Cardiovascular Magnetic Resonance Imaging of Myocardial Fibrosis in Dermatomyositis

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Cardiac involvement in dermatomyositis (DM) is infrequent but is associated with poor prognosis¹. Myocardial fibrosis has been observed in histopathological studies of DM and

has been attributed to myocardial inflammation². Reports describing its noninvasive *in vivo* visualization in DM are lacking. Fibrosis represents a common substrate of ventricular arrhythmias in an array of non-ischemic cardiomyopathies and possibly in DM as well³.

A 58-year-old woman with DM first diagnosed 6 years earlier was referred for cardiovascular magnetic resonance imaging (MRI) because of progressive exertional dyspnea. Examination revealed bilateral atrial enlargement, which was much more pronounced in the right atrium (steady-state free precession). In addition, an area of myocardial late enhancement (inversion-recovery gradient echo) was noted in the basal posterolateral wall, together with a small pericardial effusion (Figure 1). There was no evidence of myocardial edema (T2-weighted image) or hyperemia (early contrast-enhancement), considering the skeletal muscle as an internal reference. Coronary angiography revealed no significant coronary artery stenosis, indicating that the fibrosis was non-ischemic.

The unique ability of cardiovascular MRI to visualize focal myocardial fibrosis noninvasively in this setting may offer a novel method of risk stratification.

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Figure 1. Myocardial late enhancement was noted in the basal posterolateral wall (A: circle), together with a small pericardial effusion (B: arrows). RA: right atrium; RV: right ventricle; LA: left atrium; LV: left ventricle.