

Case Report

Breast Angiosarcoma in a Young Patient, Case Report

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Abstract

We show a case report on primary breast angiosarcoma, clinical presentation, histopathology findings, and a brief revision about breast angiosarcoma. The patient develops a rapidly grown breast tumor, without diagnosis until she arrived at oncology gynecology services. She is 22 years old; this kind of angiosarcoma is not relational to radiotherapy or family cancer syndrome.

Ower group performed a simple mastectomy plus major pectoral resection, no axillary dissection was necessary and chemotherapy treatment. The patient has 5 months of free period disease without relapse symptoms. Ultrasound or mammography is not much useful in diagnosis about it. Magnetic resonance is more helpful in non-epithelial breast tumors. Free border resection, the grade of the tumor, and suspect to the aggressive tumor are strong points to consider in this kind of breast tumor. Chemotherapy until now is not beneficial in adjuvant treatment just palliative election. Survival media time is 35 months to 40 months after treatment.

Keywords: Breast angiosarcoma; Primary angiosarcoma; Breast rare tumors; Cancer

Introduction

Angiosarcomas of the breast are an infrequent clinical condition, accounting for 0.04% in all breast cancers. Luini in Milan estimated an incidence between 0.002 to 0.005 per year. Breast angiosarcomas are divided into two categories, primary and secondary (radiation-associated) [1]. Those tumors share similar histological features, clinical appearance is alike, but with radiotherapy and surgery precedent and the prognosis is bad in both cases. The clinical presentation in primary angiosarcoma is two decades or three before than secondary angiosarcoma [2,3]. There is the association between primary angiosarcoma and onset pregnancy but do not is solid this affirmation maybe is a coincidence in the age group and some authors are reported primary angiosarcoma men [4].

Primary angiosarcoma is a malignant vascular neoplasm that arises within the breast parenchyma, while secondary angiosarcoma usually arises in the cutaneous tissue and might invade the breast parenchyma secondarily. In most series, patients with secondary angiosarcoma had significantly shorter survival time than those who

had primary angiosarcoma and were more prone to local and distant recurrence. Ming and coworkers reported that results of a clinical trial suggested that secondary breast angiosarcoma had more aggressive tumor phenotype than primary angiosarcoma at presentation [5].

Clinical Case

The patient is a 22 years old healthy female, In December 2019 began with pain and increased volume in the right breast; after that she takes an appointment with a general gynecologist and prescribes him, topic progestagen cream without response. The tumor was continuous growing on breast and pain, then get oncology gynecology by an ower group and we performed a cut needle biopsy with report primary breast angiosarcoma. A total mastectomy with pectoralis major muscle was performed without complications and she was discharged from hospital 2 days after surgery (Figure 1).

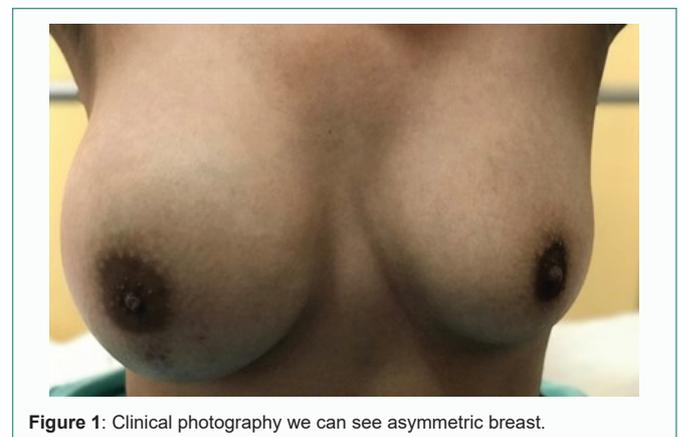


Figure 1: Clinical photography we can see asymmetric breast.

Citation: Valdespino VE, Matamoros IL, Hernandez MR, Cordova GM, Quesada AC, Gomez VV. Breast Angiosarcoma in a Young Patient, Case Report. Surg Clin J. 2020; 2(2): 1021.

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Publisher Name: Medtext Publications LLC

Manuscript compiled: Jun 22th, 2020

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The pathological final report is breast angiosarcoma high grade, 10 cm major diameter, all edges were tumor negatives more than 2 cm. No axillary dissection was performed. We used to scan tomography because obtain magnetic resonance in some hospitals in Mexico is difficult (Figure 2). The patient received radiotherapy 50 Gy in a field breast with toxicity grade I - II without complications. After 5 months of free period disease, she does not have signals of local recurrence or axillar or distant metastasis however the patient has relapse risk distant or local.

We present a case report and a review of breast angiosarcoma. Angiosarcoma breast is reported from 17 years to 84 years old in a female patient the present case is on a very young patient, this disease is very infrequent in young persons and de clinical presentations do not have risk factors in primary breast angiosarcoma. She does not report history about family cancer.

Discussion

Primary sarcomas of the breast are an infrequent clinical condition they are an incidence of fewer less than 1% of all breast cancers. Sarcomas arise from mesenchymal tissue of the mammary gland one of the most common histological subtypes of primary sarcoma of the breast is angiosarcoma. There is primary angiosarcoma this tumor develops during the third or fourth decades of life (Figure 3). And the secondary breast angiosarcoma, radiation-induced soft tissue sarcoma, after conservative surgical treatment and radiotherapy on the breast.

Adem and his team found that angiosarcoma and fibrosarcoma in the breast are most frequent breast sarcomas, and the size tumor is the most important survival prognostic factor, and they do not launch axillar nodal metastasis [6].

Primary angiosarcoma of the breast typically presents as a palpable breast mass in women in the third and fourth decade of life. Secondary angiosarcomas of the breast are generally found in older women following breast cancer treatment. Secondary angiosarcoma may present with skin lesions such as nodules or discoloration, due to damage of lymphatic or vascular channels following surgery and irradiation. There is typically a latency period of 10 years to 20 years following breast conservation and radiotherapy [7].

There are Angiosarcoma Project (ASC project), an initiative enabling US and Canadian patients to remotely share their clinical information and biospecimens for research. There are Angiosarcoma Project (ASC project) an initiative from the US and Canada, it can help to share remotely clinical information about angiosarcoma for current or futures research on this topic [8]. This group found PIK3CA mutated genes for angiosarcoma has taken more importance; this topic in particular for breast angiosarcoma, they found the response of pembrolixumab in those kinds of patient's even complete responses in two patients in metastatic disease. Targeted inhibition of the PIK3, signaling pathway can have deep effects on cancer cell growth through numerous mechanisms [9]. How is the best target the PIK3 pathway in cancer treatment remains a topic of active investigation in particular for breast angiosarcoma?

Relapse metastatic angiosarcoma for a medical oncologist is a serious challenge, standard chemotherapy nor have targeted agents demonstrated good and durable responses. Drugs used for metastatic angiosarcoma include doxorubicin, paclitaxel, docetaxel, and bevacizumab. Small case series have documented durable benefits of metronomic chemotherapy in relapsing angiosarcoma [10].

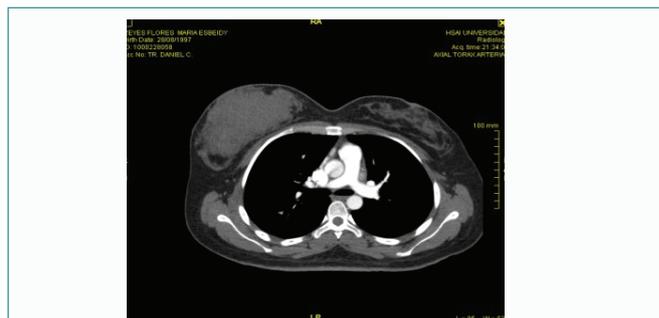


Figure 2: Chest Tomography, we can look a great breast tumor.

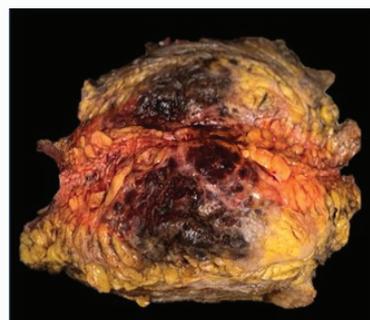


Figure 3: Whole breast tumor with ink.

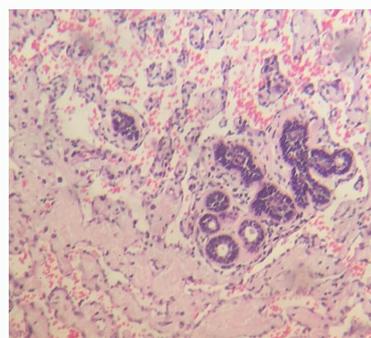


Figure 4: Lobular breast tissue surrounded by several vascular canal anastomosis (microphotograph).

Sher and coworkers reported that adjuvant chemotherapy using anthracycline and ifosfamide or gemcitabine and a taxane did not significantly improve recurrence-free survival compared with patients who did not receive chemotherapy treatment [11]. However, the administration of palliative chemotherapy resulted in a 48% response rate. In the case of secondary angiosarcomas induced by radiation treatment docetaxel showed promise response [12]. In other small series of Biswas and workers reported a median overall survival was 37.4 months (range, 8.7 - 92.8 months). The median RFS was 17.9 months (range, 2.5 - 69.4 months). They used to as chemotherapy paclitaxel and gemcitabine like palliative treatment. In addition to it, Four out of 5 patients had negative mammograms [13].

In Sri Lanka reported a bilateral synchronic angiosarcoma after surgery and radiotherapy (secondary breast angiosarcoma) in a patient of 62 years old, she develops those tumors with only 5 years of history to radiotherapy, the difference with frequent radio induced tumor between 10 or 20 years [14]. Ragavan in Singapore and coworkers give treatment to 13 patients with angiosarcoma, 5 patients

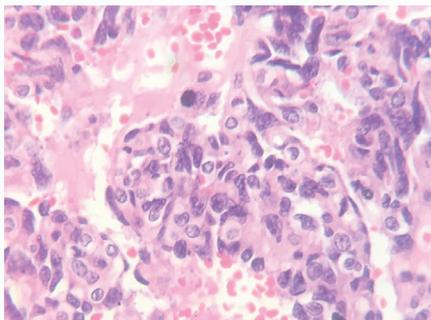


Figure 5: Tumor solid form with a vascular branch growing and mitotic activity.

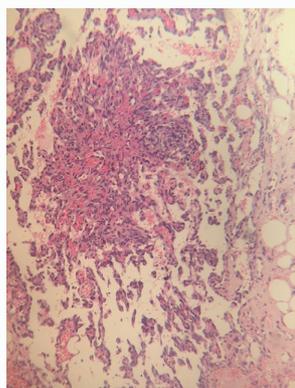


Figure 6: Microphotograph show intravascular papillary growing.

received any axillary surgical treatment, and they do not detect axillary nodal metastasis and neither axillary relapse. Recurrence was usual to the bone, liver, ovary, contralateral breast, peritoneum [15].

Breast angiosarcoma has limited available literature with small and inconclusive results, but surgery treatment with wide and clear tumor border is the best treatment, tumor size and grade are principal prognostic factors (like others sarcomas) adjuvant treatment is unclear in results, axillary nodal treatment is not necessary, molecular trigger therapy someday will be a useful treatment in the metastatic stage. The tumor cells from breast angiosarcoma proved to be immunoreactive to vascular markers (CD31 and CD34) and negative to epithelial markers; is helpful to confirm breast angiosarcoma but is very important suspect it with the clinical appearance [16].

However we do not use immunochemistry in the present case. Mammographic and sonographic findings are non-specific and can be normal in up to a third of cases. Sonographic findings are non-specific, and the mass can be hyperechoic, hypoechoic, or heterogeneous. Heterogeneous sonographic appearances may be due to hemorrhage of the tumor into the surrounding stroma. In secondary angiosarcoma, dermal lesions may be difficult to distinguish from post-irradiation skin thickening, skin thickening could be the only stage found in ultrasound [17]. In primary angiosarcoma, a heterogeneous mass is demonstrated on MR imaging with low signal intensity on T1 images and high signal on T2 images (Figures 4-6). On dynamic phase imaging, kinetic enhancement depends on tumor grade. High-grade tumors may demonstrate rapid enhancement with washout kinetics, but lower-grade tumors usually demonstrate plateau or persistent enhancement kinetics [18].

Conclusion

Breast angiosarcoma is an aggressive tumor with poor survival, five year overall rates range 38% to 78%, and 5 years free survival between 38% and 56%. Size tumor looks to be the most important factor in survival, (less than 5 cm). Early diagnosis with correct (with R0 or negative margins) surgical treatment can give patients a good quality of life, but in a rare tumor we need more reports also clinical essays if we want to improve the life of these patients. Axillary nodal dissection is not helpful for this tumor. As well as advanced tumor grade (grade III) is associated with significantly worse overall survival. Adjuvant treatment is not much useful for radiotherapy or chemotherapy. The early clinical suspect is the key and does not delay treatment until now is the correct behavior.

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