A Broken Heart Syndrome in an Electrocardiogram with a short PR-Interval

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Introduction

Takotsubo disease is being widely studied in recent times and different terms have been used to describe the disorder. This form of cardiomyopathy is of a non-ischemic nature characterized by sudden temporary weakening of the myocardium (1, 2). Such sudden weakness looks be triggered by physical or emotional stress (3), as in the case of the death of a loved one. For this reason the disease is often also known as “broken heart syndrome”. It has also been reported in cases of partial drowning.

The typical presentation of Takotsubo cardiomyopathy comprises sudden-onset congestive heart failure and/or chest pain associated with electrocardiographic abnormalities suggestive of acute myocardial ischemia of the anterior wall (4). Throughout the course of the evaluation of the patient, we often observe a bulging of the left ventricular apex with basal hypercontractility of the left ventricle (as established by echocardiography). This is the main characteristic of Takotsubo disease, which means “octopus trap” in Japanese (the first case of the disorder being reported in Japan).

The underlying cause appears to involve high levels of circulating catecholamines (mainly adrenaline / epinephrine) (1,3,4), but this aspect has not been studied in depth. Further studies are therefore needed to confirm this origin of the disease.

Another very important feature in Takotsubo disease is that the blood levels of the typical cardiac markers of myocardial infarction are normal or only very discreetly elevated.

The evaluation of patients with Takotsubo cardiomyopathy typically includes coronary angiography, which may not reveal significant obstructions capable of causing left ventricular dysfunction.

Among survivors of the initial presentation of the disease, left ventricular function is seen to improve within two months (4).

Takotsubo cardiomyopathy looks more common in postmenopausal women, and patients usually present a recent history of severe physical or emotional stress (1-4).

A Short PR-Interval (5, 6) may be associated with an otherwise normal electrocardiogram or an untold number of bizarre electrocardiographic abnormalities. Clinically, the individual may be asymptomatic or experience a variety of complex arrhythmias, which may be disabling but rarely cause sudden death. The PR-interval starts from the beginning of the P-wave (sinoatrial node depolarization), and includes the whole P-wave, i.e., the full duration of atrial depolarization. This is followed by a flat segment until depolarization reaches the atrioventricular (AV) node, creating an electrical interlude. The AV node delays conduction of the electrical impulse long enough for the ventricles to be filled by atrial contraction before the ventricles themselves contract.

The PR-interval ends as ventricular depolarization begins (the start of the QRS complex). Thus, the PR-interval represents the time it takes for the atria to depolarize and transmit electrical communication to the ventricles. It is measured from the beginning of the P-wave to the beginning of the QRS complex. The normal PR-interval measures 0.12 to 0.20 seconds in length. A short AV conduction time, whether associated to a normal or abnormal QRS complex, is correlated to an increased incidence of paroxysmal rapid heart beats. A considerable number of patients have a short PR-interval, a normal QRS complex and bouts of tachycardia.

Evidence is presented suggesting the action of endocrine and autonomic nervous system factors in the origin of both the short PR-interval and Takotsubo disease.

Nevertheless, the coexistence between Takotsubo disease and a short PR-interval has been little studied to date (7).

In 1952, Lown, Ganong and Levine (L-G-L) described an abnormal shortening of the PR-interval regarded as a pre-excitation syndrome, a case of true accelerated atrioventricular conduction. L-G-L syndrome may affect about 1/50,000 persons. In the absence of significant structural heart disease, the mortality rate appears to be low, though fatalities are not uncommon. Patients may experience an acute episode of tachycardia or a history of symptoms suggestive of paroxysmal tachycardia. Sudden death is caused by the presence of ventricular fibrillation; it not previously diagnosed and not has been treated correctly.
Short Communication

We report a case in which this association can be seen.

A 54-year-old Caucasian women had lost her husband 6 days ago. She was rushed to hospital due to sudden dyspnea and oppressive chest pain suggestive of acute coronary syndrome. She was transferred to the emergency room of our center by ambulance.

Once in the emergency room, the cardiologists requested an electrocardiogram, echocardiography, and subsequently angiography. A blood test was also requested, placing special emphasis on cardiac markers. Blood testing showed no increase in cardiac markers. Troponin (I, T) proved normal, in the same way as CPK, CK, LDH (this being another common feature of Takotsubo disease). Repeated cardiac marker testing 4 and 8 hours after first determination also proved normal, thus confirming severe myocardial suffering rather than manifest muscle damage as such.

The electrocardiogram showed signs of cardiac ischemia (Figure A).

Echocardiography in turn showed acute left ventricular dilatation (Figure B).

Angiography revealed no obstructive lesions in any coronary artery (Figure C).

Ventriculogram revealed the typical image of ventricular “ballooning” (Fig. D).

Important findings were the fact that her husband had died 6 days earlier, as well as a clinical history of sudden tachycardias from her childhood, fundamentally at rest.

Following stabilization of the acute event, the patient was closely monitored.

The blood cardiac marker levels remained normal. The electrocardiographic abnormalities also normalized, except for the length of the PR-interval. The echocardiographic anomalies were not seen to persist.

The case was diagnosed as Takotsubo disease; however, a short PR-interval, less 0.120 sec. in length, can be seen on all 12 leads on her ECG tracing (she was confirmed like a LGL pattern by posterior genetic studies; she was positive for PRKAG2 gene).

Following the prescription of appropriate treatment for cardiac ischemia, the patient was discharged from hospital. However, the short PR-interval was not diagnosed.

More often than we think, this feature might be present but is overlooked.

The presence of a short PR-interval would be an added risk factor for new cardiac events such as sudden tachycardia in bursts (8, 9), and particularly life-threatening as ventricular tachycardia or fibrillation.

We have already published more than one case with these features (7, 8).

It therefore must be stressed that this disorder exists and should not overlooked. The presence of a short PR-interval is also present in other cardiac diseases as in the Wellens echocardiographic pattern, as we have shown (7).

Summarizing

Takotsubo disease is being increasingly studied, but we do not know still its fundamental etiology. Its occurrence is not uncommon in individuals with previous cardiac electrical disorders. The prognosis may be bleak when both circumstances manifest in the same individual. Evaluation of an altered PR-interval always should be made before hospital discharge.

References

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Illustrations

Illustration 1

The electrocardiogram showed signs of cardiac ischemia

Illustration 2

In the second image, we can see on echocardiography, the typical dilation of the left ventricle in cases of Takotsubo.
Illustration 3

In the third image, we can see a normal angiography, unobstructed.

Illustration 4

In the fourth image, we can see the typical image of ballooning from Ventriculogram.
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