Multiple cardiac fatty deposits in a patient with tuberous sclerosis complex

Masahiro Kimura,1 Takao Kato,2 Kanae Miyake,3 Takeshi Kimura2

DESCRIPTION
Tuberous sclerosis complex (TSC) is a rare autosomal dominant genetic disease characterised by systemic hamartomas or benign tumours which affects multiple organs including brains, kidneys, heart, liver, eyes, lungs and skin due to inappropriate activation of mammalian target of rapamycin (mTOR) signalling.1 About half of patients have multiple cardiac rhabdomyomas during perinatal and early-neonatal period, but most of them would regress after the age of two. Angiomyolipoma (AML) is the most common tumour of the kidney and usually develops from adolescence.

A 55-year-old man with TSC attended to our hospital and had a CT for routine follow-up with renal AML. He had a history of cardiac rhabdomyoma and also had a subependymal hamartoma, liver haemangiomas, adenoma sebaceum of the skin and angiofibroma of toes. During watchful waiting, multiple fatty foci in the heart were detected by CT (figure 1) without any ECG or ultrasonic echocardiography change. Differential diagnosis of the cardiac fatty-containing lesions associated with TSC included fatty replacement after spontaneous remission of rhabdomyomas,2 3 or cardiac metastasis of AML.4 5 Then, a cardiac magnetic resonance (CMR) was performed to investigate the tissue characterisations and locations of cardiac lesions (figure 2). The lesions had chemical shift on turbo fast low-angle shot images and exhibited homogeneous high intensity on half-Fourier single-shot turbo spin-echo images, which consisted with they being fat. No abnormal enhancement was seen on perfusion images nor late gadolinium enhancement images. CMR suggested all were pure fatty lesions without evidence of residual rhabdomyoma. Most of them were located in the subintimal area of biventricular wall without mass effect, but some of them protruded into cardiac chambers without any blood-flow restriction. Although he had no episode of haemorrhage, pain or renal dysfunction, multiple AMLs in bilateral kidneys had grew slowly and reached to a maximum diameter of 17 cm. Then 5 mg/kg of everolimus, an mTOR-inhibitor, was administered and CT images revealed significant volume reduction of renal AML, while the size of cardiac masses unchanged.

In our case, CMR was helpful to characterise the cardiac lesions as pure fatty changes, and different response to everolimus between renal AMLs and cardiac lesions indicated an unlikelihood of cardiac metastasis of AML. Though a histological biopsy is required for the definitive diagnosis, it seems impractical to perform due to their focal deposition. CMR also provides detailed information about location and size of fatty deposit, then we concluded the observation strategy was reasonable for cardiac lesions. However, long-term prognosis of this fat-containing change in the heart is still unknown, so careful follow-up is needed.
Learning points

- Multiple cardiac fatty deposits are sometimes seen in patients with tuberous sclerosis complex and their prognosis is usually benign.
- Cardiac magnetic resonance imaging is useful to characterise lesions and provide locational information of cardiac fatty deposit.
- Evelomus did not affect cardiac lesions, suggesting the diagnosis of them as fatty replacement of rhabdomyomas and not metastasis of angiomyolipoma.

Contributors
MK wrote the manuscript, and TKato, KM and TKimura were supervisors of the manuscript.

Funding
The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests
None declared.

Patient consent for publication
Obtained.

Provenance and peer review
Not commissioned; externally peer reviewed.

ORCID iD
Masahiro Kimura http://orcid.org/0000-0002-9790-4609

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