Tako-tsubo syndrome and atrial fibrillation. New trigger for cardiomyopathy. A clinical case

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SOUHRN

Takotsubo syndrom (TS) je znám i pod označením stresová kardiomyopatie. Spouštěči TS jsou vnější nebo vnitřní, fyzický nebo psychický stres spojený s katecholaminovou bouří, nadměrnou sympatickou aktivitou nervového systému nebo s dysfunkcí mikrovaskulatury v přítomnosti systémového zánětu. Popisujeme případ pacienta s takotsubo kardiomyopatií spojenou s fibrilací síní a s fibromuskulární dysplazií levé přední sestupné větve.

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ABSTRACT

Takotsubo syndrome (TS) is well-known to be a stress-induced cardiomyopathy. TS is triggered by external or internal, physical or emotional stress associated with a catecholamine storm, overactivity of sympathetic nerves, or microvascular dysfunction in the setting of systemic inflammation. We report the case of a patient with Takotsubo cardiomyopathy associated with atrial fibrillation and fibromuscular dysplasia of the left anterior descending artery.

Introduction

TS is a reversible cardiomyopathy characterized by a transient left ventricular dysfunction similar to an acute myocardial infarction but in the absence of obstructive coronary artery disease (CAD). Takotsubo syndrome (TS) is described as a pathology with a good prognosis characterized by the presence of a transient left ventricular wall disease without significant culprit obstructive CAD.^{1–16} The “golden standard” for a definitive diagnosis is invasive coronary angiography and ventriculography, with an integrated multi-imaging approach, such as echocardiography, first-line non-invasive technique, and cardiac magnetic resonance (CMR), in order to discriminate TS from other acute cardiac syndromes with troponin elevation and ventricular dysfunction.^{16–19} Important hallmarks of TS are a relatively

small increase in T/I troponin, the left ventricle “apical ballooning” (wall motion dyskinesia characterized by apical akinesis and basal hyperkinesis) at echocardiography and ventriculography associated with normal epicardial coronary vessels at angiography.^{3,7,14,19–22} In addition there may be catecholamines (epinephrine, norepinephrine, and dopamine) and NT-proBNP elevation.

Case presentation

A 73-year-old woman presented with sudden onset chest pain and dyspnea about six hours after a rapid atrial fibrillation presenting with palpitations. In her history there was arterial hypertension treated with ACE inhibitors and dyslipidemia treated with statins for 10 years. The

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Fig. 1 – (A) ECG: Atrial fibrillation; inferior wall ST segment elevation, and negative T waves in lateral leads (suggestive of inferior-lateral myocardial infarction “STEMI”). (B) TTE revealed apical ballooning, also evaluated with global longitudinal strain (GLS) which is reduced in the recovery phase.

electrocardiography indicated “high-rate atrial fibrillation with ST segment elevation in inferior leads and negative T waves in lateral leads, suggestive of inferior-lateral myocardial infarction, and QT prolongation”, “inferior wall ST segment elevation myocardial infarction”, T wave inversion in apical wall and QT prolongation (Fig. 1A). The clinical course included a severe pulmonary edema complicated by cardiogenic shock.^{1,7,8,19–21} Coronary angiography ruled out obstructive lesion in the major coronary arteries, but revealed fibromuscular dysplasia of the distal tract of the left anterior descending artery. The ventriculography showed remarkable ventricular dilation. Trans-thoracic echocardiography (TTE) revealed a left ventricle (LV) apical ballooning with hyperkinetic basal segments also evaluated and confirmed with global longitudinal strain (GLS, Fig. 1B). LV ejection fraction (EF) evaluated by echocardiography was 35% (in old TTE, one year ago, EF 50%). In the acute phase she was treated with beta-blockers, furosemide and amiodarone i.v. without clinical improvement; therefore we decided to perform electrical cardioversion which restored sinus rhythm. Furthermore, during the hospitalization and immediately after the acute phase she was treated with her usual therapy for hypertension (ACE inhibitors) and dyslipidemia.

Pulmonary congestion disappeared and the patient was dismissed from hospital and got back to her normal life with her past home therapy. Amiodarone has not been prescribed as home therapy due to the presence of QT prolongation on the ECG. Her LV systolic function normalized within one week (EF 50%), except for a residual inferior-apical hypokinesis. She had no more symptoms.

Discussion

This case focuses on the relationship between TS, atrial fibrillation (AF) and coronary fibromuscular dysplasia. In the medical literature, Stiermaier et al.²³ report that in TS patients, atrial fibrillation is frequent and associated with increased long-term mortality rates. A meta-analysis pro-

duced by Prasitlumkun et al.²⁴ suggests that the presence of AF is a prognostic factor for all-cause mortality among patients with TS compared to those lacking it. However, there is no case reported in literature that could define AF as a trigger event. There are several cases of TS associated with coronary anomalies,²⁵ but none of these have atrial fibrillation as a trigger. Even though we don't know the exact etiology of TS, the onset is preceded by a trigger event in the majority of patients (70–80%). Stressor trigger is identified in about 70% of cases. The typical patient with Takotsubo syndrome is a post-menopausal women who has experienced severe, unexpected emotional or physical stress. A genetic predisposition has been suggested related to polymorphisms in relevant candidate genes, such as α_1 -, β_1 -, and β_2 -adrenergic receptors, GRK5, and estrogen receptors. During a paroxysmal atrial fibrillation, psychological stress and/or physical pain can stimulate central/autonomic nervous systems, and increase bioavailability of cortisol and circulating catecholamines, which may affect the myocardium. So, from a pathophysiological point of view it may be a convergence between the classical stressor events and the AF: they both provoke the release of catecholamines which appear to have a central role in the development of Takotsubo syndrome. Also the presence of fibromuscular dysplasia of left anterior descending coronary during atrial fibrillation could result in coronary vasospasm cause circumscribed left ventricular contraction abnormalities found in TS.

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None.

Conflict of interests

None.

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