Type 1 Brugada Phenocopy in a Patient with Stubborn Congestive Heart Failure
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Introduction
Brugada syndrome (BrS) is an inherited cardiac ion channelopathy, which can induce malignant arrhythmias and sudden cardiac death (SCD). The BrS ECG is characterized by an abnormal ST-segment elevation of at least 2 mm (0.2 mV) in leads V1-3, including three types. Type-1 manifests a coved pattern, which obtained the most diagnostic significance [1-3]. Brugada phenocopy (BrP) is referred to as a clinical situation that has an identical ECG pattern to congenital BrS but is induced by various other clinical factors. In this report, we describe the case of severe heart failure, in which the Type-1 Brugada pattern was recorded.

Case Report
A 71-year-old female was hospitalized with urinary tract infection, accompanied by fatigue and exertional dyspnea complaint for a week. Examination on admission: WBC 11.54×10⁹/L, BNP 2373.76 pg/ml, Crea 515 μmol/L, ECG: sinus rhythm (71 bpm). When the patient was routinely disposed with anti-infection, reinforcement myocardial systole, diuretic, and other symptomatic measures for 7 days, the BNP gradually decreased to 630.69 pg/ml, with discomfort complaint relieved. On the 16th day, the patient has repeated episodes of atrial fibrillation (AF) with rapid ventricular rate (150-160 bpm), accompanied by the symptoms of chest tightness, dyspnea, and orthopnea later, with an abnormal increase of BNP levels to 1950.82 pg/ml. On the 18th day, the patient occurred palpitations again. The ECG (Figure 2) showed the rapid AF (132 bpm) combined with Brugada pattern. The patient never had a similar BrP ECG before, and had no SCD family history. Echocardiography showed that the ejection fraction (EF) was reduced to 47%. The BNP > 6000 pg/ml, NT-pro BNP > 35000 pg/ml, Na⁺ 139mmol/L, K⁺ 4.1 mmol/L, Ca²⁺ 1.99 mmol/L, D-dimer: 1.26 μg/ml (<0.5), cTnI: 0.063 μg/L. The 18-lead ECG retested 4 hours later, showed that the BrP still existed, and no ST-segment mirror changes were observed in leads V7-9. Continuous intravenous recombinant human brain natriuretic peptide (rhBNP), lanatoside C (cedilanid), and diuretic to counter heart failure, as well as anti-infection treatment. 3 days later, the patient’s NT-pro BNP was significantly decreased, the ECG showed sinus rhythm (90 bpm), and the symptoms improved dramatically. The patient was discharged home on the 35th day, with regular follow-ups in the heart failure clinic. Up until now (2 years later), no arrhythmic events have occurred.

Discussion
BrS and BrP
BrS is an inherited cardiac ion channelopathy with a potential risk of death. However, BrP changes refer to the influence of some specific clinical
conditions [4], such as fever, metabolic disorder, mechanical compression, myocardial ischemia, pulmonary embolism, and even some drugs. Nevertheless, it is the most difficult to distinguish the true congenital BrS owing to identical ECG patterns. Patients with only BrP and no family history indicate a low probability of true congenital BrS. Most significantly, patients with BrP obtain a negative induced reaction with sodium channel blockers, while patients with BrS obtain a positive induced reaction [5]. For BrS, the primary goal is to prevent SCD by requiring ICD once definitely diagnosed. For BrP, once the inducement factors are corrected, it disappears. Hence, patients with BrP usually gain a better prognosis than BrS.

The Differential Diagnosis of Brugada Wave

Electrolyte Disturbance: In hypercalcemia, the accelerated flow velocity of calcium shortened the 2 phases of action potential duration (APD), which could shorten or even make the ST-segment not obvious. Hyperkalemia generally causes some changes in ECG pattern in which a prolonged QTc interval with a peak T wave [6]. However, the patient has normal electrolytes.

Myocardial Infarction: The ECG of anterior myocardial infarction is manifested as an abnormal ST-segment elevation in leads V1-3. In this case, the retested 18-lead ECG showed that ST-segment elevation in V1-2 still existed, but no ST-segment mirror changes were observed in leads V7-9. Combined with no apparently abnormal cTnl, myocardial infarction was not considered.

Right Bundle Branch Block (RBBB): The RBBB ECG is characterized as a broad R wave or rsR’ morphology of QRS complex in leads V1-2, and a broad S wave in leads I, avL, and V5-6. In this case, the ECG showed apparently J point upward and “coved” elevation of ST-segment, while the RBBB commonly presented ST-segment depression without shortening in the lead V1-2.

The Etiology Analysis of BrP in this Case

In this case, we tend to consider that BrP is caused by myocardial ischemia due to acute heart failure. Repeated and persistent episodes of tachycardia deteriorate cardiac systolic function. Within only 1 day, the level of BNP aggravated sharply from 1950.82 pg/ml to more than 6000 pg/ml, with an obvious change of stubborn and congestive heart failure. Myocardial ischemia is a well-known and published etiology of BrP. In this case, the mechanism could be related to the transient ion channel dysfunction. The outward potassium current (Ito) is increased, and the balance between inward and outward currents is broken, inducing the J-point and ST-segment elevations in leads V1-2 [7].

Conclusion

Whether Brugada syndrome or Brugada phenocopy, further examination should be recommended to get a definitive diagnosis and make appropriate therapeutic decisions.

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References


Figure 1: ECG tracings documented. (A) On admission, the 12-lead ECG showed sinus rhythm, without RBBB or ST-segment elevation in leads V1-2. (B) On the 16th day, the 12-lead ECG repeatedly showed episodes of rapid AF, without RBBB or ST-segment elevation in leads V1-2.

Figure 2: On the 18th day, the 18-lead ECG showed a “coved-like” ST-segment elevation in leads V1-2, combined with rapid AF.


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