Transcoronary Chemical Ablation of Atrioventricular Conduction

Pedro Brugada, MD, Hans de Swart, MD, Joep Smeets, MD, and Hein J.J. Wellens, MD

In seven patients with symptomatic atrial fibrillation and uncontrollable ventricular rates, selective catheterization of the atrioventricular (AV) nodal artery was performed to chemically destroy the AV node. Ethanol at a concentration of 96% and a dose of 0.5–2 ml was used after selective catheterization of the AV nodal artery had demonstrated temporary AV block after the administration of isotonic iced saline. Complete AV block was produced in five patients and AV conduction was sufficiently modified to control symptoms in the remaining two patients. A minimal enzyme rise occurred in six patients. A severe complication in the remaining patient occurred when, after 2 ml ethanol in the AV nodal artery, occlusion developed in the midnight coronary artery that led to an inferior wall myocardial infarction. It is concluded that the AV nodal artery can be selectively catheterized using presently available angioplasty techniques. Ethanol can be used to destroy the AV node and block AV conduction. (Circulation 1990;81:757–761)

We have recently shown that it is possible to identify and selectively catheterize a small coronary artery providing blood supply to an arrhythmogenic area or pathway.1–2 Using this technique, we have successfully ablated, with 96% ethanol, ventricular tachycardia in patients with incessant ventricular tachycardia after myocardial infarction.2 In our previous report,1 we had selectively catheterized the atrioventricular (AV) nodal artery in one patient having circus-movement tachycardia using the AV node–His pathway and an accessory AV pathway. Administration of iced saline resulted in termination of the arrhythmia by block in AV nodal conduction.

The AV nodal artery arises in most humans from the right coronary artery and is a readily identifiable vessel.3 Electrical ablation of the AV conduction system has been successfully used for several years to control ventricular rates in patients with atrial fibrillation not controllable with antiarrhythmic drugs.4–5 In some patients, however, electrical ablation of the AV conduction system can fail to create AV block or sufficiently modify AV conduction. Using the same techniques as previously described,1–2 we undertook chemical ablation of AV conduction in seven patients with symptomatic uncontrollable ventricular rates during atrial fibrillation. The feasibility of this procedure, but also the occurrence of a severe complication in one patient, are the reasons for this report.

Methods

Seven patients were studied. Their clinical characteristics are summarized in Table 1. All patients had failed multiple antiarrhythm drug trials to control ventricular rates during atrial fibrillation. In four patients, three to seven attempts of electrical ablation of the AV system were performed without lasting success. Three patients had a permanent pacemaker implanted. Possible damage to the pacemaker by renewed attempts of electrical ablation was an additional reason to undertake chemical ablation in these patients. The last two patients gave preference to chemical ablation above electrical ablation or surgery. The last patient had undergone successful surgical ablation of an accessory pathway. All patients had normal coronary arteries and were free from any structural heart disease. The experimental nature of this procedure was explained to all patients, and they gave informed consent. The protocol was approved by the Medical Ethical Committee of our University.

Coronary angiography using the Judkins technique was first performed in all patients. All seven had normal coronary arteries. The AV nodal artery originated from the right coronary artery in all patients except in patient 3 in whom it originated from a dominant circumflex coronary artery. The AV nodal...
artery was a discrete single vessel in five patients but two branches next to each other were observed in patients 2 and 3. Selective catheterization of the AV nodal artery was performed using previously described methodology.\textsuperscript{2,3} In summary, a 0.014-in. angioplasty guide wire was advanced through an 8F guiding catheter as far as possible into the AV nodal artery. Thereafter, a 2.2–4.0F lumen catheter with a metallic marker at the tip was advanced over the guide wire as far as possible into the AV nodal artery. The guide wire was, thereafter, removed. Contrast material was given to confirm catheterization of the vessel identified as the AV nodal artery. During the procedure, an electrode catheter was inserted in the right ventricular apex and connected to an on-demand external pacemaker in all patients. Nitroglycerin was continuously infused intravenously during the procedure. Heparin was also given at a dose of 1 mg/kg body wt (equivalent to 100 units heparin per kilogram body weight). Confirmation of selective catheterization of the AV nodal artery was done by observing the occurrence of transient AV block on selective catheterization of the artery, administration of contrast material, and administration of iced saline. An example is shown in Figure 1.

After confirmation of the identification of the AV nodal artery, 96% ethanol was given at a dose of 0.5–1 ml in 2–4 seconds except in patient 4, who received a total of 3 ml in two sessions. The reasons for the larger dose in the last patient will be discussed later.

After AV block was created, catheters with the exception of the pacing lead were removed, and the patient was transferred to the coronary care unit for 48 hours. A permanent pacemaker was implanted 2–3 days later when required.

### Results

Transient complete AV block occurred on catheterization of the AV nodal artery in all patients. Conduction recovered after a few seconds to minutes in all patients, although ventricular rates remained below the initial heart rates when the catheter was maintained in place. Iced saline and contrast material (Figure 1) resulted in transient complete AV block in all patients.

Administration of 96% ethanol led to complete AV block in all patients. All patients showed ventricular extrasystoles before the development of AV block as illustrated in Figure 1. AV block remained complete in patients 1, 2, 4, 6, and 7, after follow-up of 1–7 months (mean, 4 months). In patients 3 and 5, conduction recovered within 24 hours but the ventricular rate during atrial fibrillation decreased from a mean of 170 to 100 beats/min and from 160 to 80 beats/min, respectively, with pronounced symptomatic improvement after 2 and 5 months' follow-up.

### Complications and Course of Events in Patient 4

As previously described,\textsuperscript{2} all patients complained of short-lasting (a few seconds) chest pain during administration of ethanol. The oxalacetyl transaminase rose to 85, 96, 65, 37, 29, and 220 units (normal at our laboratory, <40 units) in patients 1, 2, 3, 5, 6, and 7, respectively. No other complications occurred. Echocardiograms were normal in all patients except patient 4, before and after ablation. Because of the severe complication occurring in patient 4, the course of events in this patient will be described in detail.

Selective catheterization of the AV nodal artery was performed without any particular problem in this patient. Administration of iced saline and contrast material resulted in transient complete AV block, as in the other patients. A dose of 1 ml ethanol was given, and complete AV block was created. The patient was transferred to the coronary care unit. Twelve hours later, AV conduction resumed with the same ventricular rates as before the procedure. Selective catheterization of the same AV nodal artery was repeated 1 day later using the same approach. Complete AV block was again easily produced by administering iced saline or contrast material. Because of the experience during the first attempt, a dose of 2 ml 96% ethanol was given in the AV nodal artery. Complete AV block occurred, associated with chest pain. Chest pain, however, persisted and was accompanied by marked ST segment elevation in the inferior leads during pacemaker rhythm. An angiography of the right coronary artery showed that the contrast material stopped in the proximal part of the right coronary artery (Figure 2) but, farther on, the artery was patent as shown by injection of contrast material more distally using a perfusion catheter (Figure 2). The vessels distal to the main right coronary artery could not be visualized (Figure 2), suggesting occlusion of the peripheral branches. Retrospective analysis of the data suggested the following sequence of events: At the time of the first procedure, no chest pain developed during administration of ethanol, no enzyme rise occurred, and complete AV block did not persist. This suggests that, for whatever reason, the alcohol did not reach the distal end of the selective catheter. During the second procedure, a dose of ethanol was given that in retrospect was probably too large. This was followed by the development of an

### Table 1. Clinical Characteristics of the Patients Studied

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age/sex</th>
<th>Indication for TCA-AVC</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>70/F</td>
<td>Failed electrical ablation</td>
</tr>
<tr>
<td>2</td>
<td>64/F</td>
<td>Failed electrical ablation</td>
</tr>
<tr>
<td>3</td>
<td>64/M</td>
<td>Failed electrical ablation, permanent pacemaker</td>
</tr>
<tr>
<td>4</td>
<td>69/F</td>
<td>Permanent pacemaker</td>
</tr>
<tr>
<td>5</td>
<td>60/M</td>
<td>Failed electrical ablation, permanent pacemaker</td>
</tr>
<tr>
<td>6</td>
<td>69/M</td>
<td>Patient preference</td>
</tr>
<tr>
<td>7</td>
<td>54/M</td>
<td>Previous arrhythmia surgery, patient preference</td>
</tr>
</tbody>
</table>

F, female; M, male; TCA-AVC, transcoronary chemical ablation of atrioventricular conduction.
Figure 1. Panel A shows right coronary angiogram in left anterior oblique projection. Atrioventricular nodal artery is indicated by arrow. Panel B shows selective catheterization of atrioventricular nodal artery with 2.5F catheter (arrow). Contrast is given showing myocardial staining and no backflow. Panel C shows effects of contrast material on ventricular rate during atrial fibrillation. Transient complete atrioventricular block and pacemaker rhythm is observed in right part of electrocardiographic strip. Leads I, II, and III are shown. Panel D shows effects of administration of 0.5 ml 96% ethanol. After series of premature ventricular beats, complete atrioventricular block and pacemaker rhythm occur. Panel E shows escape rhythm after interruption of pacing.
inferior wall myocardial infarction with a maximal oxalacetyl transaminase rise to 363 units. The left ventricular ejection fraction was 50%. The right ventricular ejection fraction was 25%, indicating that infarction of the right ventricle also occurred. One week later, this patient's course was complicated by late ventricular fibrillation from which she was successfully resuscitated. Treatment with amiodarone was started, and the patient discharged in complete AV block 2 weeks later. At 5 months' follow-up, the patient remains in complete AV block, and her functional class for dyspnea is I.

Discussion

The results of this study indicate that transcoronary chemical ablation of the AV conduction system can be done successfully; however, the complications occurring in patient 4 warn us about the possible risks of this new technique. Although the procedure was uneventful and successful in six patients, the complications in patient 4 indicate the dangers of manipulation of the coronary arteries and the dangers of a too-large volume of ethanol. Transcoronary chemical ablation with ethanol of the AV node can be done uneventfully. To avoid complications like the ones observed in patient 4, however, several precautions seem necessary. The catheter should be fully wedged in the artery to prevent any backflow. Catheters of different diameters and with a distally located balloon might be helpful for that purpose.

References

4. Gallagher JJ, Svenson RK, Kasell JH, German LD, Bardy GH, Broughton A, Critelli G: Catheter technique for closed-chest ablation of the atrioventricular conduction system: A thera-


**KEY WORDS** • percutaneous ablation • transcoronary ablation • atrioventricular block • atrial fibrillation • cardiac arrhythmias
Transcoronary chemical ablation of atrioventricular conduction.
P Brugada, H de Swart, J Smeets and H J Wellens

Circulation. 1990;81:757-761
doi: 10.1161/01.CIR.81.3.757
Circulation is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 1990 American Heart Association, Inc. All rights reserved.
Print ISSN: 0009-7322. Online ISSN: 1524-4539

The online version of this article, along with updated information and services, is located on
the World Wide Web at:
http://circ.ahajournals.org/content/81/3/757

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally
published in Circulation can be obtained via RightsLink, a service of the Copyright Clearance Center, not the
Editorial Office. Once the online version of the published article for which permission is being requested is
located, click Request Permissions in the middle column of the Web page under Services. Further
information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to Circulation is online at:
http://circ.ahajournals.org/subscriptions/