Is the value of QT dispersion a valid method to foresee the risk of sudden death? A study in Becker patients

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Sudden cardiac death is a dramatic, undesirable event that can often result from cardiomyopathies. To investigate the validity of the QT dispersion (QTd) in revealing regional heterogeneity of repolarisation, with consequent possibility of sudden death,1 we evaluated ECGs of patients affected by Becker muscular dystrophy (BMD). This is an X linked recessive muscular dystrophy caused by dystrophin anomalies in striated muscles with myocardial involvement and consequent dilated cardiomyopathy, ventricular arrhythmias and, in 30% of cases, sudden cardiac death.1

METHODS
The study was retrospective using the clinically validated database of ECGs and echocardiograms (echo recorded together with the ECG) from 30 BMD patients (mean (SD) age 25 (10) years) with variable stages of myocardial involvement, and 26, age matched controls. All subjects underwent a physical examination, blood analyses, and M mode and two dimensional echocardiography. The diagnosis of the type of muscular dystrophy in Becker patients was confirmed by DNA analysis (polymerase chain reaction) and by reduced dystrophin labelling from the immunohistochemical examination of biopsy samples.

We excluded the ECGs from subjects with SFT anomalies on the 12 lead ECG, and with sustained ventricular arrhythmias at 24 hour Holter monitoring, electrolyte imbalance, or QRS duration longer than 120 ms and from subjects under cardiovascular treatment.

The standard 12 lead ECG, recorded at a paper speed of 25 mm/s (gain 10 mm = 1 mV) on a three channel recorder, was analysed for each subject by a blinded observer using a digitiser connected to a computerised system. QRS, RR, QT, JT (the latter obtained from the formula QT-QRS) were measured over three consecutive cycles. When U waves were present, the end of the T wave was considered the nadir of the curve between the T and the U waves. The QTd was defined as the difference between the maximum and minimum QT values. QT and JT were further corrected according to Bazett’s formula

$$QTc = \frac{QT}{\sqrt{RRI}}$$

$$JTc = \frac{JT}{\sqrt{RRI}}$$

The QTC and JTc values were calculated in Becker patients who died suddenly, the survivors and those from the remaining BMD population. The results obtained comparing the electrocardiographic data from the two groups was found in RR, QT, JT, and JTc. A significant increase (p < 0.003) in QRS duration and a highly significant difference (p = 0.0001) in ventricular recovery time dispersion indexes (QTc, QTd) were observed in Becker patients compared to normal subjects.

A scattergram was made to compare the values of QTd calculated in Becker patients who died suddenly, the survivors and control group.

The relationship between echocardiographic parameters and QTd in Becker patients were statistically evaluated using the linear regression and correlation analysis. Electrocardiographic data are presented as mean (SD). Mann-Whitney non-parametric test and Student t test for unpaired data were chosen to compare the electrocardiographic data from the two groups. The same statistical tests were used to compare the ECG and echo data obtained from the eight BMD patients who died suddenly and those from the remaining BMD population. A probability value of p < 0.05 for both Mann-Whitney non-parametric test and Student t test for unpaired data was considered significant. No significant linear correlation was found between echocardiographic parameters (ejection fraction, fibre shortening) and QTd in Becker patients.

RESULTS
Findings from the electrocardiographic and echocardiographic parameters in BMD patients and in controls are presented in table 1. No significant difference between the two groups was found in RR, QT, JT, and JTc. A significant increase (p < 0.003) in QRS duration and a highly significant difference (p = 0.0001) in ventricular recovery time dispersion indexes (QTc, QTd) were observed in Becker patients compared to normal subjects.

The results obtained comparing the electrocardiographic and echocardiographic parameters from the eight Becker patients who suddenly died and from those who survived lend
weight to the study. In fact, although no sustained ventricular arrhythmias (>30) were seen on the 24 hour Holter monitoring—recorded at the same time of the examined ECG—in patients who died and no differences were noted in their clinical conditions, or in echocardiographic parameters compared to the survivors (one patient with heart failure among the eight who died suddenly and one among the survivors), a highly significant difference \( (p = 0.0001) \) is seen between the QTd in the two groups. Figure 1 shows that none of the Becker patients who died had QTd values < 100 ms, and six of them had values higher than the maximum QTd value recorded in the survivors. This latter observation shows the need for careful monitoring of Becker patients; a 24 hour Holter must be performed every month when the QTd value reaches 100 ms and, if it reveals runs of ventricular arrhythmias, pharmacological treatment with antiarrhythmic drugs (amiodarone) should be given. If pharmacotherapy is ineffective, a cardioverter-defibrillator should be implanted.

Our findings demonstrate that the analysis of dispersion of ventricular activation in the 12 lead surface ECG (QTd) was significantly increased in BMD patients; this parameter is principally related to non-homogeneous electrical activity. The case with which the QTd can be calculated, and the possibility of using the QTd value as a valid parameter to predict the risk of sudden death, may provide opportunities through anti-arrhythmic drug administration, or an implanted cardioverter-defibrillator, to prevent this dramatic event.

**IMAGES IN CARDIOLOGY**

**Magnetic resonance angiography in the evaluation of aortic pseudoaneurysm**

Magnetic resonance angiography (MRA) and transoesophageal echocardiography together are very useful imaging techniques in the evaluation of thoracic aortic pseudoaneurysms. A 63 year old hypertensive man was admitted to a community hospital by his primary care physician with three weeks history of night sweats, chills, fever, fatigue, and generalised myalgia. Blood cultures done in the outpatient clinic revealed group D salmonella bacteraemia. Transthoracic echocardiogram, abdominal computed tomography (CT), and ultrasound were normal. CT of the chest showed increased density within the anterior mediastinum, suspicious of an aortic pseudoaneurysm.

MRA of the thoracic aorta (left) showed two infected pseudoaneurysms, one in the descending aorta starting below the aortic arch (A), extending 7 cm caudally, and another smaller one in the lesser curve of the aortic arch (B). The latter (B) was missed on transoesophageal echocardiography, although the former (A) was seen clearly. Preoperative coronary angiography showed critical three vessel disease. The patient successfully underwent resection of aneurysms showing aortic pseudoaneurysm.

Once invariably a fatal condition, infected aortic pseudoaneurysms can now be cured in most cases if diagnosed early and treated with appropriate surgical technique and antibiotic. Both MRA and transoesophageal echocardiography complement each other in the evaluation of this condition.
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