Congenital cutaneous pedicled macrocystic lymphatic malformation on the upper extremity: A rare case report and review of the literature

INTRODUCTION

Lymphatic malformations are one of several categories of vascular tumor [1]. Tumors of lymphatic origin can exist in multiple forms. In particular, a lymphatic malformation can be microcystic, macrocystic, or mixed. Among macrocystic lymphatic malformations, cutaneous lymphatic malformations on the extremities are extremely rare. These cutaneous malformations more commonly occur in the neck, axilla, or groin area [2,3].

A lymphatic malformation may cause symptoms such as infection, bleeding, and rupture, as well as cosmetic problems [4]. Therefore, physicians who deal with lymphatic malformations should actively consider curative options. Consensus on the optimal treatment of these tumors is lacking, but many treatment methods have been attempted, such as observation, electrocauterization, aspiration, excision, and cryotherapy. Intralesional injections (sclerosing therapy), radiation, laser exposure, and ligation have also been tried [5].

Recently, we experienced a rare case of a congenital cutaneous pedicled macrocystic lymphatic malformation that was eliminated by electrocauterization.

CASE REPORT

A 4-day-old female infant with a congenital cutaneous mass on the dorsal area of her left first metacarpophalangeal joint presented to our outpatient clinic (Figs. 1, 2). Upon presentation, bleeding was noted in the area of the pedicle, but no rupture was observed. No additional lesions were identified near the mass or in surrounding areas. Clinicians considered performing additional imaging studies, including magnetic resonance imaging; however, the guardians did not provide permission for the patient to be sedated, considering that she was only 4 days old. The stalk of the mass measured...
less than 2 mm and had no palpable inner tissue. Due to bleeding around the pedicle, the procedure was planned promptly to avoid rupture. We excised the mass without rupture, using an electrocautery device to cut the pedicle gently with minimal bleeding. The resected mass was measured as 2.8 cm × 1.9 cm and was determined by a pathologist to be a lymphangioma (Figs. 3, 4).

A simple wet dressing was applied for 1 week, and the wound subsequently healed completely. No remnants of soft tissue bulged out from the lesion where the pedicle was previously, and no hump-like deformity was found. The patient presented for a follow-up appointment 3 and a half months later, and no recurrence of the deformity or lingering wound was noted at the surgical site (Fig. 5).

**DISCUSSION**

Tumors of lymphatic origin can be divided into microcystic, macrocystic, and mixed lymphatic malformations. Microcystic lym-
phatic malformations are limited to the dermal region and tend to extend across a wider area. Mixed lymphatic malformations are located more deeply and are associated with diffuse swelling. These usually develop on the tongue, lips, and neck. Macrocystic lymphatic malformations also have multiform characteristics and usually appear as a bulge under the skin; these are more easily detectable than mixed lymphatic malformations [5]. Among macrocystic lymphatic malformations, cutaneous lymphatic malformations on the extremities are extremely rare. These cutaneous malformations more commonly occur in the neck, axilla, or groin area [2].

No consensus exists on the optimal treatment of cutaneous lymphatic malformations. For this reason, Ha et al. [5] concluded that treatment should be individualized according to the size and location of the tumor and the complications that might result from treatment. Flanagan and Helwig [6] came to a similar conclusion after treating 158 cases of cutaneous lymphatic malformations, in which they focused more on surgical excision. Primary surgical excision was sufficient to treat 75% of their patients. However, none of the 158 cases of cutaneous lymphatic malformations presented a pedicled cystic formation. The surgical approach in the case presented here was easier than most of the other cases reported due to its narrow stalk.

Cutaneous lymphatic malformations are easy to diagnose and cure. In contrast, other types of lymphatic malformations may be more difficult to identify and treat. For example, intra-abdominal lymphatic malformations are difficult to approach and are usually diagnosed radiologically because they are asymptomatic [7]. Furthermore, surgical treatment of patients with lymphatic malformations hidden inside the body may have higher morbidity than treatment of patients with cutaneous lymphatic malformations. For these reasons, patients with cutaneous lymphatic malformations are relatively fortunate, because aggressive treatment is possible with low morbidity.

A few reports of macrocystic lymphatic malformations exist in the literature. Flanagan and Helwig [6] reported a case of bulging on the ring and little finger of the right hand, but it was less superficial than the case described in this report. They also presented a case of a superficial lymphatic malformation that was excised not on the hand, but on the right shin [6]. Patoulas et al. [3] reported rare cystic lymphatic malformations on the inguinal and scrotal areas, but not on the extremities.

The etiology of lymphatic malformations is unclear. However, it is certain that lymphatic malformations consist of an accumulation of fluid resulting from an abnormal drainage pattern of lymphatic vessels [8]. In the head and neck region, the incidence rate of this condition is roughly 1 in 2,000 to 4,000 live births, but it is only rarely seen on the upper extremities as a macrocystic lesion [8]. Okazaki et al. [9] reported their experiences with 128 cases of lymphatic malformation, none of which were cutaneous cystic lymphatic malformations on the upper extremities. Kato et al. [10] surveyed 153 cases and concluded that macrocystic lymphatic malformations instead tend to occur on the head and neck area.

Diagnostic tools can be utilized to detect lymphatic malformations, such as magnetic resonance imaging, grayscale ultrasonography, or Doppler ultrasonography [8]. While no consensus currently exists on the best method of treatment of this tumor, the main option used to eliminate a malformation is surgical treatment. Ogita et al. [11] first presented intracystic injection of OK-432 (group A Streptococcus pyogenes, human origin) for sclerotherapy. Currently, molecular targeting and phototherapy are gaining increasing popularity as curative options [8]. However, for macrocystic lymphatic malformations, especially for cutaneous cases like this, surgery is still the main option. Alternatively, if the macrocystic lymphatic malformation is located more deeply, sclerosing therapy using OK-432 injection can be a good alternative.

In conclusion, this is the first report of a cutaneous macrocystic lymphatic malformation of the hand. It can manifest similarly to polydactyly, and the treatment options for these conditions are also similar. As lymphatic malformations have a high recurrence rate, surgical excision should be conducted with periodical follow-up.

NOTES

Conflict of interest
No potential conflict of interest relevant to this article was reported.

Ethical approval
The study was approved by the Institutional Review Board of Hallym Sacred Heart Hospital (IRB No. 2019-04-016-001) and performed in accordance with the principles of the Declaration of Helsinki.

Patient consent
The patient's guardian provided written informed consent for the publication and the use of the patient's images.

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REFERENCES


