Primary antiphospholipid syndrome in a male with myocardial infarction with non-obstructive coronary arteries and a history of stroke

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A 55-year-old man with obesity, hypertension, hyperlipidemia, and a history of right-sided ischemic stroke at the age of 45 was referred to a cardiology outpatient center after hospitalization for non-ST-segment elevation myocardial infarction with non-obstructive coronary arteries (Figure 1A; Supplementary material, Video S1–S6). Non-ST-segment elevation myocardial infarction with non-obstructive coronary arteries was diagnosed based on typical symptoms and dynamics of high-sensitive cardiac troponin T levels: 4.62 ng/l on admission and 114.80 ng/l after 6 hours (normal: 0–14 ng/l). No ischemic changes were observed on the electrocardiogram. Other causes of increased serum high-sensitive cardiac troponin T were excluded. Transthoracic echocardiography showed normal ventricular function, left ventricular ejection fraction of 55%, without significant valve defects.

Three months later, at a follow-up outpatient center visit, he was completely free of cardiovascular symptoms. The electrocardiogram showed pathologic Q waves in the lateral leads, and transthoracic echocardiography showed no abnormalities. Cardiac magnetic resonance imaging showed normal ventricular function, left ventricular ejection fraction of 55%, without significant valve defects.

Since we suspected antiphospholipid syndrome (APS), the plasma levels of antiphospholipid antibodies (aPL) were measured. The lupus anticoagulant in the dilute Russell’s viper venom time (dRVVT) and activated partial thromboplastin time (aPTT) assays was absent. The anticardiolipin antibodies (aCL) IgM were elevated (32.5 MPL [normal, 0–17.0 MPL]) while aCL IgG as well as anti-β2-glycoprotein I antibodies (aβ2GPI) IgM and IgG were normal. A weakly positive titer of antinuclear antibodies (ANA1 1:320) was also found. Single-positive APS was diagnosed.

According to the current recommendations [1–3], aPL levels were reassessed after ≥12 weeks. Elevated aCL IgM were found to persist both after 8 months (32.5 MPL [normal, 0–17.0 MPL]) while aCL IgG as well as anti-β2-glycoprotein I antibodies (aβ2GPI) IgM and IgG were normal. A weakly positive titer of antinuclear antibodies (ANA3 1:320) was also found. Single-positive APS was diagnosed.

Magnetic resonance imaging of the brain showed an extensive area of malacic lesions in the right temporal lobe. Moreover, in the cerebral hemispheres, there were single small areas of raised signal in the sequence with a long time of repetition, consistent with non-specific demyelinating lesions, primarily ischemic (Figure 1C).
In chronic pharmacotherapy, aspirin 75 mg/d, metoprolol succinate 25 mg/d, valsartan 80 mg bid, and rosuvastatin 10 mg/d were used. APS is characterized by venous, arterial, or microvascular thrombosis and/or adverse pregnancy outcomes in the presence of persistent laboratory evidence of aPL [1–3]. As acquired thrombophilia, APS can be diagnosed at any age but is 5-fold less common in men [1–3]. Acute myocardial infarction is a very rare (2.8%) manifestation of APS [4, 5]. Despite indications to use warfarin in APS patients with arterial thromboembolism, our patient was treated with aspirin, given his relatively low aCL levels, with close ambulatory surveillance and follow-up outpatient visits every 6 months. It is also important to search for other cardiovascular risk factors and their appropriate treatment, which can significantly improve the prognosis of APS patients.

**Figure 1.** Coronary angiography showing non-obstructed epicardial coronary arteries (A). The timeline of subsequent diagnostic and therapeutic stages (B). Magnetic resonance imaging brain FLAIR sequence showing an extensive area of malacic lesions in the right temporal lobe (blue arrow). Furthermore, there were single small areas of raised signal in the right cerebral hemisphere, consistent with nonspecific demyelinating lesions, primarily ischemic (yellow arrows) (C).

Abbreviations: AF, atrial fibrillation; aPTT, activated partial thromboplastin time; CT, computed tomography; cTnT, cardiac troponin T; dRVVT, dilute Russell’s viper venom time; LA, left atrium, LVEF, left ventricular ejection fraction; MINOCA, myocardial infarction with non-obstructive coronary arteries; MRI, magnetic resonance imaging; TTE, transthoracic echocardiography; vWF, von Willebrand factor.
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