Severe Heart Failure Associated With Tachycardia-Induced Cardiomyopathy Due to Incessant Atrioventricular Re-Entrant Tachycardia

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ABSTRACT

Short RP interval atrioventricular re-entrant tachycardias do not typically present as an incessant form. We present 2 cases of incessant atrioventricular re-entrant tachycardias leading to tachycardia-induced cardiomyopathy with severe heart failure presentation in middle-aged adults. Both underwent accessory pathway ablation and recovered normal left ventricle function before hospital discharge. (Level of Difficulty: Intermediate.) (J Am Coll Cardiol Case Rep 2021;3:479–83) © 2021 Published by Elsevier on behalf of the American College of Cardiology Foundation. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).
**CASE SERIES**

**PATIENT 1.** A 53-year-old woman with a history of depressive disorder presented to our emergency department for asthenia, shortness of breath, nausea, and continuous palpitations evolving for 2 months. She had no cardiac history.

At admission, she was normotensive and tachycardic with a regular heart rate of 180 beats/min. She was in New York Heart Association functional class IV with mild congestive heart failure signs (Table 1).

Electrocardiogram (ECG) revealed a regular 1:1 narrow QRS complex tachycardia with short RP interval (RP<PR) and RP >90 ms compatible with AVRT, atypical atrioventricular nodal re-entrant tachycardia, or atrial tachycardia (2) (Figure 1A).

Blood tests showed a significant increase in N-terminal pro-B-type natriuretic peptide, aminotransferases, and cholestatic enzymes (Table 1). Chest radiography showed bilateral venous congestion and alveolar edema (Figure 1C).

A transthoracic echocardiography (TTE) performed during the tachycardia revealed a global hypokinesia with left ventricle ejection fraction (LVEF) reduced to 20% (Video 1).

Vagal maneuvers and 2 adenosine rapid injections failed to permanently stop the tachycardia due to incessant recurrent onset after terminations (Figure 1B). Finally, the tachycardia was stopped with a 300 mg Amiodarone injection. There was no pre-excitation on baseline ECG.

The patient was admitted to our cardiology unit. She was treated with intravenous Furosemide allowing rapid clinical and biological improvement.

A control TTE was performed after 3 days of permanent sinus rhythm and revealed a normalization of LVEF (65%) (Video 2) with a moderately dilated LV (Table 1).

An electrophysiological study was performed the same day. The narrow QRS complex tachycardia was easily induced without isoproterenol administration allowing the diagnosis of an orthodromic AVRT using a left anterolateral AP, successfully ablated. Intravenous amiodarone infusion was necessary to perform the electrophysiological study because of substantial atrial vulnerability leading to atrial fibrillation conversion during AVRT or even slow pacing maneuvers.

The patient was discharged the next day. She was then asymptomatic and biological parameters were normalizing (Table 1).

**PATIENT 2.** A 59-year-old man without any cardiac history was admitted to our emergency unit for dyspnea, asthenia, and right hypochondrium pain for several weeks. His medical history was a hemorrhagic rectocolitis treated with corticosteroid.

At admission, the patient was normotensive and tachycardic. He presented congestive heart failure signs with pulmonary edema, lower limbs edema, and hepatojugular reflux (Table 1). ECG also revealed a 1:1 narrow QRS complex tachycardia (170 beats/min) with PR>RP>90 ms (Figure 2A).

Tachycardia was incessant, sometimes resumed with vagal maneuvers or adenosine injection, but with spontaneous recurrence as an orthodromic or antidromic form (prompted by vagal depressive effect on nodal conduction) after a few seconds or minutes despite a 900 mg Amiodarone injection. The morphology of pre-excitation strongly suggested the presence of a left anterolateral AP (Figure 2B).

Blood tests also showed a significant increase in N-terminal pro-B-type natriuretic peptide, aminotransferases, and cholestatic enzymes (Table 1).

The patient was admitted to our unit and treated with intravenous Furosemide. Orthodromic tachycardia was incessant despite the continuation of Amiodarone administration.

A TTE performed the first day during the few minutes in sinus rhythm revealed a discreetly dilated LV with a severely depressed LV function (30% LVEF) and mild mitral regurgitation (Videos 3 and 4, Table 1). Coronary angiography results were normal.

An electrophysiological study was performed 3 days after admission, which confirmed the diagnosis of AVRT using a left anterolateral AP. The AP ablation definitely stopped the tachycardia and restored a normal sinus rhythm.

Two days after ablation, a control TTE showed a normalization of LVEF (56%) (Video 5, Table 1). At discharge, congestive heart failure signs had disappeared, and biological parameters were continuously improving (Table 1). The patient had no recurrence of supraventricular tachycardia or heart failure.

**DISCUSSION**

We describe here 2 cases of tachycardia-induced cardiomyopathy due to incessant orthodromic AVRT with heart failure presentation.

Although permanent junctional reciprocating tachycardias are commonly incessant and associated...
TABLE 1  Patient Characteristics, Clinical, Biological, and Echocardiographic Data

|                         | Patient 1                          | Patient 2                          |
|-------------------------|------------------------------------|------------------------------------|
| Age, yrs                | 53                                 | 59                                 |
| Sex                     | Female                             | Male                               |
| Body mass index, kg/m²  | 26.7                               | 20.1                               |
| Accessory pathway       | Left lateral                       | Left anterolateral                 |
| Physical examination at admission | Rales at pulmonary bases, nausea, palpitations | Rales up to one-half lung fields, peripheral edema, hepatopulmonary reflux, right hypochondrium pain |
| Heart rate, beats/min   | 180                                | 170                                |
| Blood pressure, mm Hg   | 120/70                             | 105/83                             |
| SpO₂, %                 | 98                                 | 97                                 |
| Temperature, ºC         | 37.2                               | 37.1                               |
| Blood test              |                                    |                                    |
| NT-proBNP, ng/ml        | 4,199                              | 2,188                              |
| T-Troponin, ng/l        | 17                                 | 24                                 |
| AST (N <35 U/l)         | 99                                 | 63                                 |
| ALT (N <35 U/l)         | 318                                | 117                                |
| GGT (N <40 U/l)         | 449                                | 270                                |
| ALP (N <105 U/l)        | 188                                | 123                                |
| Bilirubin, µmol/l       | 23                                 | 41                                 |
| Creatinine, µmol/l      | 95                                 | 83                                 |
| Lactate, µmol/l         | 1.5                                | 1.7                                |
| TSH, mU/l               | 0.58                               | 2.67                               |
| Hemoglobin, g/dl        | 14                                 | 12                                 |
| TTE                     |                                    |                                    |
| Rhythm                  | Tachycardia 180 beats/min          | Sinus rhythm                       |
| LVEDD, mm (LVEDDi, mm/m²) | 50 (26)                         | 56 (29)                            |
| LVEDV, ml (LVEDVi, ml/m²) | -                             | 120 (62)                          |
| LVEF, %                 | 20                                 | 57 (32)                            |
| Left atrium volume, ml/m² |                            | 141 (77)                          |
| sPAP, mm Hg             | 20                                 | 150 (83)                           |
| Valvular heart disease  | -                                  | Mild MR                            |

*With Simpson biplane method.

ALT = alanine aminotransferase; ALP = alkaline phosphatase; AST = aspartate aminotransferase; GGT = gamma-glutamyltransferase; LVEDD = left ventricle end-diastolic diameter; LVEDVi = indexed left ventricle end-diastolic volume; LVEDV = left ventricle end-diastolic volume; LVEF = left ventricle ejection fraction; MR = mitral regurgitations; NT-proBNP = N-terminal pro-B-type natriuretic peptide; sPAP = systolic pulmonary artery pressure; TSH = thyroid stimulating hormone; TTE = transthoracic echocardiography.

with TCM (5), incessant forms of lateral AP-related AVRT with short RP interval (RP<PR) are unusual. This may be the consequence, in these two 50-year-old patients, of a progressive change in electrophysiological characteristics occurring in the atrioventricular node or the AP over time (6). However, this spontaneous evolution occurs rather exceptionally.

Among supraventricular tachycardias, incessant form and slower ventricular rate have been shown to be associated with the development of TCM (7). Patients with rapid and paroxysmal tachycardias are indeed more symptomatic and are diagnosed earlier. In these 2 cases, both the incessant presentation as well as a delay in diagnosis led to heart failure. Diagnosis delay may derive from various reasons. Patient 1 was diagnosed with “spasmophilia” since childhood and later developed depression and anxiety disorders. Therefore, these “crisis” had always been attributed to anxiety by caregivers and the patient herself. For the first time we could correlate her symptoms with tachycardia. On the other hand, Patient 2 never experienced palpitations and was referred because of dyspnea and asthenia for a couple of months as well as abnormal liver blood test prescribed by his family doctor.

Medical antiarrhythmic treatment failed to permanently stop the arrhythmia in these patients because of incessant recurrence. Therefore, rapid catheter ablation was the only therapeutic option.

Both patients presented typical heart failure signs, initial deep LV dysfunction and signs of hepatic congestion. It is also interesting to note the rapid recovery of LV function after successful arrhythmia management contrasting with the severity of initial presentations. To confirm the diagnosis of TCM, efforts were made to exclude other potential causes of
FIGURE 1  Patient 1 Electrocardiograms and Chest Radiography

(A) Initial electrocardiogram: regular narrow QRS complex tachycardia (180 beats/min) with short RP interval (RP < PR) and RP > 90 ms. (B) Recurrent onset of the tachycardia after termination by adenosine injection. (C) Chest radiography: bilateral venous congestion and alveolar edema.

FIGURE 2  Patient 2 Electrocardiograms

(A) Initial electrocardiogram: regular narrow QRS complex tachycardia (170 beats/min) with short RP interval (RP < PR) and RP > 90 ms. (B) Tachycardia restarted in an antidromic form after adenosine injection. Positive V₅, negative D₁-aVL, and positive inferior leads suggest a left anterolateral accessory pathway.
heart failure (2). For patient 1, noninvasive criteria, such as absence of risk factors for cardiovascular disease, normal T-troponin level, and normal LV wall motion at day 3 echocardiography, allowed us to reasonably rule out associated ischemic cardiomyopathy. Coronary angiography was normal for Patient 2. In addition, 2-dimensional echocardiographies did not reveal any valvular heart disease, LV hypertrophy, or argument for infiltrative heart disease. Blood pressure monitoring was normal during both hospitalizations. Rapid improvement of LVEF and barely dilated LV dimensions usually suggest the absence of underlying dilated cardiomyopathy (3), but a long-term follow-up is necessary to ensure the absence of early hidden cardiac disease.

This case series highlights that incessant AVRTs, like other supraventricular tachycardias, can lead to tachycardia-induced cardiomyopathy with severe heart failure presentation. Therefore, cardiologists should always strive to document arrhythmias when at an early, paroxysmal stage with all existing means. Furthermore, mildly symptomatic patients known to experience paroxysmal AVRTs, when an ablation is not immediately considered, should be informed about a potentially rare but severe evolution and followed up closely.

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KEY WORDS accessory pathway, arrhythmia-induced cardiomyopathy, orthodromic re-entrant tachycardia, palpitation, supraventricular tachycardia, Wolff-Parkinson-White

APPENDIX For supplemental videos, please see the online version of this paper.