Case Report

Recurrence Leiomyosarcoma of the Popliteal Vein: A Rare Soft Tissue Sarcoma

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1. Introduction

Sarcoma is a malignant tumor that arises from mesenchymal cells and constitutes approximately 1% of all malignancies in adults. About 510% of all soft tissue sarcomas are leiomyosarcoma (LMS). LMS is a soft tissue sarcoma that originates from smooth muscle cells, and it can be divided into three subtypes according to the site of presentation: soft tissue LMS, cutaneous LMS, and vascular LMS [1]. Vascular LMS is the least common subtype of LMS. The first case of vascular LMS was reported by Perl in 1871, and the tumor originated from the inferior vena cava [2–4]. About one-third of vascular LMS is located in the extremities, most commonly in the saphenous vein (25%) [1]. Vascular LMS originating from the popliteal vein is very rare, and to the best of our knowledge, only nine cases have been reported to date.

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close or positive [1, 7]. The role of chemotherapy for systemic control remains unclear.

We herein report a case of popliteal LMS and present a literature review of previously reported cases. Although the frequency of this type of tumor is low, it should be considered in a patient who presents with a mass at the popliteal fossa and intermittent claudication.

2. Case Presentation

A 49-year-old woman presented with a soft tissue mass at the posterior aspect of the right proximal leg in 2019. Tumor removal was performed by the general physician without tissue pathologic examination or evaluation of the surgical margin. One year later, she returned with a recurrent mass at the same site of the previous surgery. The mass had exhibited slow growth. She had mild pain and intermittent claudication without a history of an edematous leg. Physical examination revealed an ill-defined mass that was located at the posterior aspect of the right proximal leg and extended to the popliteal fossa. It was $10 \times 10$ cm in size, movable, and mildly tender (Figures 1(a) and 1(b)).

Plain radiographs revealed a soft tissue mass on the posterior aspect of the right proximal leg immediately distal to the popliteal fossa without any bone lesions. Magnetic resonance imaging (MRI) showed a deep-seated heterogeneous soft tissue mass originating from the popliteal vein (Figures 2(a), 2(b), and 2(c)).

Chest computed tomography (CT) for staging revealed no pulmonary metastasis. A core needle biopsy was performed. The pathological report indicated a malignant soft tissue tumor that was immunohistochemically positive for smooth muscle actin, epithelial membrane antigen, vimentin, caldesmon, and desmin, compatible with LMS (Figures 3(a), 3(b), and 3(c)).

The patient underwent wide en bloc resection of the tumor, including the segment of the involved popliteal vein, without venous reconstruction (Figures 4(a), 4(b), 4(c), 4(d), and 4(e)).

The surgical wound was covered by a split-thickness skin graft. The final pathology report from the surgical specimen revealed a typical intersecting 90-degree angle fascicular arrangement of spindle cells as well as scattered pleomorphic hyperchromatic cells with typical cigar-shaped nuclei and eosinophilic cytoplasm, compatible with LMS (Figures 5(a), 5(b), 5(c), and 5(d)).

All resected margins were free of tumor tissue (Figures 6(a), 6(b), 6(c), 6(d), 6(e), and 6(f)).

The patient received no other adjuvant treatments. The postoperative course was uneventful, and the patient was discharged within 1 week of surgery. No complications, adverse outcomes, local recurrence, or distant metastasis was reported at 16 months after surgery. The patient had good functional outcomes (Figures 7(a) and 7(b)).

3. Discussion

Vascular LMS is the least common subtype of LMS and originates from the tunica media of the vessel wall [1]. The popliteal vein is a very uncommon location of vascular LMS, and
to the best of our knowledge, only nine cases have been reported to date [2–7]. The clinical presentation of vascular LMS of the popliteal vein includes a mass at the popliteal fossa, intermittent claudication, and pain with varying degrees of intensity. Some patients’ presentation may mimic deep vein thrombosis (DVT), but patients with DVT usually have risk factors for DVT [3]. In an otherwise healthy patient who presents with intermittent claudication and leg edema without risk factors for DVT, a vascular tumor or tumor that compresses and obstructs the venous circulation should be considered, although such tumors are very rare [3, 4, 8].

Figure 2: Magnetic resonance imaging. (a) Sagittal view with gadolinium contrast. (b) Axial view with gadolinium contrast. (c) Axial T2-weighted image.
The mass can grow rapidly and involve adjacent structures such as bone, nerves, and other vessels. Affected patients usually have a history of leg edema because of the obstruction of venous return, but some patients have no history of an edematous leg because of the development of collateral circulation, especially patients with a slowly growing mass. Basu et al. [6] reported a case of a 35-year-old woman with LMS of the left popliteal vein without leg edema. The tumor was resected without vascular reconstruction. Killoran et al. [3] and Karamoshos et al. [4] reported cases of LMS of the popliteal vein in patients with a history of leg edema. En bloc resection of the popliteal vein tumor with vascular reconstruction was performed in both cases. In our case, the patient presented with recurrence of a slowly growing mass at the right popliteal fossa and no history of leg edema. She underwent wide en bloc resection of the tumor, including the segment of the involved popliteal vein, without venous reconstruction. She had good oncologic and functional outcomes without complications.

MRI is the imaging technique of choice to identify tumor characteristics and determine the tumor size, extension, and anatomic relationships [3, 9]. MRI is also very useful for preoperative planning. However, MRI is not useful in differentiating benign from malignant lesions. CT angiography (CTA) is also helpful to identify intraluminal tumors and their collateral circulation for preoperative planning [3, 5]. In patients with adequate collateral circulation, wide resection without vascular reconstruction is suitable. In our case, we did not use CTA for preoperative planning because it was not available at that time. We found that the popliteal vein was totally occluded by the tumor intraoperatively, but the patient had no history of leg edema. Therefore, we considered that she already had an adequate collateral circulation. We considered performing wide en bloc resection without vascular reconstruction. Basu et al. [6] also reported a case of a 35-year-old woman with LMS of the left popliteal vein without leg edema, and the investigation did not include CTA. The patient underwent wide resection without vascular reconstruction.

The definitive diagnosis of LMS is achieved by histopathology and immunohistochemistry. The histopathological study of vascular LMS is similar to that of LMS at any location. The findings are characterized by pleomorphic hyperchromatic spindle-shaped cells with typical cigar-shaped nuclei and eosinophilic cytoplasm [1, 10]. Immunohistochemical studies are very helpful for a definitive diagnosis; they are usually positive for smooth muscle actin, vimentin, desmin, calponin, and smooth muscle myosin heavy chains and negative for S-100, inhibin alpha, and CD117 [1].

The mainstay of treatment is wide en bloc resection of the tumor, including the involved segment [1, 4, 7]. Venous reconstruction after resection is not necessary in patients who have no history of leg edema and in whom CTA reveals collateral circulation [6]. Radiotherapy is an important

Figure 3: There is positive immunoreactivity for (a) desmin, (b) smooth muscle actin, and (c) caldesmon, which supports the diagnosis of leiomyosarcoma.
Figure 4: Tumor removal. (a) Incision. (b) Tumor after en bloc excision (superficial view). (c) Tumor after en bloc excision (deep margin). (d) Popliteal vein involved by tumor. (e) Surgical wound covered by split-thickness skin graft.
treatment for local control when surgical margins are positive or close [1, 7]. The efficacy of chemotherapy for systemic control is still unclear. In our case, we performed wide en bloc resection of the tumor, including the segment of the involved popliteal vein, without venous reconstruction. The patient did not receive radiotherapy or chemotherapy because the surgical margin was negative and there was no evidence of metastasis.

The 5-year survival rate of vascular LMS is about 32% [3, 8], and approximately 10% of patients present with lung or liver metastasis at the time of diagnosis because the vascular tumors have direct access to the blood circulation [3, 11]. Vascular LMS generally has a poor prognosis.

We have summarized the clinical data of nine patients with LMS of the popliteal vein from our literature review in Table 1.

4. Conclusion
Vascular LMS at the popliteal vein is uncommon but should be considered as a differential diagnosis in a patient who presents with a mass at the popliteal fossa. The mainstay of treatment is wide en bloc resection of the tumor, including the involved segment of the vein. Venous reconstruction after resection is not necessary in chronic cases without a history of an edematous leg.
Figure 6: (a) Low power magnification of leiomyosarcoma reveals typically intersecting 90-degree angle fascicular arrangement of spindle cells. (b) High magnification area shows scattered pleomorphic hyperchromatic cells with typically "cigar-shaped" nuclei (arrowhead) and eosinophilic cytoplasm. (c) and (d) Reveals increase mitotic activity with atypical forms. (e) and (f) Sections from the main vessel show the neoplasm derived from the tunica media of the vessel (arrowhead) and high magnification reveals significantly cytologic atypia.
Data Availability

The datasets used or analyzed during the current study are available from the corresponding author upon reasonable request.

Ethical Approval

This retrospective chart review study involving human participants was in accordance with the ethical standards of the institutional and national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards. The Human Research Ethics Committee of the Khon Kaen Hospital acknowledged and approved this study (Approval Number: KEXP65030).

Consent

The patient acknowledged and signed informed consent regarding publishing her data and photographs.

Conflicts of Interest

The authors declare that they have no competing interests.

Authors’ Contributions

TP: Conceptualization, methodology, validation, formal analysis, investigation, data curation, writing—original draft, visualization, supervision, project administration. SS: Validation,
formal analysis, investigation, writing—original draft, supervision, project administration. KS and CS: Methodology, validation, formal analysis, writing—review and editing. TC: Data curation, visualization, writing—review and editing.

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