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Recovery pattern of left ventricular dysfunction following radiofrequency ablation of incessant supraventricular tachycardia in infants and children

J V De Giovanni, A Dindar, M J Griffith, R A Edgar, E D Silove, O Stumper, J G C Wright

Abstract

**Objective**—To assess recovery pattern of left ventricular function secondary to incessant tachycardia after radiofrequency ablation in a group of infants and children.

**Design and setting**—A combined prospective and retrospective echocardiographic study carried out in a tertiary paediatric cardiac centre.

**Patients**—Echocardiographic evaluation of left ventricular size and function in nine children with incessant tachycardia, before and after successful radiofrequency ablation. Age at ablation ranged from 2 months to 12.5 years (mean 4.1 years). Recovery of left ventricular function was analysed in relation to age at ablation (group I < 18 months, group II > 18 months).

**Main outcome measure**—Ventricular recovery pattern.

**Results**—Seven of the nine children had left ventricular dysfunction; six of these also had left ventricular dilatation. All children with left ventricular dysfunction had normalisation of ejection fraction and fractional shortening; left ventricular dilatation also improved, but the improvement occurred after recovery of function. There was a shorter recovery time for left ventricular function in younger (group I) than in older children (group II) (mean (SD) 5.7 (7.2) months vs 31.3 (5.2) months (p < 0.002).

**Conclusions**—Tachycardia induced cardiomyopathy is reversible following curative treatment with radiofrequency. Recovery of left ventricular systolic function precedes recovery of left ventricular dilatation. Time course to recovery is shorter in younger children.

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Keywords: radiofrequency ablation; left ventricular dysfunction; incessant tachycardia; echocardiography; children

Incessant or persistent supraventricular tachycardia can lead to a clinical picture of dilated cardiomyopathy with an enlarged left ventricle and reduced systolic function. The duration and the rate of the arrhythmia determine the onset of tachycardia induced cardiomyopathy.

Reversibility of cardiac dysfunction has been shown following control of the arrhythmia or after successful curative treatment of the underlying substrate, either by surgery or by catheter techniques. The pattern and time scale of recovery following curative treatment of incessant tachycardia with radiofrequency ablation in children is not documented, however.

Radiofrequency (RF) ablation has been shown to be a very effective method of treatment for several types of arrhythmias in children, including the incessant forms. Some indications for RF ablation include drug refractory or life threatening arrhythmias and haemodynamic compromise resulting from the tachycardia. Although there are some reservations about RF ablation in infants, numerous studies have shown its effectiveness and its early and medium term safety, justifying its use in certain clinical situations.

In this study we examined the pattern of recovery of left ventricular dysfunction, including the time scale to normalisation, using echocardiography in nine infants and children who had RF ablation for incessant supraventricular tachycardia.

**Methods**

**Patients**

Between January 1992 and October 1996, nine children (five boys and four girls) presented with incessant tachycardia, which was unresponsive or only partially controlled by antiarrhythmic drugs (table 1). The age at presentation ranged from 1 day to 12.5 years (mean 2.3 years). Seven of the nine patients had left ventricular dysfunction; six of these also had left ventricular dilatation. The two with normal systolic function may have had diastolic dysfunction, as all the children had symptoms or signs of heart failure.

The type of tachycardia was diagnosed on the surface electrocardiogram, and the incessant or persistent nature confirmed on ward or Holter monitoring. Patients were subdivided into two groups: group I consisted of four children aged less than 18 months at the time of ablation; group II included the five older children. One patient had focal atrial tachycardia (FAT). Eight had the permanent form of junctional reciprocating tachycardia (PJRT).

**Group I**

All four children in this group had PJRT with cycle lengths between 280 and 360 ms. The
mean interval between diagnosis and ablation was 3.5 months (range from antenatal to 7 months). The age range at ablation was 2 to 17 months (mean 6 months). The three youngest patients (cases 2, 8, and 9) were treated, unsuccessfully, with up to three different antiarrhythmic drugs (including digoxin, verapamil, propranolol, sotalol, and amiodarone) and were therefore subjected to RF ablation. The mean age at ablation for these three patients was 2.5 months and the mean weight 4.7 kg. The fourth patient in this group (patient 1) presented in late infancy with heart failure. Early arrhythmia recurrence required a further seven months of medical treatment, at which point a second RF ablation was successful. All were treated with antiarrhythmic drugs.

**Group II**

Group II comprised four children with PJRT with cycle lengths from 360 to 440 ms and one child with FAT. The mean interval between the diagnosis of tachycardia and ablation was 35.8 months (range 2 weeks to 61 months). Their ages at the time of ablation ranged from 4.8 years to 12.5 years (mean 6.9 years). None was controlled with antiarrhythmic drugs and all received antiarrhythmic drugs.

### Table 1  Patient data

<table>
<thead>
<tr>
<th>Patient</th>
<th>Sex</th>
<th>Age at diagnosis</th>
<th>Presentation</th>
<th>Diagnosis</th>
<th>Age at ablation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>M</td>
<td>10 months</td>
<td>CHF</td>
<td>PJRT</td>
<td>17 months</td>
</tr>
<tr>
<td>2</td>
<td>F</td>
<td>Antenatal</td>
<td>Fetal hydrops</td>
<td>PJRT</td>
<td>2.5 months</td>
</tr>
<tr>
<td>3</td>
<td>F</td>
<td>1 month</td>
<td>CHF</td>
<td>PJRT</td>
<td>5 years</td>
</tr>
<tr>
<td>4</td>
<td>M</td>
<td>12 years</td>
<td>CHF</td>
<td>PJRT</td>
<td>12 years</td>
</tr>
<tr>
<td>5</td>
<td>M</td>
<td>5 months</td>
<td>CHF</td>
<td>PJRT</td>
<td>5 years 6 months</td>
</tr>
<tr>
<td>6</td>
<td>M</td>
<td>1 month</td>
<td>CHF</td>
<td>PJRT</td>
<td>4 years 10 months</td>
</tr>
<tr>
<td>7</td>
<td>F</td>
<td>7 years</td>
<td>CHF</td>
<td>FAT</td>
<td>7 years</td>
</tr>
<tr>
<td>8</td>
<td>F</td>
<td>1 month</td>
<td>CHF</td>
<td>PJRT</td>
<td>2.5 months</td>
</tr>
<tr>
<td>9</td>
<td>M</td>
<td>Antenatal</td>
<td>Tachycardia</td>
<td>PJRT</td>
<td>2 months</td>
</tr>
</tbody>
</table>

CHF, congestive heart failure; FAT, focal atrial tachycardia; PJRT, permanent junctional reciprocating tachycardia.

## Statistical Analysis

All values were expressed as mean (SD). The comparison between echocardiographic measurements and between groups on the basis of recovery time was performed using the Student’s t test; z scores were calculated to express enlargement of left ventricular systolic and diastolic dimensions compared with normal values and to assess recovery of these variables independently of changes in body weight. The z score was calculated as [observed dimension − mean normal dimension] ÷ standard deviation around the mean dimension.

### Results

#### Outcome of Ablation

Ten ablation procedures were performed on the nine children. In the patient with FAT, the earliest intracardiac atrial activation was in the roof of the right atrium and measured −29 ms compared with the surface P wave. In the PJRT group, the tachycardia cycle length and the RP and PR intervals were 360 (52.9) ms, 227 (19.2) ms, 132 (40.8) ms, respectively, and the
shortest VA interval during tachycardia was in the right posterior or right posteroseptal region.

One child required a repeat procedure seven months later for early recurrence. Complete AV block occurred in two patients, transiently in one (lasting 20 seconds) and persistent in the other (the patient was aged 2.5 months at ablation), requiring permanent transvenous pacing. No other complications were encountered.

FOLLOW UP

Eight patients were in sinus rhythm and one in paced rhythm following RF ablation. The mean (SD) follow up period from the time of ablation was 13.4 (10.6) months for group I and 47.4 (10.7) months for group II.

LEFT VENTRICULAR DYSFUNCTION:

ECHOCARDIOGRAPHIC EVALUATION

Seven of the nine children (77.8%) had left ventricular dysfunction, all of whom recovered following successful ablation. Six of these also had left ventricular enlargement (group I 50%; group II 80%), all of whom improved, four with full recovery at the time of writing. Two patients, one with fetal hydrops and another presenting with heart failure in infancy, had normal ventricular size and function before ablation, despite heart failure symptoms. These two were excluded from statistical analysis of recovery time in view of their normal preablation values.

Following RF ablation, left ventricular ejection fraction increased from 42.6(17.9)% to 71.1(3.0)% (p < 0.003) and shortening fraction from 19.8(8.9)% to 35.2(3.15)% (p < 0.002) after a mean period of 20.7 months (fig 1A and B). Subgroup analysis showed a mean recovery time of 5.7 (7.2) months for group I and 31.3 (5.2) months for group II (p < 0.002) (fig 2).

The measured LVESD and LVEDD, which were correlated with body weight, were expressed as z scores to take into account the expected changes in left ventricular size with the patient’s growth. Figure 3 shows the z score changes before and at a mean of 20.7 months after the catheter ablation. The pre- and postablation z scores were 2.97 (2.69) and 0.5 (0.73) for LVESD (p < 0.02) and 2.41 (2.23) and 0.76 (1.3) for LVEDD (p < 0.07, NS).

Figure 2 Time interval to recovery of left ventricular ejection fraction after radiofrequency ablation in seven children with left ventricular dysfunction.

Figure 3 (A) Left ventricular end diastolic dimension before and after radiofrequency ablation, expressed in z scores. (B) Left ventricular end systolic dimension before and after radiofrequency ablation, expressed in z scores.

Discussion

BACKGROUND

Incessant tachycardia is a well recognised cause of “cardiomyopathy,” consisting of left ventricular enlargement and impairment of systolic function. This phenomenon is particularly apparent in children in whom specific types of incessant arrhythmias, for example PJRT/FAT, are usually present in otherwise structurally normal hearts.

CLINICAL AND EXPERIMENTAL DATA

Several clinical and experimental studies have shown a causal link between the heart rate and left ventricular dysfunction. Tachycardia cycle length and duration of the arrhythmia contribute to left ventricular dysfunction. Many investigators have also observed regression of the cardiomyopathy following normalisation of heart rate in patients whose arrhythmia was either controlled or cured. There have been similar findings in experimental models (dogs and pigs) using fast atrial pacing. Damiano et al, using radiouclide angiography, reported significant reduction in ejection fraction in 13 dogs that were paced for three months at a rate of 190 beats/min. After eight weeks of pacing, there was no significant change in cardiac output or stroke volume but there was marked increase in left ventricular dimension to compensate for the reduced contraction. Moreover, five of the 13 dogs were followed up for eight weeks after the pacing period and these showed recovery of ejection fraction but left ventricular dimension remained abnormal, indicating secondary changes which do not recover immediately on control of tachycardia. Coleman et al showed...
depletion of myocardial energy stores in paced dogs, giving a plausible explanation for the left ventricular dysfunction.24 Similar studies on pigs25–27 confirmed the haemodynamics but also showed histological and biochemical consequences of persistent tachycardia both on immature and fully grown animals. In these pig studies, persistence of left ventricular dilatation and left ventricular diastolic dysfunction following cessation of pacing indicate potentially serious changes to the myocardium, even though systolic function returned to normal and an early cure for incessant tachycardia would seem appropriate to preserve the myocardium.

CURRENT STUDY: RECOVERY OF CARDIOMYOPATHY
Systolic dysfunction is the first manifestation of tachycardia cardiomyopathy, followed by left ventricular dilatation. Two toddlers in group I who had impaired ventricular function before ablation did not show left ventricular dilatation on the echocardiogram, whereas four of five children in group II had left ventricular dilatation; the only one in group II (patient 6) who did not show left ventricular enlargement had persistent tachycardia—that is, intermittent rather than incessant. This suggests that the longer the duration of the tachycardia, the more damaging the effect on the myocardium.

This study confirms that curing the tachycardia, in our case using RF ablation, results in resolution of the cardiomyopathy. Left ventricular systolic function, as assessed by echocardiography, recovered fully in all seven who had dysfunction before ablation. The time scale for recovery varied considerably between groups I and II, implying that the older children had worse “myocardial damage” (very likely a reflection of longer duration of tachycardia). Two children in group II (patients 4 and 5) who were initially diagnosed as suffering from primary cardiomyopathy were successfully treated for incessant tachycardia, resulting in recovery of systolic function. This recovery took up to three years; cardiac enlargement, however, persisted despite marked improvement in LVEDD. On the other hand, two infants in group I (patients 8 and 9) who had left ventricular dysfunction but with no cardiac enlargement, showed normalisation of ejection fraction and fractional shortening within weeks of successful ablation. Thus the more advanced the cardiomyopathy, the slower and more incomplete was the recovery.

AVAILABLE TREATMENT

Ventricular dysfunction caused by incessant tachycardia does not usually improve with medical treatment.8 11 Curative procedures have included catheter ablation, initially using direct current and, more recently, radiofrequency or surgery. RF ablation is a recognised therapeutic curative technique for many tachycardia substrates, with a high success rate and few complications even in children.11 14–16 Although complications have occurred, the learning curve for this relatively new technique must be taken into account. An injudicious number of radiofrequency applications, however, can inevitably lead to serious complications.9 10 There have only been isolated case reports of successful catheter ablation for incessant tachycardia in children with recovery of left ventricular function,11 and only one study addressing recovery time.12 The potential long term adverse effects of RF ablation have to be balanced against the recognised effects of incessant tachycardia on left ventricular function. It was technically feasible to perform the ablation in very young children whose average weight was 4.7 kg, and with an average of 1.5 radiofrequency applications. Complete AV block is undoubtedly a major complication. In our patient who needed a pacemaker, PIR was abolished within three seconds of the first radiofrequency application, but junctional tachycardia led to catheter movement resulting in AV block. The use of temperature controlled ablation catheters, back up pacing, or cessation of energy delivery during junctional tachycardia may reduce this risk. None of the patients has had a recurrence of the arrhythmia over a modest follow up period, and all are asymptomatic.

CONCLUSION

Incessant tachycardia frequently causes “cardiomyopathy,” resulting in left ventricular dysfunction followed by cardiac enlargement. These changes are reversible following cure of the tachycardia; however, the recovery time is related to the duration of the tachycardia, is significantly shorter in infants, and cardiac enlargement takes longer to resolve than cardiac function. RF ablation provides an effective cure for children with incessant supraventricular tachycardia and should be resorted to at an early stage, particularly if there is left ventricular dysfunction, in order to minimise myocardial damage from the arrhythmia.


**IMAGES IN CARDIOLOGY**

**Congenital morphologies of the aortic valve**

A collection of the different congenital morphologies of the aortic valve seen by echocardiography. From right to left: monocuspid, bicuspid, tricuspid, and multicuspid valves are shown with systolic view on the upper and diastolic view on the lower side.

M ZUBER
R JENNI
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