Introduction

Takotsubo cardiomyopathy, also known as Transient apical ballooning syndrome, stress-induced cardiomyopathy and broken-heart syndrome, is a rare non-ischemic cardiomyopathy that presents as an acute coronary syndrome without evidence of obstructive atherosclerotic coronary disease. Its name is derived from the Japanese Takotsubo – an octopus trap, resembling the elliptical shape of the very typical akinetic left ventricular apex during systole on imaging studies.  

It is nowadays increasingly recognized as a new disease entity when faced with normal coronary arteries on angiography with the very typical left ventriculogram, often presenting with acute heart failure, arrhythmias or rarely ventricular rupture.

Case presentation

A 20 previously healthy year old female (including no known drug allergies) presented for an elective breast lumpectomy. After anaesthetic induction, the patient suddenly experienced a twenty second interval of wide complex irregular tachycardia associated with unrecordable blood pressure. The patient was urgently intubated and resuscitated. There was clinical pulmonary edema present. Blood gases revealed metabolic acidosis with severe hypoxia, despite a high oxygen flow rate. An urgent echocardiogram was done which showed ventricular dilatation with apical and anteroseptal hypokinesia. The atri were normal and the base was spared.

Once in the Intensive Care Unit, inotropes and fluids were given to improve blood pressure and oxygen saturation. The patient was kept sedated with 3mg of midazolam and morphine. The electrocardiogram revealed sinus tachycardia with early onset left bundle branch block with absent elevations in creatinine kinase and troponins.

An echocardiogram on the fifth day (after the acute event) showed normal left ventricular dimensions, global and regional contractility with an ejection fraction of 61%. The patient’s parameters eventually normalised and she was discharged home on an angiotensin converting enzyme (ACE) inhibitor and advised to limit physical activity for the next few weeks. A scheduled coronary angiogram was refused by the patient. A review echocardiogram at outpatients was organised at one and three months post-discharge.

Discussion

Aetiology and Pathogenesis

A variety of psychological and physiological stressors (including anaesthesia) have been implicated in the literature, with one study revealing that such precipitants were present in 61% of cases. Two major pathogenic mechanisms have been proposed: a) catecholamine cardiotoxicity as a primary or secondary phenomenon and b) neurogenic stunned myocardium causing epicardial coronary arterial spasm as a result of an exaggerated sympathetic response.

Focal myocytolysis is consistently present in myocardial histology, characteristically absent in myocardial infarction. In addition, biopsies also often display contraction band necrosis, though the catecholamine-mediated myocardial stunning causing the insult is not consistent.

More evidence for an exaggerated sympathetic response has emerged from studies in ovariectomised female rats in which the syndrome of local apical ballooning provoked by restraining stress could be prevented by β-blockade and attenuated by oestrogen suppletion. However, considering the duration of akinia the and the multivessel coronary

Table 1: Mayo Clinic major criteria for diagnosing Takotsubo cardiomyopa

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<th>Major Criteria</th>
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<td>1. On echocardiography/ventriculogram, transient dyskinesis/akinesis of the left ventricular mid-segments with or without apical involvement with no single arterial territory involved, with or without a stressful trigger and accompanied with a massive decrease in left ventricular ejection fraction</td>
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<td>2. Absence of obstructive coronary disease or angiographic evidence of acute plaque rupture</td>
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<td>3. New electrocardiographic abnormalities</td>
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<tr>
<td>ST-segment elevation and/or T-Wave inversion OR Modest elevations in cardiac troponins</td>
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<td>4. Absence of phaeochromotycoma or myocarditis</td>
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Interestingly, a number of cases have been reported that are similar in presentation to Takotsubo cardiomyopathy but do not however manifest the typical elliptical shape of the left ventricular apex, a presentation described as an ‘inverted Takotsbo’.16,23
spasm required for such an extensive apical wall abnormality, conventional coronary vasospasm due to sympathetic over-response seems improbable.6  

The apical myocardium manifestations could be explained by its high vulnerability towards adrenergic aggression,12 a statement consistent with similar wall abnormalities observed in phaeochromocytoma-related cardiomyopathy.13 However, recently several other forms (Types I-V) of stress-induced cardiomyopathy have been described.14-16  

Epidemiologically  

The syndrome has a higher preponderance for the female gender in the over sixties, with estrogen possibly playing a role.17,18 Reports in children and young adults have also been reported.19,20 A genetic role might also be possible after an isolated report described the disease in two sisters.21  

Clinical Features and Investigations  

The patient’s presentations are very non-specific, with sudden onset of symptoms resembling an acute STEMI, with or without cardiogenic shock and arrhythmias.1 The Mayo Clinic have drafted up four major criteria which have to be present in order to diagnose Takotsubo cardiomyopathy.22  

Management  

The specific treatment of the condition is still largely empirical due to the limited availability of controlled data. Drugs including diuretics, β-blockers such as carvedilol, and ACE inhibitors are often used until recovery of LV function, with no evidence available for their use after recovery. Most importantly, anti-platelets should be considered until a thrombotic pathogenesis is excluded, during the apical akinesis or dyskinesis interval to resolve the cardioembolic risk. Some physicians also consider inotropes or intra-aortic balloon counterpulsation, with the latter being the preferred option due to the potential role of catecholamine excess in the pathogenesis. Follow-up echocardiographic evaluation is routinely performed to ensure resolution of the left ventricular dysfunction and improvement in the ejection fraction.24,25  

Prognosis  

Although the prognosis for most patients with this syndrome is favorable, with complete recovery of ventricular function within 1 to 4 weeks, several cases of fatal outcomes have been reported.17,26 The evolution, although mainly uneventful, can be complicated, rarely, by left ventricular rupture and ventricular tachycardia, possibly causing sudden death. The recurrence of this syndrome seems to be rare.1  

Conclusion  

The incidence of “broken heart syndrome” has not as yet been ascertained with the prevalence likely to be under-estimated because of the low level of awareness and infrequent diagnosis. It is nowadays been increasingly recognized in clinical practice which is why more research is needed to determine the exact pathogenesis, increase awareness and optimize management of the syndrome, especially in the acute setting, and identify those subjects prone to this potentially lethal condition.  

References may be accessed at www.thesynapse.net