Cryoablation time-dependent dose–response effect at minimal temperatures (−80°C): an experimental study

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Aims
To establish a temporal safety window for cryoablation at minimal temperatures and to assess the electrophysiological and histological changes as a function of the application duration.

Methods and results
Twenty mini-pigs underwent AV nodal cryoablation at −80°C without prior cryomapping. The duration of the cryoapplication following atrioventricular block (AVB) was randomized to 0, 10, 20, 40, or 60 s. Atrioventricular block was obtained in all animals after a median of 3 (1–8 interquartile range) applications. One week later, AV nodal conduction fully recovered in animals with application duration ≤10 s, whereas persistent AVB incidence increased as a function of time in animals with longer applications duration. Cryoablation application duration following AVB was the only independent predictor of persistent AVB (OR, 1.116; 95% CI, 1.013–1.229; P = 0.026). There was no difference in lesion location or size between animals with vs. those without persistent AVB at 1 week. However, animals randomized to longer application duration demonstrated higher degree of cell destruction and fibrotic content.

Conclusion
In this closed-chest pig model, there was a relation between cryoapplication duration following AVB at −80°C and recovery of conduction. A safety window of at least 10 s was observed in all cases.

Keywords
Cryothermal ablation • Atrioventricular block • Perinodal ablation • Atrioventricular node • Supraventricular tachycardias

Introduction
Cryothermal ablation is increasingly used in high-risk perinodal ablations, due to the absence of undesired persistent atrioventricular block (AVB) cases reported in the literature.1 The ability of this technology to create reversible lesions at temperatures of −30°C (cryomapping) is responsible in part for this excellent safety profile.1–9 However, the electrophysiological effects may not always be predicted by prior cryomapping at −30°C and adverse effects can subsequently be observed during cryoablation (at −70° to −80°C).2–9 In this case, anecdotal reports suggest that, if the application is immediately interrupted, there is usually complete recovery of conduction.2–9 We hypothesized that the reversibility of cryothermal application at the AV node is dependent, not only on the minimal temperature at the catheter tip, but also on the duration of the application. Therefore, the
objectives of this study were to: (i) establish a temporal safety window during AV nodal cryoablation at minimal temperatures (−80°C); (ii) evaluate the residual electrophysiological AV nodal impairment after reversible lesions at −80°C; and (iii) assess the histological changes related to the occurrence of AVB as a function of the duration of the cryoablation application.

Methods
The experimental protocol was approved by the Ethics Committee of Animal Experimentation of our institution. Twenty mini-pigs of either sex weighing 30 ± 4 kg were used in this study.

Electrophysiological study
Three quadripolar catheters were positioned using fluoroscopy to the right atrium, the His bundle, and the right ventricular apex for recording and pacing. Surface electrocardiographic leads and filtered bipolar intracardiac electrograms were recorded at 100 mm/s paper speed. Atrial and ventricular stimulation protocols included pacing trains at a fixed cycle length and the extrastimulus technique.

Atrioventricular conduction properties were assessed at baseline prior to cryoenergy application, 1 h post-ablation and 1 week later. The following parameters were recorded: AH and HV intervals, AV nodal Wenckebach cycle length, AV nodal effective refractory period.

Cryoaulation protocol
Percutaneous AV nodal cryoablation was performed using a 9 Fr catheter (Freezor® Max, CryoCath Technologies Inc., Montreal, Canada). The cryocatheter has a distal 8 mm tip cooling electrode and three proximal ring electrodes with 3.5–2 spacing. The principles of the cryoablation system have been described in detail.1,4,10 The cryoablation catheter was positioned at the triangle of Koch and cryoenergy at minimal temperature of −80°C (without prior cryomapping) was delivered at sites with large low right atrial and small His bundle recordings, aiming to achieve AVB, defined as the occurrence of two consecutive P-waves not conducted to the ventricle. Prior to energy delivery, the duration of the cryoablation following AVB was randomized to 0, 10, 20, 40, or 60 s (Figure 1). The randomization table was designed to obtain equally distributed groups (Group 1: 0 s, Group 2: 10 s, Group 3: 20 s, Group 4: 40 s, Group 5: 60 s). The cryoapplication was maintained until the temperature reached −80°C and stopped if no AVB was obtained 30 s after the minimal temperature was reached. In that case, the catheter was allowed to rewarm and was repositioned to perform another cryoapplication. Afterwards, a permanent pacemaker was implanted and animals were kept alive for 1 week. The outcome variables after cryoablation were: (i) acute AVB: complete AV 1 h after the cryoapplication delivery; (ii) acute AV conduction recovery: 1:1 atrial conduction to the ventricle up to 1 h after the cryoapplication delivery; (iii) delayed AV conduction recovery: AV conduction recovery that occurred between the end of the procedure up to 1 week later; and (iv) persistent AVB: complete AVB 1 week after the cryoablation procedure. The following procedural variables corresponding to the effective cryoapplication (i.e. causing AVB) were recorded.

- Time to effect: interval from onset of cryoablation application to AVB occurrence.
- Temperature at AVB: catheter tip temperature at the time of AVB occurrence.
- Nadir temperature: minimal temperature reached during the application.
- Time to cryoablation temperature: interval from onset of cryoablation application to catheter tip temperature ≤ −80°C.
- Critical cryoablation application duration: cryoablation application duration interval from onset of AVB until the end of the application.

Histology
Animals were sacrificed at 7 days with a lethal injection of pentobarbital. Their hearts were explanted, rinsed, and fixed in 10% formalin. At least four blocks of tissue of the AV junctions were dehydrated and embedded in paraffin. Specimens were serially sectioned perpendicular to the septal leaflet of the tricuspid valve at a thickness of 7–10 µm with a microtome (Microm HM310) and stained with Masson’s trichrome. Cryolesions size was measured using image analysis software (SigmaScanPro 5.0, Jandel Scientific, San Rafael, CA, USA). The geometry of the cryoablation lesions’ area was calculated as $A = (1/4)\pi(W/2)D$, and the volume was calculated as $V = (1/2)(4/3)\pi(W/2)^2D$, where $W$, lesion width and $D$, lesion depth.11

Morphometric analysis at light microscopy was restricted to experiments in which AV conduction block was obtained after a single cryoapplication and to controls. It was performed using digital images (512 x 512 pixels) from each tissue block by using a linear measurement software program (SigmaScan Pro 5.0). Twenty-two measurements of AV nodal and His myocytes diameter (μm) were obtained from each tissue block. For each measurement, the widest cross-sectional myocyte diameter was selected and averaged to allow comparison with controls. To quantify the amount of myocyte/connexitive tissue, morphometry was performed using a grid of vertical and horizontal lines providing 121 intersections or points. The total number of intersection points was defined as 100%. The proportion of tissue corresponding to connective tissue matrix in each specimen was expressed as connective tissue density, defined as the percentage of intersection points overlying within the tissue extension limited by the grid. Connective tissue density was estimated in a minimum of eight different randomly chosen fields of the AV node and His bundle for each specimen and values were then averaged. Blood vessels and perivascular interstitial tissue were excluded from connective tissue quantification. To quantify the extent of myolysis in the specialized tissue, only myocytes in which the nucleus was present in the plane of the section were considered and at least 12 myocytes per section were analysed. Myocytes were scored by morphometry as: (i) normal myocytes, without myolysis, (ii) mildly myolytic if myolysis involved between 15 and 35% of the sarcoplasm, and (iii) severely myolytic if >40% of the sarcomeres were absent.

In four animals, an electrophysiological study was performed but no cryoapplications were delivered and they were used as control specimens.

Statistical analysis
Results are reported for the entire sample ($n = 20$) unless otherwise specified. Continuous variables are reported as mean ± SD or median and interquartile range (IQR), depending on whether they were normally or non-normally distributed. Categorical variables are reported as number and percentage. Comparisons were made using Student’s t-test and the Fisher’s exact test, as appropriate. In case of non-normally distributed continuous variables, Wilcoxon test was used. Statistically significant variables in the univariate analysis were included in a stepwise logistic regression model. Analyses were performed with SPSS v14.0 and statistical significance was established at $P < 0.05$. 
Results

A total of 107 cryoablation applications were delivered at the level of the AV node. Atrioventricular block was obtained in all animals after a median of 3 (1–8 IQR) cryoablation applications. A single application was needed to obtain AVB in six (30%) animals, two to four in six (30%), and five or more applications in the remaining. The mean minimal temperature recorded during the applications that caused AVB was $-82.3 \pm 0.5^\circ$C. The median temperature at AVB occurrence was $-80^\circ$C (range $-50$ to $-85^\circ$C). The mean time to effect was $31.8 \pm 18.7$ s and the mean time to cryoablation temperature $\leq -80^\circ$C was $22.2 \pm 4.1$ s. Mean application duration according to assigned group was: Group 1: $3.75 \pm 2.22$, Group 2: $14 \pm 1.82$, Group 3: $22.5 \pm 3.32$, Group 4: $41.5 \pm 12.37$, Group 5: $65 \pm 6$ s.

Atrioventricular conduction acutely recovered in 10 animals after a median of 55 (range 30–1500) seconds following AVB. Two additional animals with acute AVB, had delayed AV conduction recovery 1 week after the ablation procedure. There were no cases of delayed AVB in the group of animals with acute AV conduction recovery following cryoablation. Thus, eight animals had persistent AVB 1 week after the procedure.

Predictors of acute atrioventricular conduction recovery

Atrioventricular nodal conduction fully recovered in all animals randomized to cryoablation application duration of 0 and 10 s following the second-blocked P-wave. No cases of acute AVB occurred with applications duration lasting $\leq 15$ s. In univariate analyses, several variables were associated with the occurrence of acute conduction recovery, whereas lesion size was not (Table 1). In multivariate analysis, critical cryoablation application duration from AVB onset remained as the only significant predictor of acute AV conduction recovery (OR, 0.887; 95% CI, 0.799–0.984; $P = 0.024$).

Predictors of persistent atrioventricular block occurrence

One week after the cryoablation procedure, there were no cases of persistent AVB in animals randomized to application duration of 0 and 10 s following the second-blocked P-wave, whereas the probability of presenting persistent AVB increased as a function of time in animals with longer application duration. Persistent AVB occurred in one (25%), three (75%), and four (100%) animals assigned to critical cryoapplication duration of 20, 40, and 60 s, respectively. The first case of persistent AVB occurred after a 16 s application duration following the second-blocked P-wave. Several variables were related to the occurrence of persistent AVB in the univariate analyses (Table 1). In a multiple logistic regression model, critical application duration from AVB onset was the only independent predictor of persistent AVB (OR, 1.116; 95% CI, 1.013–1.229; $P = 0.026$).
Atrioventricular nodal conduction characteristics following transient atrioventricular block

A significant prolongation of the AH interval was the only electrophysiological parameter that was affected by the cryoablation application in animals with AV nodal conduction recovery (Table 2). Other parameters of AV nodal function fully normalized at 60 min and at 7 days after cryoablation.

Histology

On qualitative analysis, all cryoablated specimens were consistently characterized by extensive myocardial damage involving the AV junction walls. Cryolesions were well-circumscribed discrete lesions with sharp borders, dense areas of fibrotic tissue, and contraction band necrosis (Figure 2). Myocardial tissue displayed shrunk cells with degenerated cytoplasm, decreased staining, coagulation necrosis, lymphocytes, and macrophages infiltration (Figure 3). Overall (n = 20), there was no difference in lesion size between animals with vs. without AVB acutely or 1 week after the cryoablation procedure (Table 1).

In six animals, AV conduction block was obtained after a single cryoapplication. Of those, AV nodal conduction acutely recovered in animals randomized to application duration of 0 and 10 s (n = 2), whereas AVB persisted >1 h after the procedure in the remaining animals randomized to 20–60 s application duration (Table 3). Histologically, all six animals demonstrated a lesion that affected the proximal AV conduction system, both the penetrating bundle and a portion of the AV node. There was no difference in lesion location at the AV node level or in lesion size between animals with AV nodal conduction recovery vs. persistent AVB at 1 week (Figure 2). Morphometric analysis of the cryolesions showed loss of cellular viability throughout the wall thickness. However, in animals that subsequently had acute AV conduction recovery (#6, #7), isolated viable cells surrounded by necrotic tissue were observed within the lesions of the specialized AV node and penetrating His bundle (Figure 3). Moreover, the degree of myolytic changes differed between animals with vs. without persistent AVB at 1 week (Table 3 and Figure 3). Animals with AV conduction recovery showed only mild myolytic changes, whereas severe myolytic changes (>40% myolytic fibres), both at the AV node level and the penetrating His bundle, were observed in animals with persistent AVB (#3, #19, #22). Similarly, animals with persistent AVB at 1 week demonstrated higher collagen content when compared with animals with AV

### Table 1 Univariate predictors of acute atrioventricular nodal conduction recovery and persistent atrioventricular block at 1 week

<table>
<thead>
<tr>
<th>Predictor</th>
<th>Yes (n = 10)</th>
<th>No (n = 10)</th>
<th>P-value</th>
<th>Yes (n = 12)</th>
<th>No (n = 8)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of applications</td>
<td>6 (11)</td>
<td>3 (3)</td>
<td>0.131</td>
<td>3.5 (10)</td>
<td>3 (6)</td>
<td>0.635</td>
</tr>
<tr>
<td>Time to effect</td>
<td>40 (32)</td>
<td>19 (18)</td>
<td>0.016</td>
<td>34.5 (31)</td>
<td>19 (24.3)</td>
<td>0.031</td>
</tr>
<tr>
<td>Time to cryoablation temperature</td>
<td>21.55 ± 5.14</td>
<td>23 ± 4.27</td>
<td>0.507</td>
<td>21.83 ± 5.32</td>
<td>22.75 ± 3.88</td>
<td>0.662</td>
</tr>
<tr>
<td>Critical cryoablation application duration</td>
<td>13 (18)</td>
<td>49 (42.5)</td>
<td>0.003</td>
<td>14 (17.75)</td>
<td>48 (33.25)</td>
<td>0.010</td>
</tr>
<tr>
<td>Temperature at AVB</td>
<td>−80 (0)</td>
<td>−80 (13)</td>
<td>0.045</td>
<td>−80 (0)</td>
<td>−78.5 (14)</td>
<td>0.037</td>
</tr>
<tr>
<td>End of lesion</td>
<td>−83 (5)</td>
<td>−82 (3.5)</td>
<td>0.614</td>
<td>−82.5 (4.5)</td>
<td>−82 (4.25)</td>
<td>0.694</td>
</tr>
<tr>
<td>Lesion area</td>
<td>2.95 (0.7)</td>
<td>3 (1.85)</td>
<td>0.682</td>
<td>2.9 (0.5)</td>
<td>3.3 (2.35)</td>
<td>0.281</td>
</tr>
<tr>
<td>Lesion volume</td>
<td>16.05 ± 7.67</td>
<td>14.7 ± 8.56</td>
<td>0.724</td>
<td>14.94 ± 6.85</td>
<td>16.06 ± 9.63</td>
<td>0.768</td>
</tr>
<tr>
<td>Lesion volume</td>
<td>39.45 ± 22.67</td>
<td>40.91 ± 24.63</td>
<td>0.894</td>
<td>40.38 ± 23.15</td>
<td>40 ± 24.56</td>
<td>0.973</td>
</tr>
</tbody>
</table>

Mean ± SD; median (interquartile range).

### Table 2 Atrioventricular nodal conduction characteristics before and following cryoablation in animals with atrioventricular nodal conduction recovery

<table>
<thead>
<tr>
<th>Predictor</th>
<th>Baseline</th>
<th>Acute (n = 10)</th>
<th>P-value&lt;sup&gt;a&lt;/sup&gt;</th>
<th>1 Week (n = 12)</th>
<th>P-value&lt;sup&gt;b&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>AH interval</td>
<td>28.1 ± 6.5</td>
<td>32.3 ± 6.8</td>
<td>0.04</td>
<td>32.3 ± 6.5</td>
<td>ns</td>
</tr>
<tr>
<td>HV interval</td>
<td>32.8 ± 6.8</td>
<td>32.8 ± 6.8</td>
<td>ns</td>
<td>32.3 ± 6.5</td>
<td>ns</td>
</tr>
<tr>
<td>Wenckebach cycle</td>
<td>271.1 ± 38.9</td>
<td>268.9 ± 50.8</td>
<td>ns</td>
<td>259.5 ± 43.5</td>
<td>ns</td>
</tr>
<tr>
<td>AVN refractory period 600</td>
<td>297.8 ± 33.1</td>
<td>297.8 ± 33.1</td>
<td>ns</td>
<td>283.6 ± 38.3</td>
<td>ns</td>
</tr>
<tr>
<td>AVN refractory period 400</td>
<td>245 ± 30.1</td>
<td>223.7 ± 93.5</td>
<td>ns</td>
<td>251 ± 28.1</td>
<td>ns</td>
</tr>
</tbody>
</table>

<sup>a</sup>Baseline vs. acute.

<sup>b</sup>Baseline vs. 1 week.
conduction recovery and controls. Interestingly, case #17 had delayed AV conduction recovery at 1 week and showed intermediate myolytic changes and fibrotic content level when compared with animals with acute recovery and persistent AVB. Finally, a progressive increase in cell size was observed from controls to persistent AVB animals.

**Discussion**

The main findings of the present study are that in this closed-chest pig model: (i) there is a safety window of at least 10 s during cryoablation at minimal temperatures (−80°C), where complete recovery of the AV node electrophysiological properties is always observed; (ii) the main determinant of the electrophysiological effects of cryoablation of the AV node at minimal temperatures is application duration, but not lesion size; (iii) changes at the microscopic level, such as cell destruction and fibrosis, explain the irreversible electrophysiological effects demonstrated by applications duration lasting >15 s.

**Undesired atrioventricular block during cryoablation at −80°C**

As opposed to radiofrequency ablation, there are no cases reported in the literature of undesired persistent AVB during cryoablation. For this reason, cryoablation has become a very attractive alternative to radiofrequency during perinodal ablations, particularly in young individuals in whom AVB would be a disastrous outcome. In this setting, cryomapping at −30°C creates reversible lesions that enable testing the functional effects of ablation before the formation of a permanent lesion. However, the electrophysiological effects may not always be predicted by prior cryomapping at −30°C and adverse effects can be observed during cryoablation (at −70 to −80°C). We and others have reported the occurrence of transient high-grade AVB during cryoablation at sites where previous cryomapping was judged to be ‘safe’ as verified using a navigation system. Although gradual normalization of the electrophysiological properties has been observed when the application is immediately interrupted, no information exists to date regarding the safety profile of cryoablation in this situation.

**Application duration as a determinant of the electrophysiological effect**

Permanence of tissue destruction with cryothermal energy is related to time-dependent variables and tissue temperature. Holman et al. first reported the occurrence of undesired AVB during selective cryosurgical ablation (−60°C) of the perinodal tissue that completely resolved over time if the cryothermic exposure was immediately terminated. They also observed that despite prolongation of AH and VA intervals, Wenckebach point, and AV nodal effective refractory period in the immediate post-operative period, all these variables normalized by 14 days after surgery except for the AH interval. In our study, only a significant prolongation in the AH interval following transient AVB at −80°C was observed acutely and 1 week later. Also in agreement with them, no instances of persistent AVB were observed during follow-up in our study after applications lasting <16 s.

Several clinical studies in humans have reported the occurrence of undesired transient AVB during cryoablation applications at temperatures ≤−70°C. In all cases AVB resolved completely, usually within a few seconds or minutes, with PR interval and Wenckebach point returning to baseline. In those studies, no patient had persistent AVB or required pacemaker implantation. The present study confirms the benign outcome of undesired complete AVB that occurs during cryoablation at −80°C. Moreover, the observation that there were two cases of delayed AV conduction recovery and no cases of AV nodal conduction impairment 1 week following cryoablation suggests that long-term deterioration is unlikely. However, although either acute or delayed AV nodal conduction recovery can be observed with applications lasting >15 s, the probability of persistent AVB occurrence significantly increased over that application duration, indicating...
that, in fact, permanent AV nodal block occurs with long enough cryoenergy applications. Other authors have failed to obtain persistent AVB in a significant proportion of patients with clinical indication of AV node ablation, despite delivering long duration cryoenergy applications preceded by multiple cryomapping attempts. Several circumstances may explain their results, such as smaller catheter tip size (4–6 mm), catheter tip–heart size mismatch, tissue oedema as a consequence of multiple consecutive cryomapping and cryoablation attempts, and higher temperatures during cryoablation. Our results confirm the ability of long enough cryoablation applications (>60 s) to consistently produce persistent AVB when large catheters (8 mm) are used and very low temperatures (<−80 °C) are reached.

**Characteristics of the cryolesion**

Several hypotheses have been proposed to explain the reversibility of the electrophysiological effects during cryoablation such as iceball enlargement and temperature gradient effect on tissues in the proximity of the catheter tip, but none has systematically analysed the histological correlates of reversibility of the electrophysiological effects. In the present in vivo study using cryoablation at minimal temperatures (−80 °C), we found no differences in lesion location or in lesion size between animals with vs. those without persistent AVB, application duration being the only independent predictor of AV nodal conduction recovery. Separate analysis of experiments in which AVB was obtained with a single application confirmed these results. At minimal temperatures, cryothermal energy created a well-defined lesion with sharp borders displaying the characteristic features previously described (Figure 2). We performed morphometric analysis of the lesions to get further insights into the factors determining normalization of the electrophysiological conduction properties. The degree of myolytic changes and collagen content of the lesion increased as a function of the critical cryoapplication duration, thereby determining the electrophysiological effect. Interestingly, strands of isolated viable cells surrounded by necrotic tissue were observed within the lesions of the specialized conduction system in animals that subsequently showed AV conduction recovery (Figure 3). Similar findings were observed by Fujino et al. in animals demonstrating late AV conduction recovery at 6 weeks following percutaneous AVB cryoablation attempts. Thus, applications lasting <15 s from the electrophysiological effect onset create a well-defined lesion, but enable some viable cells to survive. However, the long-term outcome of these lesions remains unknown and studies are needed to establish the risk of delayed (>1 week) AVB development under these conditions.
<table>
<thead>
<tr>
<th>Experiment #</th>
<th>Randomized group assignment</th>
<th>Critical application duration (s)</th>
<th>AVN conduction after ablation</th>
<th>Myocyte diameter (µm)</th>
<th>CTD/field (%)</th>
<th>Myolyis AV node/field (%)</th>
<th>Myolyis His bundle/field (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Controls n = 4</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>7.5 ± 1.2 (5–9)</td>
<td>20 ± 4 (24–35)</td>
<td>8 ± 2 (5–12)</td>
<td>7 ± 3 (3–12)</td>
</tr>
<tr>
<td>#7</td>
<td>Group 1</td>
<td>1</td>
<td>Acute AV conduction recovery</td>
<td>12 ± 2 (7–18)</td>
<td>35 ± 4 (28–45)</td>
<td>21 ± 4 (16–29)</td>
<td>25 ± 4 (18–35)</td>
</tr>
<tr>
<td>#6</td>
<td>Group 2</td>
<td>15</td>
<td>Acute AV conduction recovery</td>
<td>13 ± 2 (8–18)</td>
<td>30 ± 4 (23–46)</td>
<td>25 ± 4 (19–33)</td>
<td>22 ± 3 (18–40)</td>
</tr>
<tr>
<td>#17</td>
<td>Group 3</td>
<td>23</td>
<td>Delayed AV conduction recovery</td>
<td>15 ± 2 (9–20)</td>
<td>40 ± 4 (33–50)</td>
<td>35 ± 7 (25–48)</td>
<td>32 ± 4 (26–43)</td>
</tr>
<tr>
<td>#3</td>
<td>Group 3</td>
<td>20</td>
<td>Persistent AVB</td>
<td>15 ± 2 (10–22)</td>
<td>58 ± 4 (48–75)</td>
<td>65 ± 5 (55–85)</td>
<td>68 ± 5 (55–95)</td>
</tr>
<tr>
<td>#19</td>
<td>Group 5</td>
<td>60</td>
<td>Persistent AVB</td>
<td>17 ± 2 (9–24)</td>
<td>62 ± 4 (50–80)</td>
<td>60 ± 5 (55–85)</td>
<td>66 ± 5 (55–98)</td>
</tr>
<tr>
<td>#22</td>
<td>Group 5</td>
<td>60</td>
<td>Persistent AVB</td>
<td>16 ± 2 (9–20)</td>
<td>65 ± 5 (50–83)</td>
<td>77 ± 5 (65–98)</td>
<td>85 ± 5 (72–98)</td>
</tr>
</tbody>
</table>

Mean ± SD (range). AVB, atrioventricular block; CTD, connective tissue density; NA, not applicable.

**References**


**Limitations**

Several limitations in our work should be discussed. First, lesion size can depend on several conditions such as electrode orientation, contact pressure, and blood velocity. Second, more than one cryoablation application was needed to obtain AVB in two-thirds of the experiments. To avoid bias caused by the possibility of lesion confluence and the effect of consecutive freeze–thaw cycles, morphometric analysis was limited to experiments in which AVB was obtained with a single energy application controlled for in the present study. 8 For this reason, application duration was randomly assigned to partially overcome this limitation. Second, more than one cryoablation application was needed to confirm the favourable outcome of reversible lesions created at minimal temperatures in the model.

**Clinical implications**

More aggressive strategies are used with the aim of increasing success rates and reducing recurrences during cryoablation. In patients with accessory pathways, it has been suggested that substituting the tip cryoablation catheters are more likely to achieve a high success rate. Our results suggest that even persistent AVB could be expected during perinodal cryoablation. A similar approach has been suggested to address the need for more aggressive strategies, as these might be useful in further defining the role of this energy for ablation of accessory substrates at high risk of AVB.

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**Conflict of Interest**

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