INTRODUCTION

Electrocardiographic abnormalities in patients with the Brugada Syndrome may be transient, and ajmaline or flecainide tests can uncover such abnormalities.\textsuperscript{1,2} These tests may, however, cause conduction disorders and arrhythmias.\textsuperscript{3,4}

We report electrocardiographic findings in a patient who presented variable QTc prolongation and T-wave alternans.

CASE REPORT

A 56-year-old man with no history of cardiovascular disease was admitted to the emergency service after receiving an electric shock. The electrocardiogram (ECG) showed a prominent J-wave and ST-segment elevation of up to 5 mm in leads V1 to V3 (Figure 1A). Total creatine kinase (CK) increased but the echocardiogram was normal. After 24 hours, the prominent J wave was still observed unchanged in leads V1 and V2, but it had disappeared from lead V3 (Figure 1B and 2A).

The patient was suspected of having the Brugada syndrome,\textsuperscript{5} so an intravenous ajmaline test was performed (50 mg). After 60 seconds, J-point and ST-segment elevation in leads V1 to V3 were seen (Figure 2B). The maximum elevation of 12 mm occurred in V2 (Figure 2C). After 5 minutes, the QTc intervals were prolonged from 372 to 539 ms and the T wave became negative (Figure 3A), followed by T-wave alternans (Figure 3B). Finally, the QTc interval stabilized at 539 ms (Figure 3C).

The ECG returned to normal after administration of isoproterenol (Figure 3D). During one year of follow-up, the QTc values were normal and the ECG traces were similar to that of Figure 2A.

DISCUSSION

The Brugada syndrome can coexist with hereditary long QT syndrome (LQT3) and patients’ ECGs can show both afflictions.\textsuperscript{6} The Brugada syndrome appears
when the heart rate is faster and the QTc interval is smaller, whereas LQT3 presents at slower heart rates and a longer QTc interval.\(^7\)

Sodium channel blockers lead to a decrease in phase 0 and 1 amplitudes in the epicardium, with loss of the action potential dome and action potential shortening. This creates an electrical gradient between the epicardium and the endocardium, leading to a marked ST elevation in the ECG.\(^6\) However, when the effect wears off, variations in the duration and amplitude of the subepicardial action potentials appear. Moreover, these variations are independent of heart rate.
rate. Thus, the QTc intervals in the ECG are unevenly prolonged and T-wave alternans appears. Administration of isoproterenol shortens the duration of the subepicardial action potentials, improves intramyocardial conduction and normalizes the QTc interval.

REFERENCES