

Research Letter

Nonsustained Ventricular Tachycardia at the Time of Implantation Predicts Appropriate Therapies on Rapid Ventricular Arrhythmia in Primary Prevention Patients With Nonischemic Cardiomyopathy

Results From the Very-High-Rate Registry

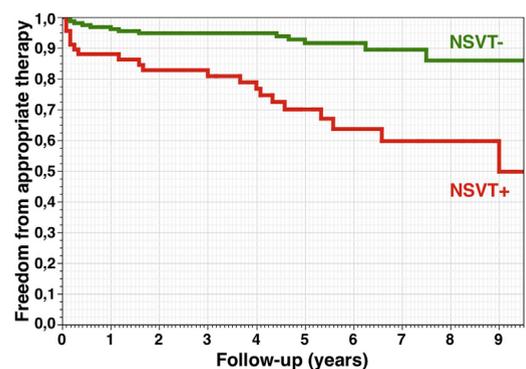
Recent results from the Danish trial call into question the benefits of implantable cardioverter-defibrillator (ICD) in primary prevention for patients with nonischemic cardiomyopathy (NICM) and a reduced left ventricular ejection fraction (LVEF), considering that a majority of these well-treated patients also received cardiac resynchronization therapy (CRT) (1). We sought to evaluate the factors associated with appropriate therapies for ventricular arrhythmias over 220 beats/min in NICM patients included in the VH-RATE (Very High Rate study) study (2,3).

Consecutive NICM patients with a reduced LVEF despite optimal medical therapy, who underwent the implantation of an ICD in primary prevention of sudden cardiac death, were included. Devices were programmed with identical tachycardia settings: a monitoring zone starting at a rate of 170 beats/min or above, and a ventricular fibrillation zone at 220 beats/min or above, with nominal settings for the number of detection intervals, according to the manufacturer (average 18 ± 4 detection intervals). Shock therapies were programmed at maximum output and anti-tachycardia pacing (ATP) before or during charging, when available. A 24-h electrocardiographic Holter monitoring was performed in all patients just before implantation to detect the presence of nonsustained ventricular tachycardia (NSVT) episodes (≥ 3 consecutive complexes and < 30 s). Time to first appropriate therapy (ATP or shock) was recorded. Follow-up ended with death, heart transplantation, or ICD removal.

A total of 223 patients were included (60 ± 11 years of age, LVEF $24 \pm 7\%$, 26% single chamber, 22% dual chamber, and 52% CRT devices). Patients were well treated, with 97% having a beta-blocker, although information on renin-angiotensin-aldosterone system inhibitors was not available. During a mean follow-up of 4.7 ± 2.5 years (median 4.7 years [interquartile range: 3.2 years]) (i.e., 1,051 patient-years), 35 (16%) patients experienced at least 1 appropriate therapy, including 24 (11%) patients with shocks (11 for ventricular fibrillation [VF], 10 for fast monomorphic ventricular tachycardia [VT], and 3 for both VT and VF), and 11 (5%) patients with ATP-only therapies.

After adjustment for age; sex; LVEF; New York Heart Association functional class; single-chamber, dual-chamber, and biventricular devices; history of atrial fibrillation; NSVT at the time of implantation; complete atrioventricular block; and treatment with a beta-blocker and amiodarone, patients with NSVT ($n = 67$, 30%) was the only independent factor associated with a higher risk of appropriate therapy over 220 beats/min (adjusted hazard ratio: 4.81; 95% confidence

FIGURE 1 Freedom From Appropriate Therapy in Patients With or Without NSVT on 24-h Holter Monitoring at the Time of Implantable Cardioverter-Defibrillator Implantation



Freedom from appropriate therapy over 220 beats/min in patients with nonischemic cardiomyopathy according to the presence (NSVT+) or absence (NSVT-) of NSVT (≥ 3 consecutive complexes) at the time of implantable cardioverter-defibrillator implantation. NSVT = nonsustained ventricular tachycardia.

interval: 2.33 to 10.30; $p < 0.0001$) (Figure 1). NSVT on 24-h Holter monitoring at the time of implantation was the only independent factor associated with a higher risk of appropriate shock on VF or polymorphic VT (adjusted hazard ratio: 5.21; 95% confidence interval: 1.63 to 18.80; $p = 0.005$). Patients with NSVT were also more likely to have NSVT episodes detected in the monitoring zone (170 to 220 beats/min) during follow-up (14.9% vs. 6.4%; $p = 0.04$).

Presence of NSVT alone identifies a subgroup (30%) of NICM patients (37% implanted with a CRT device) at very-high risk of rapid and potentially life-threatening ventricular arrhythmia (5.0 events per 100 patient-years vs. 1.7 events per 100 patient-years; $p < 0.0001$) (4). Even if ICD treated fast ventricular arrhythmia is not an equivalent endpoint as all-cause mortality, our findings suggest that NSVT may predict a higher risk population of NICM patients who will actually benefit from implantation of a defibrillator in primary prevention of sudden cardiac death. A new randomized study in that population may be needed (5).

*Nicolas Clementy, MD
 Arnaud Bisson, MD
 Farid Challal, MD
 Clementine Andre, MD
 Bertrand Pierre, MD
 Laurent Fauchier, MD, PhD
 Dominique Babuty, MD, PhD

*Cardiology Department
 Trousseau Hospital
 François Rabelais University of Tours
 60 rue du Plat d'Étain
 37044 Tours
 France
 E-mail: nclementy@yahoo.fr

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REFERENCES

1. Køber L, Thune JJ, Nielsen JC, et al. Defibrillator implantation in patients with nonischemic systolic heart failure. *N Engl J Med* 2016;375:1221-30.
2. Clementy N, Pierre B, Lallemand B, et al. Long-term follow-up on high-rate cut-off programming for implantable cardioverter defibrillators in primary prevention patients with left ventricular systolic dysfunction. *Europace* 2012; 14:968-74.
3. Clementy N, Challal F, Marijon E, et al. Very high rate programming in primary prevention patients with reduced ejection fraction implanted with a defibrillator: results from a large multicenter control study. *Heart Rhythm* 2017;14:211-7.
4. Chen J, Johnson G, Hellkamp AS, et al. Rapid-rate nonsustained ventricular tachycardia found on implantable cardioverter-defibrillator interrogation: relationship to outcomes in the SCD-HeFT (Sudden Cardiac Death in Heart Failure Trial). *J Am Coll Cardiol* 2013;61:2161-8.
5. Kadish A, Dyer A, Daubert JP, et al. Prophylactic defibrillator implantation in patients with nonischemic dilated cardiomyopathy. *N Engl J Med* 2004; 350:2151-8.