Combined Lymphedema and Capillary Malformation of the Lower Extremity

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**Background:** Primary lymphedema and capillary malformation are independent vascular malformations that can cause overgrowth of the lower extremity. We report a series of patients who had both types of malformations affecting the same leg. The condition is unique but may be confused with other types of vascular malformation overgrowth conditions (eg, Klippel–Trenaunay and Parkes Weber).

**Methods:** Our Vascular Anomalies Center and Lymphedema Program databases were searched for patients with both capillary malformation and lymphedema. Diagnosis of lymphedema–capillary malformation was made by history, physical examination, and imaging studies. Because lymphedema–capillary malformation has phenotypical overlap with other conditions, only patients who had imaging confirming their diagnosis were included in the analysis. Clinical and radiological features, morbidity, and treatment were recorded.

**Results:** Eight patients (4 females and 4 males) had confirmed lymphedema–capillary malformation. Referring diagnosis was Klippel–Trenaunay syndrome (n = 4), diffuse capillary malformation with overgrowth (n = 3), or lymphatic malformation (n = 1). The condition was unilateral (n = 6) or bilateral (n = 2). Morbidity included infection (n = 6), difficulty fitting clothes (n = 6), bleeding or leaking vesicles (n = 5), leg length discrepancy (n = 4), and difficulty ambulating (n = 3). All patients were managed with compression regimens. Operative management was liposuction (n = 3), treatment of phlebectatic veins (n = 3), staged skin/subcutaneous excision (n = 1), and/or epiphysiodesis (n = 1).

**Conclusions:** Lymphedema and capillary malformation can occur together in the same extremity. Both conditions independently cause limb overgrowth primarily because of subcutaneous adipose deposition. Compression garments and suction-assisted lipectomy can improve the condition. Lymphedema–capillary malformation should not be confused with other vascular malformation overgrowth diseases that have different morbidities and treatments. (Plast Reconstr Surg Glob Open 2016;4:e618; doi: 10.1097/GOX.0000000000000487; Published online 10 February 2016.)

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The field of vascular anomalies is confusing because lesions often look similar, and incorrect terminology commonly is used. Diagnosis of a vascular anomaly is most difficult when a patient has multiple or combined lesions. The purpose of this study was to describe the rare combination of primary lymphedema associated with a capillary malformation of the lower extremity. These patients are typically labeled erroneously with a vascular...
malformation overgrowth syndrome (eg, Klippel–Trenaunay syndrome and Parkes Weber syndrome). Recognition of lymphedema–capillary malformation is important so that the patient is managed correctly.

METHODS
Our Vascular Anomalies Center and Lymphedema Program databases were queried for patients with both lower extremity primary lymphedema and capillary malformation. Diagnosis was made by history, physical examination, and imaging studies. Because lymphedema–capillary malformation has phenotypical overlap with other conditions clinically (eg, diffuse capillary malformation with overgrowth; congenital, lipomatous, overgrowth, vascular malformations, epidermal nevi and spinal/skeletal anomalies and/or scoliosis; Klippel–Trenaunay syndrome; and Parkes Weber syndrome), only patients who had imaging studies (eg, lymphoscintigraphy, magnetic resonance imaging [MRI], and computed tomography) confirming their diagnosis were included in the analysis. Clinical features, radiological findings, morbidity, and treatment were recorded.

RESULTS
Eight patients (4 females and 4 males) had confirmed lymphedema–capillary malformation (Figs. 1, 2). Nine other patients likely had the condition but did not have adequate photographs and/or imaging studies to ensure the diagnosis. The disease was unilateral in 6 patients and bilateral in 2 individuals. Patients were referred with a diagnosis of Klippel–Trenaunay syndrome (n = 4), diffuse capillary malformation with overgrowth (n = 3), or lymphatic malformation (n = 1). Morbidity included infection (n = 6), difficulty fitting clothes (n = 6), bleeding or leaking vesicles (n = 5), leg length discrepancy (n = 4), and difficulty ambulating (n = 5). All patients were managed with compression garments, and 7 individuals were prescribed pneumatic compression. Operative intervention was liposuction (n = 3), treatment of phlebectatic veins (n = 3), staged skin/subcutaneous excision (n = 1), and/or epiphysiodesis (n = 1).

DISCUSSION
Patients with vascular anomalies often are misdiagnosed and receive incorrect management.1,2
Although many types of vascular anomalies can cause lower extremity overgrowth, children frequently are labeled generically with “lymphedema” if there is no skin discoloration or with “Klippel–Trenaunay syndrome” if a cutaneous stain is present. In this study, we describe the association of primary lower extremity lymphedema in combination with a capillary malformation.

Primary lymphedema results from the anomalous development of the lymphatic system and is rare, affecting 1 in 100,000 persons; several mutations are known to cause the disease. Males typically present with bilateral lower extremity swelling in infancy; females most commonly develop unilateral leg edema in adolescence. Lymphatic fluid in the interstitial space causes the extremity to enlarge
because of subcutaneous adipose deposition. Children with the disease may be erroneously diagnosed with Klippel–Trenaunay syndrome or Parkes Weber syndrome. Patients with lymphedema can have psychosocial morbidity because of their deformity and an increased risk of infection. Primary lymphedema usually can be diagnosed by history and physical examination. The color of the integument appears normal; patients have pitting edema of the distal limb, and the skin of the dorsal foot is unable to be pinched between 2 fingers. Diagnostic confirmation of lymphedema requires lymphoscintigraphy, which is 92% sensitive and 100% specific for the disease.

Capillary malformation (previously referred to as “port-wine stain”) affects approximately 1 in 300 individuals; the lesion is caused by a somatic mutation in GNAQ. Rarely, a capillary malformation diffusely affects a large area of the lower extremity. Capillary malformation can cause tissues below the area of the skin to become overgrown, particularly subcutaneous adipose. Axial bone enlargement also can occur, resulting in a leg length discrepancy. Diagnosis of capillary malformation is made by physical examination. The stain darkens as the patient ages, and subcutaneous phlebectasia can develop as blood stagnates and dilates venules. Over time, capillary malformations become more similar to venous malformations histopathologically and radiologically. Patients with diffuse capillary malformation with overgrowth of the lower extremity often are erroneously labeled with “Klippel–Trenaunay syndrome” or “Parkes Weber syndrome.”

Diagnostic confirmation of lymphedema–capillary malformation requires imaging (Fig. 3). If this condition is suspected based on history and physical examination, we then obtain an MRI and lymphoscintigraphy. We usually wait until children are at least 12 months of age to obtain the MRI to reduce the risk of anesthesia. Early diagnosis is important for prognosis and management and to rule out Klippel–Trenaunay or Parkes Weber syndrome. Lymphedema only affects the skin and subcutaneous tissues and does not cause axial overgrowth. However, the capillary malformation component might cause vertical bone hypertrophy and a leg length discrepancy. MRI shows that vascular anomalies are not present in tissues below the muscle fascia; this finding does not change over time. Other features of lymphedema–capillary malformation include subcutaneous edema (lymphedema), subcutaneous phlebectasia (capillary malformation), and excess adipose hypertrophy below the skin (lymphedema and capillary malformation). Although MRI can exhibit features consistent with lymphedema (thickened skin, subcutaneous edema and adipose hypertrophy, normal subfascial compartment), diagnostic confirmation requires lymphoscintigraphy.
findings consistent with lymphatic dysfunction include delayed transit time, dermal backflow, and/or abnormal collateral vessels.\textsuperscript{13,14}

The major differential diagnosis of lymphedema–capillary malformation is Klippel–Trenaunay syndrome or Parkes Weber syndrome (Table 1). Individuals with Klippel–Trenaunay syndrome have a capillary–lymphatic–venous malformation of an overgrown extremity.\textsuperscript{20} The condition is caused by a somatic mutation in $\text{PIK3CA}$.\textsuperscript{21} The “capillary” malformation of the leg typically is a lymphatic malformation with bleeding vesicles that cause lymphorrhea. In contrast, a sporadic capillary malformation caused by a $\text{GNAQ}$ mutation does not have vesicles. Some patients with lymphedema–capillary malformation in our series seemed to have a sporadic capillary malformation without evidence of lymphatic etiology, whereas others had a stain that contained lymphatic vesicles. Children with Klippel–Trenaunay syndrome may have an enlarged contralateral foot and exhibit macrodactyly. Klippel–Trenaunay syndrome has venous and/or lymphatic malformations that affect tissues below the muscle fascia.\textsuperscript{4,5,19,20} A large embryonic vein in the subcutaneous tissue (marginal vein of Servelle) is present and pathognomonic for the condition;\textsuperscript{4,5} diversion venography usually demonstrates an intact deep venous system. Endovenous laser ablation of the marginal vein and coil occlusion of the sciatic vein are performed to prevent thromboembolism. Individuals with Klippel–Trenaunay syndrome require monitoring for a leg length discrepancy and may need a shoe lift or epiphysiodesis. Patients often undergo selective amputations to facilitate footwear. Reduction of circumferential overgrowth typically is performed using staged skin and subcutaneous excision.\textsuperscript{20} Cutaneous lymphatic vesicles are managed by sclerotherapy, carbon dioxide laser, and/or resection. Deeper lymphatic and venous anomalies may benefit from sclerotherapy.

Parkes Weber syndrome consists of a combination of a capillary malformation and diffuse arteriovenous malformation of an extremity causing overgrowth.\textsuperscript{20} The capillary malformation is consistent with a sporadic lesion or with capillary malformation–arteriovenous malformation; they typically do not bleed or cause lymphorrhea. The cutaneous stain is warmer than a typical capillary malformation and exhibits fast-flow by Doppler; the lesion may be phenotypically similar to an arteriovenous malformation. The syndrome can be sporadic or familial as a result of a mutation in $\text{RASA1}$.\textsuperscript{22} The tissue below the muscle fascia is affected, and the muscle and bone usually are overgrown.\textsuperscript{19,20} MRI exhibits diffuse subfascial microfistulae represented as flow-voids.\textsuperscript{19} The enlarged limb muscles and bones show abnormal signal and enhancement. Angiography illustrates dilated, tortuous arteries and draining veins with discrete arteriovenous shunting.\textsuperscript{19} Individuals with Parkes Weber syndrome require monitoring for a leg length discrepancy. Patients occasionally need intervention for congestive heart failure and embolization to reduce shunting. Amputation may be necessary.

It is important to differentiate lymphedema–capillary malformation from Klippel–Trenaunay and Parkes Weber syndromes because the prognosis and treatment are different. Lymphedema–capillary malformation exhibits morbidity of both lymphedema and capillary malformation: infection (lymphedema), vesicles (lymphedema), difficulty ambulating (lymphedema), leg length discrepancy (capillary malformation), phlebectasia (capillary malformation), circumferential overgrowth (lymphedema and capillary malformation), and difficulty fitting clothes (lymphedema and capillary malformation). The condition is managed similar to pa-
tients with either isolated lymphedema or capillary malformation. Individuals are prescribed custom-fitted garments and pneumatic compression to treat lymphedema. Pulsed dye laser is considered to lighten the color of the capillary malformation. Because both lymphedema and capillary malformation cause subcutaneous adipose overgrowth, patients with lymphedema–capillary malformation have circumferential enlargement of their lower extremity. Individuals in our series generally had larger extremities compared with patients with either isolated lymphedema or capillary malformation, possibly because the 2 conditions synergistically increased adipose hypertrophy. Although lymphedema does not cause axial skeletal overgrowth, diffuse capillary malformation can cause a leg length discrepancy. Four of the patients in our series had a leg length discrepancy, and thus, children with lymphedema–capillary malformation should undergo orthopedic screening. Patients with symptomatic overgrowth can be treated with suction-assisted lipectomy to reduce the size of their extremity; this intervention is used to manage patients with isolated lymphedema or capillary malformation. Because lower extremity capillary malformations cause phlebectasia, patients may require intervention for symptomatic varicosities. None of the patients with lymphedema–capillary malformation had interventions that are commonly required for Klippel–Trenaunay syndrome or Parkes Weber syndrome (eg, removal of embryonic veins, embolization, amputations, and treatment of macrodactyly).

It is unclear whether patients with lymphedema–capillary malformation have both conditions as a coincidence or whether the 2 vascular malformations are embryologically related. Capillary malformation is caused by a somatic mutation in GNAQ, whereas primary lymphedema may result from several mutations (eg, VEGFR3, SOX18, and FOXC2). We hypothesize that lymphedema–capillary malformation likely results from an anomaly that causes both hypoplastic lymphatic development and capillary malformation. Patients with lymphedema–capillary malformation might have a novel somatic mutation responsible for their condition. Alternatively, lymphedema–capillary malformation may be a phenotypic variant of primary lymphedema or an overgrowth condition with a known mutation (eg, GNAQ, PIK3CA, and RASA1).

**References**