Cavernous Lymphangiomas of the Breast Mimicking Breast Cancer

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Lymphangiomas are congenital malformations of the lymphatic system and are thought to result from failure of the lymphatic system to connect with the venous system. Lymphangiomas in the breast are rare entities, and only a few cases have been documented in the literature.1,2 Its sonographic appearance is reported as similar to that of a cluster of simple cysts or dilated ducts; however, the echogenicity depends on the size of the lymphatic channels, similar to a hemangioma. Here, we report a case of breast lymphangiomas mimicking multifocal breast cancer in a 34-year-old woman.

Case Report

A 34-year-old woman came to our hospital with a 2-month history of a palpable mass in her left breast. Physical examination revealed a movable nontender mass located in the upper outer quadrant and another soft mass measuring 1.5 cm located in the left axilla. The overlying skin was not remarkable. She had undergone simple excision of a benign mass in the left axilla 10 years previously. We performed sonography first because of her young age. On sonography, the palpable mass in the left upper outer breast was revealed as an irregular spiculated hypoechoic mass with a central cystic portion (Figure 1A). Another poorly defined oval complex echoic mass was also in the same quadrant (Figure 1B). There was no axillary lymphadenopathy, but there was a well-defined cystic mass in the left axillary palpable portion mimicking dilated ducts in accessory breast tissue (Figure 2). Poorly defined masses located in the left upper outer quadrant were sonographically suspicious for breast cancer, and sonographically guided core biopsies were performed for each mass. The pathologic diagnosis was lymphangiomas. Surgical excision was performed after sonographically guided localization of the breast lesions. The axillary mass was not excised, and it was presumed to be recurrent lymphangiomas after the previous surgery.
Gross pathologic examinations revealed poorly defined spiculated whitish gray masses. Microscopically, they were composed of numerous cavernous and cystically dilated spaces lined by endothelial cells supported by a prominent fibrocollagenous stroma (Figure 3). The final histologic diagnosis was cavernous lymphangiomas of the breast.

**Discussion**

Lymphangiomas are relatively uncommon lesions, more so in the breasts, where they are very rare. They occur predominantly in children, with up to 90% of cases presenting by the second year of life.\(^3\) Lymphangiomas are vascular malformations rather than true neoplasms, and they are produced by sequestrated primitive lymphatic tissue that fails to communicate with peripheral drainage pathways.\(^2\)\(^-\)\(^4\) Lymph vessels of the adult mammary gland originate in the interlobular connective tissue and the walls of lactiferous ducts. They communicate with the overlying cutaneous lymphatic plexus around the nipple in the subareolar plexus and then drain to the axilla.\(^3\) Cases of lymphangiomas in the breast that have been reported in the literature have mainly been located in the upper outer quadrant of the breast.\(^6\)\(^-\)\(^8\) In our case, the same distribution pattern was seen. This result may be related to the drainage pattern of the breast lymphatic system.

The mammographic findings are nonspecific. Because of the frequent location high in the axilla, it may not be possible to image these lesions. Additionally, many occur in infants and children, in whom it is desirable to avoid radiation if possible. Cystic lymphangiomas show round or lobulated densities on mammography.\(^9\)\(^-\)\(^11\) When the lymphatic channels are prominent, the sonographic appearance can be similar to that of duct ectasia that occurs within accessory breast tissue within the axilla. The lymphatic vessels in a lymphangioma can be completely collapsed with transducer pressure, whereas the ectatic ducts within accessory breast tissue cannot.\(^12\)
Usually, lymphangiomas are described as fairly well-defined multicystic masses. In our case, however, they appeared as complex echoic spiculated masses mimicking breast cancer. The echogenicity of lymphangiomas can change from the size of the lymphatic channels. When the lymphatic channels are small, the echogenic vessel walls are close together, making numerous interfaces, and the overall echogenicity resembles that of solid masses. In our case, the lymphatic channels were too small to make a well-defined cystic mass. In addition, extensive fibrosis with a prominent fibrocollagenous stroma in the periphery made the masses spiculated and hypoechoic.

Magnetic resonance imaging is the imaging modality of choice for diagnosis and evaluating the extent of disease. On magnetic resonance imaging, cystic lymphangiomas are typically seen as septated masses of low T1- and high T2-weighted signal intensity, with only septal enhancement. The signal intensity of cyst contents can be variable because of hemorrhage or proteinaceous fluid.13

Primary treatment of lymphangiomas is surgical excision. However, there may be difficulty in obtaining safe margins because of the tendency of these lesions to infiltrate surrounding tissues. The probability of recurrence is low if the lesions are completely excised.14 Sometimes, percutaneous sclerosis or a combination of percutaneous sclerosis and surgical excision is performed.

This report describes an unusual case of cavernous lymphangiomas of the breast presenting as suspicious malignant masses from an extensive fibrocollagenous stroma and small lymphatic channels. Although the incidence of this disease is extremely rare in the adult breast, a breast lymphangioma should be considered when an irregular complex echoic cystic mass is located in the upper outer quadrant of the breast. Wide surgical excision after evaluation of the disease extent is the most effective treatment modality.

References

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