Ventricular tachycardia during exercise testing as a predictor of sudden death in patients with chronic chagasic cardiomyopathy and ventricular arrhythmias

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Abstract
Objective—To verify the prognostic value of exercise induced ventricular arrhythmias in patients with chagasic cardiomyopathy.

Methods—69 consecutive patients (37 male, 32 female; age range 21–67 years) with chronic chagasic cardiomyopathy and ventricular arrhythmias (more than 10 ventricular premature complexes per hour) were evaluated during treadmill exercise testing, using the Bruce protocol. Protocol end points were peak heart rate or presence of sustained ventricular tachycardia.

Main outcome measure—Sudden cardiac death.

Results—44 patients (group I) developed ventricular tachycardia during exercise testing (five sustained and 39 non-sustained), and 25 did not (group II). After a follow up of 24 (SD 15) months sudden cardiac death occurred in seven patients in group I and in none in group II (P < 0.05).

Conclusions—Ventricular tachycardia on exercise testing is significantly associated with sudden cardiac death in patients with chronic chagasic cardiomyopathy and ventricular arrhythmias.

Keywords: chagasic cardiomyopathy, ventricular tachycardia, exercise testing

Exercise results in a series of alterations which can increase circulating catecholamines and the sympathetic drive to the heart.¹ These physiological changes can affect the myocardium and initiate a ventricular arrhythmia, specially in patients with cardiac disease and left ventricular dysfunction.² However, the clinical value of exercise testing for patients with ventricular arrhythmias and coronary heart disease is not established.

Chagas disease is one of the most important cardiac diseases in South America.³ Left ventricular dysfunction, ventricular arrhythmias, and autonomic changes are common findings in this disease⁴ and sudden death is an important problem in areas where the disease is endemic.⁵ The prognostic importance of ventricular arrhythmias during exercise testing has been studied in patients with coronary artery disease.⁶⁻¹⁰ The significance of exercise induced arrhythmias has not been reported in Chagas disease. This study was undertaken to verify the prognostic value of exercise induced ventricular arrhythmias in patients with chagasic cardiomyopathy.

Methods
DEFINITIONS
Chronic chagasic myocarditis was defined as the presence of chronic cardiomyopathy and a positive Machado Guerreiro serum complement and haemaglutinin test.

Ventricular tachycardia was defined as ≥3 sequential ventricular complexes at a rate of >100/min. If this arrhythmia lasted more than 30 seconds or resulted in cardiovascular collapse, it was defined as sustained ventricular tachycardia; if not it was considered to be non-sustained ventricular tachycardia.

PATIENTS, MATERIALS, AND PROCEDURES
The study population included 69 patients with chronic chagasic cardiomyopathy and ventricular arrhythmias. All patients had their antiarrhythmic drugs discontinued for at least five half lives and had more than 10 ventricular premature beats per hour or at least one episode of ventricular tachycardia during a 24 hour Holter monitoring. Three of these patients were on amiodarone therapy (low dose of 200 mg orally per day) and the drug was discontinued for one month before the patient entered the protocol. There were no patients receiving β blocker therapy. There were 37 men and 32 women, with ages ranging from 21 to 67 years (mean 46, SD 12, years). Thirty two patients had palpitations, 16 had syncope, and seven had clinical documentation of sustained ventricular tachycardia or ventricular fibrillation. Forty eight patients had a history of congestive heart failure; of these 39 were in New York Heart Association (NYHA) class II, and nine were in class III or IV. Each patient was in a compensated state before entering the protocol.

Echocardiography was performed in all patients and ejection fraction was calculated by the Pombo method.¹¹ Patients suspected of having coronary artery disease underwent a coronary angiography study. No patient had any other identifiable organic heart disease.

Treadmill exercise testing—Exercise testing was performed on a motor driven treadmill
(Model Funbec, ESDO-1), according to the standard Bruce protocol. Twelve-lead ECGs were performed before and after exercise. Leads II and V5 were continuously recorded during exercise and for eight minutes afterwards. The mean recording time for each patient was 17 minutes. Criteria for terminating the exercise were occurrence of maximum heart rate, sustained ventricular tachycardia, or fatigue.

Follow up—Patients were seen in an arrhythmia research clinic by one of the investigators every three months. Mean follow up was 24 (15) months.

STATISTICAL ANALYSIS
The results are presented as mean (SD). Statistical comparisons were made using the Student t test. Contingency tables were evaluated by $\chi^2$ analysis and significance was defined as a probability (P) less than 0.05 using a two tailed analysis.

Results
The results are shown in the table.

ELECTROCARDIOGRAPHY
Right bundle branch block, either isolated (six patients) or associated with left anterior hemiblock (26 patients) was present in 32 patients (46%). Left anterior hemiblock was present in 13 patients (18-9%), left bundle branch block in four patients (5-8%), and complete AV block in the His-Purkinje system in four patients; the latter four patients had VI pacemakers implanted. Sixteen patients (2.3%) did not have conduction disturbances on the surface electrocardiogram.

HOLTER MONITORING
All patients had more than 10 ventricular premature beats per hour; 43/69 (62%) had more than 100 per hour; 43/69 (62%) had non-sustained ventricular tachycardia; and 32/69 (46%) had both non-sustained ventricular tachycardia and more than 100 ventricular premature beats per hour.

EJECTION FRACTION
Ejection fraction was estimated by two dimensional echocardiography. Mean ejection fraction was 46.6 (18.6)%.

ARRHYTHMIAS DURING EXERCISE TESTING
Two patients did not have any arrhythmia during exercise; 10 (14%) had isolated premature ventricular beats, 13 (18%) had couplets, 39 (56%) had non-sustained ventricular tachycardia, and five (7%) had non-sustained and sustained ventricular tachycardia. The four patients with complete AV block were pacemaker dependent during exercise; all of them developed arrhythmias during exercise testing: one couplets, two non-sustained ventricular tachycardia, and one sustained ventricular tachycardia. There was no change in the QTc interval during exercise when compared with control values.

MEDICAL THERAPY AND CLINICAL OUTCOME
Patients were seen in an arrhythmia research clinic by one of the investigators every three months. Thirty six patients (52%) were off drugs. Antiarrhythmic drugs were prescribed to patients with clinical sustained ventricular arrhythmias, syncope, and inducible ventricular arrhythmias and for those with disabling palpitations. After completing clinical and laboratory evaluation 33 patients (48%) were on antiarrhythmic drugs because of symptoms: 23 patients on amiodarone, three on quinidine, three on propafenone, one on amiodarone and propafenone, one on disopyramide, and two on phenytoin. During a mean follow up of 24 months 12 patients died; seven of these had sudden cardiac death. Of the patients with sudden death, five were on drug therapy (three on amiodarone, one on propafenone, and one on disopyramide).

Ventricular tachycardia during exercise testing was the only variable that significantly influenced sudden cardiac death in this study population (table).

Discussion
Regardless of the mechanisms of cardiac arrhythmias the sympathetic nervous system and circulating catecholamines are very important in arrhythmogenesis. The adrenergic state can suppress or provoke cardiac arrhythmias; serious arrhythmias can be provoked or exacerbated by exercise. Also, patients with exercise induced ventricular tachycardia may be more sensitive to plasma noradrenaline than other patients. The majority of these reports are from patients with coronary heart disease; in some there is a correlation between the occurrence of exercise induced ventricular arrhythmias and significant coronary artery disease and impairment of left ventricular function. On the other hand patients with exercise induced non-sustained ventricular tachycardia and normal cardiac function have a good prognosis. However, the clinical significance of exercise testing in patients with ventricular arrhythmias remains poorly defined.

Chronic chagasic myocarditis is a cardiac
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neuromyopathy with sympathetic ganglion denervation and dysfunction of the parasympathetic autonomic control of the heart. This gradual autonomic denervation in chronic Chagas cardiomyopathy can partially explain sudden death in some patients.

Our study population was ambulatory and two thirds of them had a left ventricular ejection fraction above 40% and only 13% had class III or IV congestive heart failure. These patients with only mild left ventricular dysfunction were different from the usual referral cases, where sustained ventricular tachycardia, disabling symptoms, and severe left ventricular dysfunction were more common. Additionally, the autonomic dysfunction and the sympathetic reserve of the heart was probably better than in more advanced cases of Chagas cardiomyopathy. Whether the degree of autonomic dysfunction and catecholamine sensitivity explains the results of our study remains unclear. Nonetheless, ventricular tachycardia during exercise testing was the only variable which was significantly associated with sudden cardiac death in our study population with chronic chagasic cardiomyopathy.

The results of our study are applicable to our population with relatively well preserved ventricular function. In this particular subset, exercise testing may be able to select patients who need more aggressive anti-sudden-death therapy. However, the role of inducibility of ventricular tachycardia by programmed stimulation and its suppression with antiarrhythmic drugs needs to be assessed relative to exercise testing in these patients.