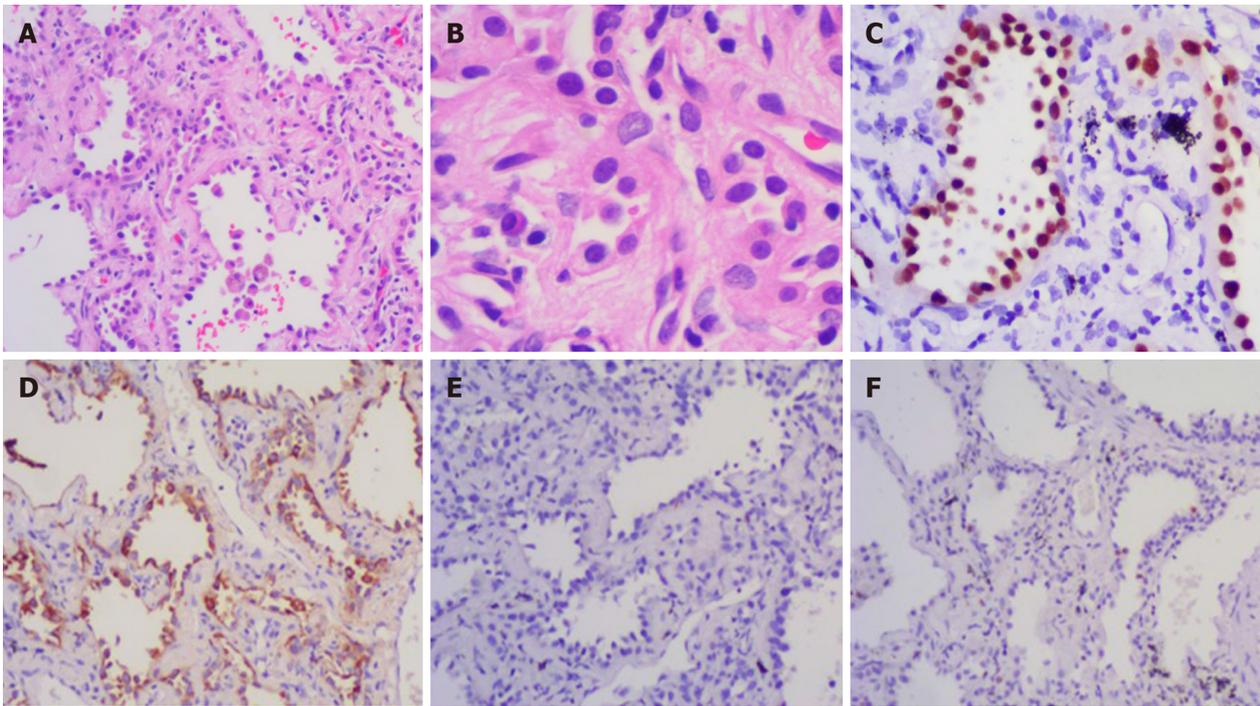


Figure 5 The tissue section showing lung adenocarcinoma. A: (100 ×) Pathological examination indicated lung adenocarcinoma; B: (400 ×) Tumour cells showed prominent atypia at high magnification; C: (200 ×) Thyroid transcription factor-1 was positive; D: (200 ×) Napsin-A was positive; E: GCDFP-15 was negative; and F: Ki-67 was focally positive.

that cytokeratin can be focally positive in malignant tumours (1%-5%), which increases with PT grade. Another study found that p40 was more specific, but less sensitive, in distinguishing sarcomatoid carcinoma from malignant PT than p63, but this study requires further validation^[11].

For primary treatment of PT, surgical resection is recommended. Previously, simple mastectomy was suggested for borderline and malignant PTs in order to reduce the recurrence rate. However, recent research has revealed that the survival after mastectomy and wide local excision with postoperative radiotherapy was equivalent^[12]. Consequently, conservative surgery is recommended, but mastectomy should be carried out according to the following reasons: Benign or borderline tumours at least 8 cm in size, malignant PTs, or positive margins^[13]. Margins of at least 1 cm with wide local excision were recommended by the National Comprehensive Cancer Network guideline for each PT grade. It is widely known that surgical margin status is an important risk factor for local recurrence. However, a meta-analysis demonstrated that margin positivity was a higher local recurrence (LR) risk only for malignant PTs, but was not associated with benign and borderline PTs^[14]. In our case with a benign PT without distant metastasis, mastectomy was the better choice due to the large size of the tumour. The other 11 cases with huge benign PTs also underwent mastectomy. If the tumour size is less than 20 cm, wide local excision with a margin of at least 1 cm is preferred as the initial treatment. Lymph node metastasis is rare in PTs^[2]. It is not necessary to routinely dissect axillary lymph nodes. However, sentinel lymph node biopsy or low-grade axillary lymph node dissection is recommended if palpable lymph nodes are detected pathologically^[15]. After mastectomy for giant benign PTs, some cases chose reconstruction with a latissimus dorsi musculocutaneous flap, transverse rectus abdominis myocutaneous flap, or skin grafting. The study by Kuo *et al*^[16] revealed that, for initial unresectable giant PTs, transcatheter arterial chemoembolization prior to surgery is recommended to improve the resectability of PTs without requiring skin grafting. In addition, postoperative chemotherapy and endocrine therapy have no significant effect on PTs, especially with regard to reducing the rate of recurrence or death.

According to a meta-analysis which included 9234 cases, the pooled LR rates for benign, borderline and malignant PTs were 8%, 13% and 18%, respectively, and the ranges of the 5-year cumulative LR risks for benign, borderline and malignant PTs were 3%-23%, 9%-55% and 14.8%-55%, respectively^[14]. Other risk factors for LR also include mitoses, tumour border, stromal cellularity, stromal atypia, stromal overgrowth, tumour necrosis, and type of surgery. Another study revealed that

patients without MED12 mutations had a higher likelihood of recurrence, whereas the disease-free survival of patients with PTs was improved with the occurrence of MED12 mutations^[17]. Compared to the primary tumour, some studies have shown that similar or lower histological grading may occur during recurrences^[18]. However, other studies have shown that the primary benign PT recurred as a malignant lesion^[18,19]. For malignant PTs, the most common sites of metastasis included the lung (70% to 80%), pleura (60% to 70%), and bone (20% to 30%)^[20].

CONCLUSION

PT is a rare fibroepithelial breast tumour. We report the unique case of a female patient who presented with a rapidly expanding breast PT. This case shows that a giant benign PT may reveal malignant features. The clinical manifestations and imaging examinations led us to misdiagnose the tumour as malignant. However, pathological diagnosis of the tumour after complete excision confirmed that it was a benign PT. Pathological and immunohistochemical results are important to differentiate this disease, and the lung nodule proved to be adenocarcinoma, rather than a metastatic tumour. These results are a reminder that we should not be fooled by appearances. All conditions should be considered to make an accurate diagnosis, in order that patients receive appropriate treatment and avoid excessive treatment^[20-29].

ACKNOWLEDGEMENTS

We would like to thank Dr. Xiao-Song Chen (Ruijin Hospital) for additional editorial assistance.

REFERENCES

- 1 **Krings G**, Bean GR, Chen YY. Fibroepithelial lesions; The WHO spectrum. *Semin Diagn Pathol* 2017; **34**: 438-452 [PMID: 28688536 DOI: 10.1053/j.semdp.2017.05.006]
- 2 **Lenhard MS**, Kahlert S, Himsel I, Ditsch N, Untch M, Bauerfeind I. Phyllodes tumour of the breast: clinical follow-up of 33 cases of this rare disease. *Eur J Obstet Gynecol Reprod Biol* 2008; **138**: 217-221 [PMID: 17868973 DOI: 10.1016/j.ejogrb.2007.08.002]
- 3 **Tan PH**, Thike AA, Tan WJ, Thu MM, Busmanis I, Li H, Chay WY, Tan MH; Phyllodes Tumour Network Singapore. Predicting clinical behaviour of breast phyllodes tumours: a nomogram based on histological criteria and surgical margins. *J Clin Pathol* 2012; **65**: 69-76 [PMID: 22049216 DOI: 10.1136/jclinpath-2011-200368]
- 4 **Islam S**, Shah J, Harnarayan P, Naraynsingh V. The largest and neglected giant phyllodes tumor of the breast-A case report and literature review. *Int J Surg Case Rep* 2016; **26**: 96-100 [PMID: 27475116 DOI: 10.1016/j.ijscr.2016.07.022]
- 5 **Dillon MF**, Quinn CM, McDermott EW, O'Doherty A, O'Higgins N, Hill AD. Needle core biopsy in the diagnosis of phyllodes neoplasm. *Surgery* 2006; **140**: 779-784 [PMID: 17084721 DOI: 10.1016/j.surg.2006.03.022]
- 6 **Kleer CG**, Giordano TJ, Braun T, Oberman HA. Pathologic, immunohistochemical, and molecular features of benign and malignant phyllodes tumors of the breast. *Mod Pathol* 2001; **14**: 185-190 [PMID: 11266524 DOI: 10.1038/modpathol.3880282]
- 7 **Bode MK**, Rissanen T, Apaja-Sarkkinen M. Ultrasonography and core needle biopsy in the differential diagnosis of fibroadenoma and tumor phyllodes. *Acta Radiol* 2007; **48**: 708-713 [PMID: 17728999 DOI: 10.1080/02841850701367911]
- 8 **Chia Y**, Thike AA, Cheok PY, Yong-Zheng Chong L, Man-Kit Tse G, Tan PH. Stromal keratin expression in phyllodes tumours of the breast: a comparison with other spindle cell breast lesions. *J Clin Pathol* 2012; **65**: 339-347 [PMID: 22259180 DOI: 10.1136/jclinpath-2011-200377]
- 9 **Barrio AV**, Clark BD, Goldberg JI, Hoque LW, Bernik SF, Flynn LW, Susnik B, Giri D, Polo K, Patil S, Van Zee KJ. Clinicopathologic features and long-term outcomes of 293 phyllodes tumors of the breast. *Ann Surg Oncol* 2007; **14**: 2961-2970 [PMID: 17562113 DOI: 10.1245/s10434-007-9439-z]
- 10 **Mishra SP**, Tiwary SK, Mishra M, Khanna AK. Phyllodes tumor of breast: a review article. *ISRN Surg* 2013; **2013**: 361469 [PMID: 23577269 DOI: 10.1155/2013/361469]
- 11 **Cimino-Mathews A**, Sharma R, Illei PB, Vang R, Argani P. A subset of malignant phyllodes tumors express p63 and p40: a diagnostic pitfall in breast core needle biopsies. *Am J Surg Pathol* 2014; **38**: 1689-1696 [PMID: 25046342 DOI: 10.1097/PAS.0000000000000301]
- 12 **Barth RJ Jr**, Wells WA, Mitchell SE, Cole BF. A prospective, multi-institutional study of adjuvant radiotherapy after resection of malignant phyllodes tumors. *Ann Surg Oncol* 2009; **16**: 2288-2294 [PMID: 19424757 DOI: 10.1245/s10434-009-0489-2]
- 13 **Wang Y**, Zhang Y, Chen G, Liu F, Liu C, Xu T, Ma Z. Huge borderline phyllodes breast tumor with repeated recurrences and progression toward more malignant phenotype: a case report and literature review. *Onco Targets Ther* 2018; **11**: 7787-7793 [PMID: 30464526 DOI: 10.2147/OTT.S171714]

- 14 **Lu Y**, Chen Y, Zhu L, Cartwright P, Song E, Jacobs L, Chen K. Local Recurrence of Benign, Borderline, and Malignant Phyllodes Tumors of the Breast: A Systematic Review and Meta-analysis. *Ann Surg Oncol* 2019; **26**: 1263-1275 [PMID: 30617873 DOI: 10.1245/s10434-018-07134-5]
- 15 **Parker SJ**, Harries SA. Phyllodes tumours. *Postgrad Med J* 2001; **77**: 428-435 [PMID: 11423590 DOI: 10.1136/pmj.77.909.428]
- 16 **Kuo CY**, Lin SH, Lee KD, Cheng SJ, Chu JS, Tu SH. Transcatheter arterial chemoembolization improves the resectability of malignant breast phyllodes tumor with angiosarcoma component: a case report. *BMC Surg* 2019; **19**: 100 [PMID: 31351458 DOI: 10.1186/s12893-019-0562-0]
- 17 **Ng CC**, Tan J, Ong CK, Lim WK, Rajasegaran V, Nasir ND, Lim JC, Thike AA, Salahuddin SA, Iqbal J, Busmanis I, Chong AP, Teh BT, Tan PH. MED12 is frequently mutated in breast phyllodes tumours: a study of 112 cases. *J Clin Pathol* 2015; **68**: 685-691 [PMID: 26018969 DOI: 10.1136/jclinpath-2015-202896]
- 18 **Borhani-Khomani K**, Talman ML, Kroman N, Tvedskov TF. Risk of Local Recurrence of Benign and Borderline Phyllodes Tumors: A Danish Population-Based Retrospective Study. *Ann Surg Oncol* 2016; **23**: 1543-1548 [PMID: 26714948 DOI: 10.1245/s10434-015-5041-y]
- 19 **Muller KE**, Tafe LJ, de Abreu FB, Peterson JD, Wells WA, Barth RJ, Marotti JD. Benign phyllodes tumor of the breast recurring as a malignant phyllodes tumor and spindle cell metaplastic carcinoma. *Hum Pathol* 2015; **46**: 327-333 [PMID: 25476122 DOI: 10.1016/j.humpath.2014.10.014]
- 20 **Miyaguni T**, Deguchi S, Teruya J, Kuniyoshi S, Tomita S, Soda N, Muto Y. Phyllodes Tumor of the Breast with a Grossly Malignant Appearance: A Case Report. *Breast Cancer* 1998; **5**: 205-208 [PMID: 11091650 DOI: 10.1007/bf02966697]
- 21 **Udapudi DG**, Vasudeva P, Srikantiah R, Virupakshappa E. Massive benign phyllodes tumor. *Breast J* 2005; **11**: 521 [PMID: 16297126 DOI: 10.1111/j.1075-122X.2005.00149.x]
- 22 **Liang MI**, Ramaswamy B, Patterson CC, McKelvey MT, Gordillo G, Nuovo GJ, Carson WE 3rd. Giant breast tumors: surgical management of phyllodes tumors, potential for reconstructive surgery and a review of literature. *World J Surg Oncol* 2008; **6**: 117 [PMID: 19014438 DOI: 10.1186/1477-7819-6-117]
- 23 **Zhao Z**, Zhang J, Chen Y, Shen L, Wang J. An 11 kg Phyllodes tumor of the breast in combination with other multiple chronic diseases: Case report and review of the literature. *Oncol Lett* 2013; **6**: 150-152 [PMID: 23946794 DOI: 10.3892/ol.2013.1361]
- 24 **Likhitmaskul T**, Asanprakit W, Charoenthammaraksa S, Lohsiriwat V, Supaporn S, Vassanasiri W, Sattaporn S. Giant benign phyllodes tumor with lactating changes in pregnancy: a case report. *Gland Surg* 2015; **4**: 339-343 [PMID: 26312220 DOI: 10.3978/j.issn.2227-684X.2015.01.09]
- 25 **Sbeih MA**, Engdahl R, Landa M, Ojutiku O, Morrison N, Depaz H. A giant phyllodes tumor causing ulceration and severe breast disfigurement: case report and review of giant phyllodes. *J Surg Case Rep* 2015; **2015** [PMID: 26703928 DOI: 10.1093/jscr/rjv162]
- 26 **Yan Z**, Gudi M, Lim SH. A large benign phyllodes tumour of the breast: A case report and literature review. *Int J Surg Case Rep* 2017; **39**: 192-195 [PMID: 28854407 DOI: 10.1016/j.ijscr.2017.08.039]
- 27 **Kallam AR**, Kanumury V, Korumilli RM, Gudeli V, Polavarapu H. Massive Benign Phyllodes Tumour of Breast Complicating Pregnancy. *J Clin Diagn Res* 2017; **11**: PD08-PD09 [PMID: 28658847 DOI: 10.7860/jcdr/2017/26277.9929]
- 28 **Rathore AH**. Huge Fungating Benign Phyllodes Tumor of Breast. *Pak J Med Sci* 2018; **34**: 770-771 [PMID: 30034456 DOI: 10.12669/pjms.343.15548]
- 29 **Benoit L**, Ilenko A, Chopier J, Buob D, Darai E, Zilberman S. A rare case of a giant ulcerated benign phyllode tumor. *J Gynecol Obstet Hum Reprod* 2019; **48**: 217-220 [PMID: 30142471 DOI: 10.1016/j.jogoh.2018.08.007]



Published by **Baishideng Publishing Group Inc**
7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA

Telephone: +1-925-3991568

E-mail: bpgoffice@wjgnet.com

Help Desk: <https://www.f6publishing.com/helpdesk>

<https://www.wjgnet.com>

