Li-Fraumeni Syndrome: Adopting a Diagnosis with an Unknown Family History

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Abstract:

Introduction: Li-Fraumeni Syndrome (LFS) is a rare, autosomal dominant disease that is typically screened for and identified in patients with a known history of cancer and family history of cancer.

Presentation of Case: Ms. DA is an adopted 39-year-old South American female with a past medical history of iron deficiency anemia, celiac disease, and polio. She presented with ductal carcinoma in situ with spindle cell atypia at age 35 and leiomyosarcoma of the ankle at age 36. Genetic counseling was offered due to her age and cancer types, with subsequent genetic testing revealing a TP53-R337H mutation, which was diagnostic of LFS. Given her high risk status, extensive cancer screenings at common LFS body sites were recommended, but she was lost to follow-up. Four years later, she returned after discovering a new right breast mass on self-examination. She was referred to a new breast surgeon, who was able to establish consistent and close follow-up while providing counseling and education regarding lifetime risk and cancer trajectory of classical LFS compared to the Brazilian LFS-subtype.

Conclusion: The importance of having a high level of suspicion for patients with an unknown family history is crucial as seen in this case. Without a biological family history, clinicians rely on their judgement to decide when further work-up is warranted. Advancements in medicine and genetic testing have increased the ability to accurately diagnose genetic diseases to help patients make life-saving decisions about their health.

Keywords: Li-Fraumeni Syndrome; LFS; breast cancer; screening; adoption

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Consent: Consent was taken from the patient’s next of kin for publication of this case report.

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Introduction

Li-Fraumeni Syndrome (LFS) is an inherited autosomal dominant malignancy that is characterized by the development of multi-system cancers due to a deficiency in the cell cycle regulator, TP53 (Figure 1)[1]. Also known as Sarcoma-Breast-Leukemia-Adrenal Gland (SBLA) Cancer Syndrome, approximately 400 families in the United States have been described in literature with this disease since its initial characterization in 1969 [1, 2]. The overall prevalence worldwide is less known, with an estimate of over 1000 multigenerational families across 172 countries. Family history plays a significant role in risk stratification and screening protocols since definitive diagnosis requires physicians to identify the risk and order DNA sequencing to identify TP53 mutations or deletions [3]. In presence of young age and absence of family history, providers need to have an even higher index of suspicion in caring for patients with possible multi-system cancers. Here we present an adopted patient with no known biological family history with a classical LFS presentation.

Figure 1 LFS and its common cancer associations
Case presentation

Ms. DA is a 39-year-old South American female with a past medical history of iron deficiency anemia, celiac disease, and polio. She was adopted as a child from Brazil and was unaware of her biological family’s medical history. She initially presented at age 35 for an early screening mammogram, which showed a well-circumscribed density in the central right breast. An image-guided core biopsy revealed ductal carcinoma in situ with spindle cell atypia, with subsequent right breast lumpectomy and whole breast radiation treatment. One year later, she returned with a new left ankle mass. Excisional biopsy and resection revealed a leiomyosarcoma with clear margins noted, with radiotherapy afterwards.

Genetic counseling was offered due to her age and cancer types, with subsequent genetic testing revealing a TP53-R337H mutation, which was diagnostic of LFS. Given her high risk status, extensive cancer screenings at common LFS body sites were recommended, but she was lost to follow-up.

Four years later, she returned after discovering a new right breast mass on self-examination. Mammogram and ultrasound confirmed a benign hematoma, likely due to trauma. She was referred to a new breast surgeon, who was able to establish consistent and close follow-up while providing counseling and education regarding lifetime risk and cancer trajectory of classical LFS compared to the Brazilian LFS-subtype. After extensive discussion, the patient opted for prophylactic bilateral mastectomy and reconstruction due to concerns of prior radiotherapy exposure to her breast. Since then, she had no new complaints and is compliant with standard LFS cancer screening schedules.

Discussion

LFS carries a primary cancer risk of up to 100%, by the age of 60 in women, and a risk of up to 73% in men [4]. Furthermore, these patients have a high risk of developing a secondary and a tertiary malignancy of about 30-57% and 38%, respectively [5, 6]. Early detection and referral for genetic screening becomes complicated in situations such as Ms. DA as LFS screening criteria consistently utilize family history as a defining feature (Figure 2)[7]. Other protocols with and without family history have also been used, with varying success rates [8-11]. Despite having an unknown family history and only fulfilling the personal history of cancer criteria, her presentation relative to her age triggered a high index of clinical suspicion and warranted further work-up.

<table>
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<tr>
<th>Criteria for TP53/Molecular Testing for LFS</th>
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<tr>
<td><strong>Classical</strong></td>
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<tr>
<td>1. Proband with sarcoma diagnosed before age 45 AND</td>
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<tr>
<td>2. First-degree relative with any cancer before age 45 AND</td>
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<tr>
<td>3. First/Second-degree relative with any cancer before age 45/sarcoma at any age</td>
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<tr>
<td><strong>Chompret</strong></td>
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<tr>
<td>1. Classical criteria, but before age 46</td>
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<tr>
<td>2. Proband with multiple tumors (except breast), 2 of LFS spectrum, 1 before 46</td>
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<tr>
<td>3. Proband with adrenocortical or choroid plexus tumor, regardless of family history</td>
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<tr>
<td><strong>Birch</strong></td>
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<tr>
<td>Similar to classical, however proband can also be diagnosed with brain or adrenocortical tumor in addition to sarcoma</td>
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<tr>
<td><strong>Eele</strong></td>
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<tr>
<td>Family history only: 2 first- or second-degree relatives with LFS spectrum malignancies, diagnosed at any age</td>
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Figure 2 Various criteria for assessing need for genetic testing in definitive diagnosis of LFS.
Subsequent cancer screenings with individuals with LFS include: annual physical, clinical, self-breast, and dermatological examinations, whole body MRI screening for secondary cancers, colorectal screening and risk-reducing bilateral mastectomy [12, 13]. High recurrences of secondary and tertiary cancers rates in LFS require patients to undergo strict surveillance studies. Of note, compared to the high lifetime primary cancer risk of most individuals/families with LFS, the Brazilian population’s specific R337H founder mutation has only a 60% lifetime risk with improved survival rates post-treatment[14].

The differential diagnosis for LFS can include BRCA1/BRCA2 mutations for breast cancer, and the family of DNA repair enzymes of MLH1, MSH2, MSH6, PMS1, and PMS2 in mismatch repair cancer (Lynch) syndrome[12, 13]. Advances in genetic screening and testing, however, has eliminated the uncertainty of these inherited cancer syndromes through accurate diagnosis. Despite its rarity and lack of supporting family history, Ms. DA was able to make potentially lifesaving decisions due to the clinical judgement of her physicians. Further studies are required to assess the utility of genetic testing in adoption cases where family history is unavailable.

Conclusion

Li Fraumeni Syndrome is a rare, autosomal dominant disease that is typically screened for and identified in patients with a known history of cancer and family history of cancer. As seen in this case, the importance of suspicion and clinical judgement for patients with an unknown family history is crucial. In the setting of global migration, physicians need to have high clinical suspicion when confronted with abnormal age to cancer presentations. Advancements in medicine and testing have increased our ability to accurately diagnose genetic diseases to help patients make life-saving decisions about their health and healthcare.

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Competing interest and funding statement

Jia Hong Chen, David I. Kaufman MD, Justin Chin, and Christine Lomiguen MD report that there are no competing interests or financial disclosures for this study.

Consent

Patient has given their informed consent for this case report to be published.

References