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Initial presentation and recurrence of metastatic rhabdomyosarcoma as breast mass

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Rhabdomyosarcoma rarely metastasizes to the breast. We report a case of a pediatric patient who initially presented with a right breast mass and pancytopenia, which was subsequently diagnosed as alveolar rhabdomyosarcoma. Despite initial favorable response to chemotherapy, a new metastatic focus was found in the contralateral breast 10 months later.

Introduction

In the pediatric and adolescent population, the overwhelming majority of breast masses are due to benign etiologies. When malignant, they tend to be metastatic disease from lymphoma, leukemia, and rhabdomyosarcoma. In both primary and metastatic cases of rhabdomyosarcoma to the breast, the histologic subtype is most commonly alveolar. Our case of metastatic alveolar rhabdomyosarcoma initially presented as a right breast mass and pancytopenia. This is an extremely rare presentation of rhabdomyosarcoma. Following an initial favorable response to chemotherapy, a new left breast metastasis was found on repeat PET/CT 10 months after diagnosis.

Case report

A 15-year-old female presented to urgent care with diffuse swelling of the right breast. She initially palpated a lump in her breast one week before presentation. The lump then quickly enlarged and became more firm, such that the right breast was noticeably larger than the left breast. The patient also had symptoms of an upper respiratory tract

infection and two episodes of epistaxis. Physical exam was positive for a large firm mass involving most of the outer

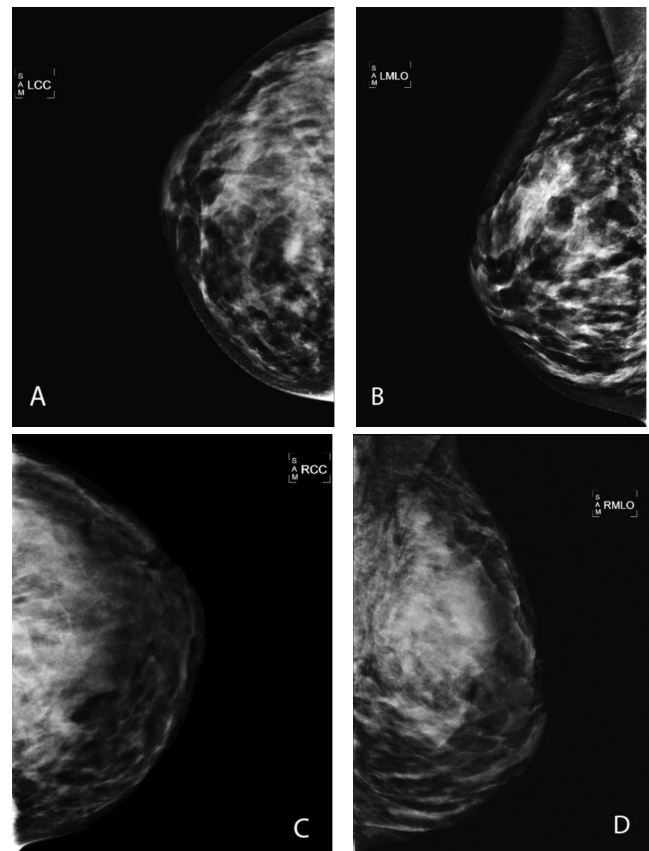


Figure 1. 15-year-old female with metastatic rhabdomyosarcoma. Bilateral mammogram demonstrates diffusely increased density within the right breast.

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right breast. In addition, laboratory analysis was remarkable for pancytopenia (WBC 4, Hgb 10, Plt 56). A subsequent mammogram showed a unilateral dense right breast (Fig. 1). Breast ultrasound revealed a 5cm hypoechoic mass at 10 o'clock (not shown). The patient was started on Keflex for possible mastitis.

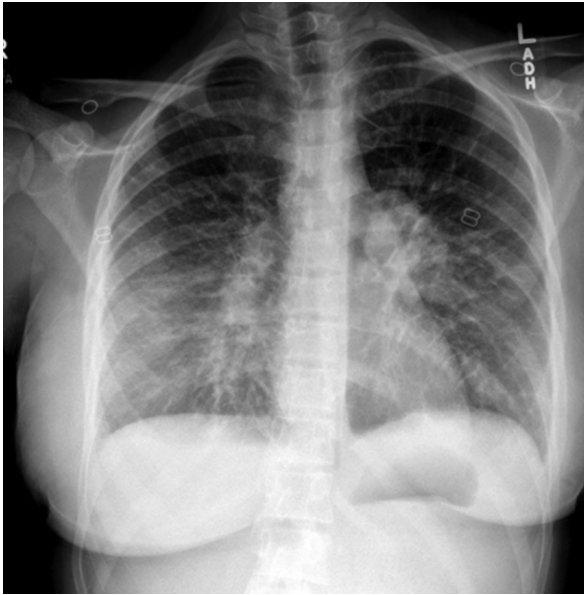


Figure 2. 15-year-old female with metastatic rhabdomyosarcoma. Frontal chest radiograph shows multiple findings including asymmetrically enlarged right breast, mediastinal lymphadenopathy, small bilateral pleural effusions, and pulmonary opacities.

One week later, the patient did not notice any improvement in her right breast swelling. Repeat laboratory evaluation confirmed persistent pancytopenia. Due to concern for malignancy, she was admitted to the hospital and referred to surgical oncology. The working diagnosis was acute myelogenous leukemia with granulocytic sarcoma of the breast. Rhabdomyosarcoma was considered but was felt to be unlikely due to the presence of pancytopenia. Bone marrow biopsy was nonspecific; immunophenotyping for myeloid leukemia was negative. The patient then underwent a palpation-guided core biopsy of the breast mass. Pathology was consistent with alveolar rhabdomyosarcoma with a FOXO1 (FKHR) gene rearrangement on chromosome 13q14.

On further questioning, the patient reported several weeks of right foot swelling and tenderness following minimal trauma. Subsequent foot radiographs demonstrated a healing second metatarsal fracture. She also complained of developing pain in her right thigh and knee. A chest radiograph was remarkable for intrathoracic lymphadenopathy and diffuse pulmonary opacities (Fig. 2). A PET scan revealed diffuse infiltration of the right breast with multiple hypermetabolic masses. Moreover, there was diffuse bone-marrow activity plus innumerable hypermetabolic pulmonary opacities, pathologically enlarged lymph nodes

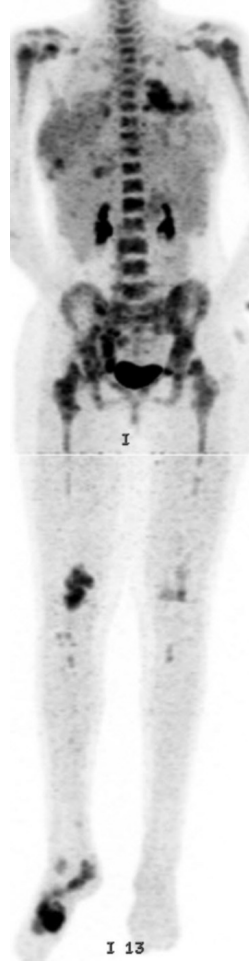


Figure 3. 15-year-old female with metastatic rhabdomyosarcoma. Maximum intensity projection (MIP) image reveals extensive hypermetabolic activity within the right breast, lungs, axial and proximal appendicular skeleton, right foot, and lymph nodes of the chest, abdomen, pelvis, right inguinal region, and right popliteal fossa.

throughout the chest/abdomen/pelvis, a large right popliteal fossa mass, and multiple soft-tissue masses in the right foot. The largest and most hypermetabolic of these soft-tissue masses was located along the plantar surface at the level of the metatarsals (Figs. 3-4). The patient was diagnosed with stage 4 rhabdomyosarcoma, with the largest right foot mass presumed to be the primary tumor site.

The patient was started on chemotherapy with etoposide, ifosfamide, and mesna per COG protocol ARST08P1 Arm P2. She also received radiation therapy to the right foot, popliteal fossa, and hemipelvis. Ten months later, a PET/CT performed after 13 cycles of chemotherapy showed decreased

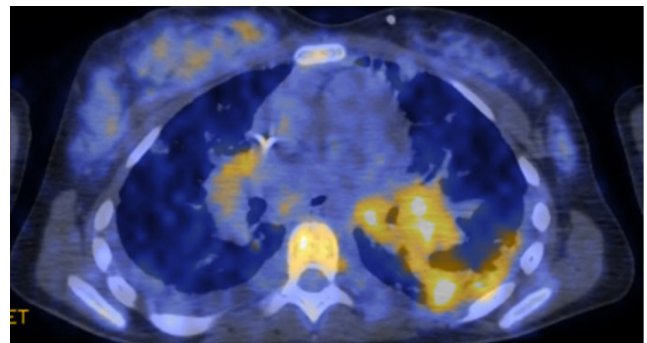


Figure 4. 15-year-old female with metastatic rhabdomyosarcoma. Fusion image of the chest better details hypermetabolic activity in a diffusely enlarged and dense right breast, extensive pulmonary involvement, and intrathoracic lymphadenopathy.

activity in the previous areas of involvement. However, a new hypermetabolic focus in the lateral left breast was identified (Fig. 5). Ultrasound confirmed the presence of a 2.2cm left axillary tail mass with suspicious features (Fig. 6). Histopathology from subsequent core-needle biopsy was

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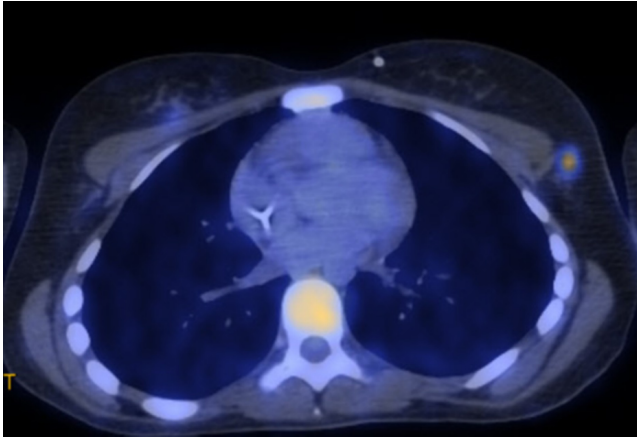


Figure 5. 15-year-old female with metastatic rhabdomyosarcoma. Fusion image from a subsequent PET/CT shows a new hypermetabolic focus in the far left lateral breast.

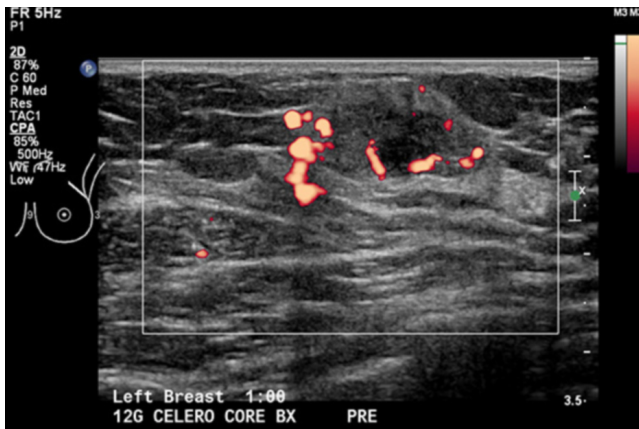


Figure 6. 15-year-old female with metastatic rhabdomyosarcoma. Correlative ultrasound demonstrates a mass with heterogeneous echogenicity in the left axillary tail with irregular margins and internal vascularity, suspicious for a new metastasis.

consistent with metastatic alveolar rhabdomyosarcoma (Figs. 7-8). The chemotherapy regimen was then altered to vincristine, irinotecan, and temozolomide. The left breast mass initially decreased in size. Unfortunately, the response was transient; followup ultrasound a month later revealed that it had grown to 4 cm. One week after the ultrasound study, the patient presented to the emergency department with dizziness and hypoesthesia on the right side of her body. MRI of the brain showed multiple intra-axial hemorrhagic metastases, the largest of which was in the left inferior parietal lobule (Fig. 9).

Discussion

In the pediatric population, breast malignancies are quite rare. The etiologies of breast enlargement and masses are most often benign. These include hormonal, infectious, or traumatic etiologies. In addition, benign tumors such as fibroadenomas outnumber malignancies. Unlike in adult

women, primary breast carcinoma in children and adolescents is exceedingly rare, with phyllodes tumor being the most common primary malignancy. Most malignancies are metastases, commonly from neuroblastoma, rhabdomyosarcoma, or hematologic malignancies (1).

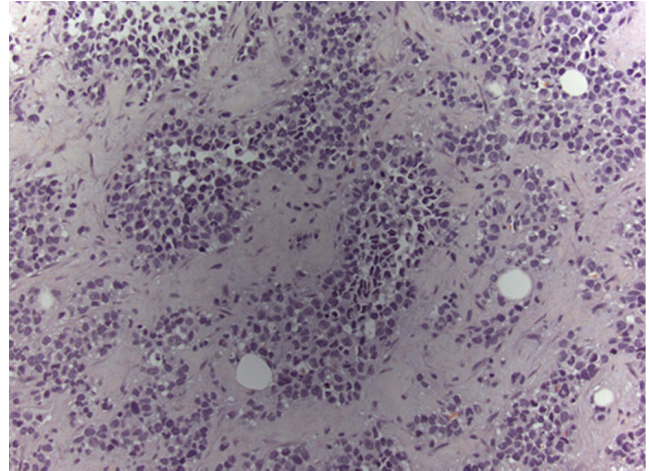


Figure 7. 15-year-old female with metastatic rhabdomyosarcoma. The core biopsy demonstrated sheets of small round blue cells, separated by a framework of dense fibrous septa, with formation of irregular "alveolar" spaces and central loss of cellular cohesion. This is consistent with a poorly differentiated small blue-round-cell tumor.

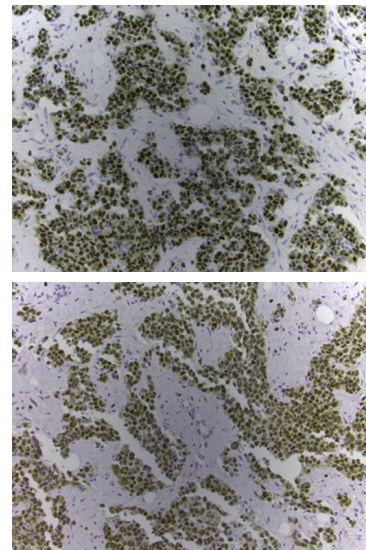


Figure 8. 15-year-old female with metastatic rhabdomyosarcoma. The tumor cells stained positive for myogenic markers, including myogenin (above) and myoD1 (below). They were negative for pancytokeratin, CD45, and S100 (not shown). The cytogenetic study (not shown) revealed a FOXO1 (FKHR) gene rearrangement on chromosome 13q14. These findings are consistent with alveolar rhabdomyosarcoma.

The most common soft-tissue sarcoma of childhood, rhabdomyosarcoma accounts for 5% of all childhood malignancies (2). These tumors arise from primitive muscle cells that fail to differentiate into normal skeletal muscle; they also occur in organs that do not contain skeletal muscle. Most cases arise from the head and neck, genitourinary organs, and extremities. The main subtypes of rhabdomyosarcoma are embryonal (with botryoid and spindle-cell variants) and alveolar. Other less common subtypes include

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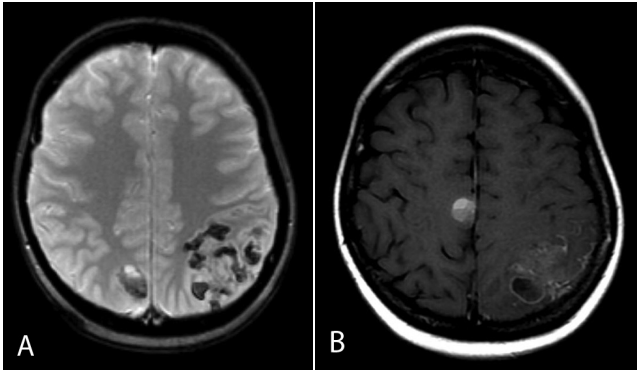


Figure 9. 15-year-old female with metastatic rhabdomyosarcoma. Gradient echo (A) demonstrates intracranial hemorrhagic metastases with fluid-fluid levels. Post-contrast (B) image shows leptomeningeal enhancement surrounding the mass in the left inferior parietal lobule.

pleomorphic (which are uncommon in children) and anaplastic (3). Embryonal rhabdomyosarcoma constitutes 60-70% of all rhabdomyosarcomas and often arises in mucosal-lined tissues of the head and neck, genitourinary organs, and gastrointestinal tracts (2). This subtype derives its name from its resemblance to developing muscle cells (hypercellular zones demonstrating myogenesis mixed with hypocellular zones that contain primitive mesenchyme) (3). In contrast, the alveolar subtype, which makes up 20 % of rhabdomyosarcoma, tends to occur in the soft tissues of the trunk and extremities (2). These tumors produce small round blue cells that are separated by fibrovascular septa. The central aggregates of tumor cells appear to be contained within “alveolar” spaces. The discovery of a PAX3-FKHR fusion gene (PAX3 gene on chromosome 2q35 and FKHR/FOXO1 gene on chromosome 13q14) allows definitive differentiation of alveolar rhabdomyosarcoma from other small round-blue-cell tumors. In terms of survival, the alveolar and pleomorphic subtypes have poorer prognosis than the embryonal subtype. The botryoid and spindle-cell variants of the embryonal subtype have the best prognosis among rhabdomyosarcomas (3).

Metastases to the breast are seen in 6% of patients with rhabdomyosarcoma (1). These are most commonly of the alveolar subtype. In one analysis that examined 19 cases of rhabdomyosarcoma with initial metastases to the breasts, all of those (18) in which the histology could be accurately determined were of the alveolar subtype. Both primary and metastatic breast rhabdomyosarcomas have been reported to most commonly occur in adolescent females or young adults, with an age range of 11.5 to 20.2 years and a median of 15.2 years (4). However, since 1997, other cases have been reported in adult women. More recently, Li et al reported five patients with either primary or secondary breast rhabdomyosarcomas whose mean age was 30 years (5).

Clinically, breast metastases often present as rapidly enlarging masses that may be tender. Their radiologic appearances are nonspecific. Although they can be bilateral

and multiple, breast metastases more commonly initially present as a single mass. Mammographically, they can also (as in the case of our patient) manifest as diffusely increased density (1). The combination of pancytopenia and breast mass, seen with our patient, is an exceedingly rare presentation of rhabdomyosarcoma (6).

Due to the radiosensitivity of developing fibroglandular tissue in pediatric patients and their dense breasts, ultrasound is preferred over mammography as the initial imaging modality in this population. The sonographic appearance of a metastatic lesion is most commonly a solid mass with variable echogenicity. As in adults, breast masses in the pediatric population require thorough workup. Although the appearances of benign and malignant masses may overlap, those with suspicious features (for example, irregular margins, posterior shadowing, antiparallel orientation) warrant further evaluation with tissue sampling (7). However, the lack of a discrete mass on ultrasound does not negate biopsy, as metastatic rhabdomyosarcoma can sonographically mimic normal fibroglandular tissue (8).

To the best of our knowledge, there is a paucity of long-term survival data in the literature for rhabdomyosarcoma with initial metastases to the breasts. Hays et al reported a 5-year Kaplan-Meier estimated survival rate of 35% in patients with breast metastases from rhabdomyosarcoma. Nevertheless, the same study reported two patients with no evidence of active disease for more than 15 years (4).

Our case emphasizes the importance of recognizing that, although rare, malignancies do occur in the pediatric and adolescent breasts. In addition, metastatic disease is significantly more common than primary breast malignancies. In these young patients, especially those with known primary malignancies such as rhabdomyosarcoma or lymphoma/leukemia, a new palpable breast mass should be viewed with suspicion. Although benign entities occur with much greater frequency in children than adult females, a breast mass should warrant careful workup in all patients.

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