Venous malformations, previously called cavernous haemangiomas, are categorized as slow-flow vascular malformations. A few cases of venous malformations with a dermatomal distribution have been described previously (1–4). We call this clinical entity “zosteriform venous malformations”, while the terms “zosteriform cavernous haemangiomas” and “unilateral dermatomal cavernous haemangiomatosis” were previously used to describe the lesions. We report here a case of venous malformations with a unilateral C6 dermatomal distribution and summarize the available literature regarding venous malformations with this characteristic distribution pattern.

CASE REPORT

A 58-year-old Japanese man presented with a 5-year history of two gradually-enlarging, bluish nodular lesions involving the right upper extremity. The patient was otherwise healthy. No pain or pruritus was noted. The lesions were sessile, spherical, smooth-surfaced, and were located on the lateral flexor surface of the right forearm and the radial aspect of the wrist joint (Fig. 1). They measured 35 mm and 25 mm in diameter, respectively. They were distributed along the C6 dermatome. No tenderness was noted, but palpable pulsations and surrounding dilated veins were found on the wrist lesion. Neither hyperhidrosis over the lesions nor hypertrophy of the right arm was detected. The patient’s family history was negative for similar lesions. Laboratory examinations were unremarkable. No visceral masses were detected in imaging studies, including computed tomography, esophagogastroduodenoscopy, and colonoscopy.

Magnetic resonance imaging (MRI) revealed two intramuscular masses, both measuring approximately 30 mm in diameter, in addition to two peripheral ones corresponding to the skin lesions (Fig. 2). All four lesions were oriented along the long axis of the extremity, strictly following radial neurovascular distribution: two intramuscular lesions followed the deep branch of the radial nerve that runs along the radius, while two superficial lesions followed the superficial branch of the radial nerve. These masses had signal intensities similar to that of skeletal muscle on T1-weighted images, and higher signal intensities in comparison with skeletal muscle on T2-weighted MRI. Inhomogeneous enhancement was observed on gadolinium-enhanced T1- and T2-weighted MRI. Needle puncture of the wrist tumour caused spurts of bright-red blood. Histological examination of an excisional biopsy specimen from the forearm tumour demonstrated multiple irregular dilated blood vessels with congested erythrocytes in the dermis and subcutaneous tissue (Fig. 3). A single layer of flat mature endothelial cells lined each vessel. A diagnosis of venous malformations was made.

DISCUSSION

Zosteriform venous malformations, first reported in 1977 by Steinway & Fretzin. (1), have a distinct distribution pattern limited to unilateral single or multiple adjacent dermatomes, further characterized by the absence of visceral lesions or enchondromas and a negative family history. Blue rubber bleb naevus syndrome is a rare disorder characterized by multiple venous malformations of the skin and gastrointestinal tract (5). Maffucci’s syndrome is also a rare sporadic disorder consisting of enchondromas and vascular anomalies including venous malformations (5). Autosomal dominantly inherited cutaneous venous malformations, caused by mutations in angiopoie-
Malformations with clearly identified intramuscular lesions followed the superficial branch and the deep branch of the radial nerve, respectively, indicating that the lesions are distributed along dermatome, but not Blaschko’s lines, because Blaschko’s lines are not applicable to intramuscular lesions. The cause of this characteristic neurovascular distribution has not yet been determined. However, an increasing body of evidence has allowed us to propose a plausible hypothesis for the development of the characteristic distribution of this disease. Borisov et al. (11) initially demonstrated that selective denervation induces loss of capillaries in skeletal muscle. Bates et al. (12) showed that overexpression of the soluble form of neuropilin-1 causes abnormally formed blood vessels as nerve defasciculation and a severe disruption in the peripheral nerve pattern. Indeed, vascular and nervous systems use similar signals and support each during development of the neurovascular system (13). Thus, we can speculate that ectopic overgrowth of venous blood vessels due to dysregulation of signalling in a particular nervous segment may be involved in the development of venous malformations along dermatomes.

Four further cases with zosteriform venous malformations have been reported thus far (1–4). In three cases lesions were located on several adjacent dermatomes. Although rapid enlargement in puberty and changes in size associated with the menstrual cycle were observed in one case (2), hormonal effects were not noted in the other cases. Although venous malformations are congenital anomalies caused by inborn error of vascular morphogenesis (14, 15), they are not always evident at birth (14). Rapid enlargement can occur in association with trauma, haemorrhage, partial resection, or hormonal changes. The late onset of venous malformations in our case is unusual. Vascular fragility associated with ageing and unnoticed minor traumas may influence the growth of the lesions.

REFERENCES