Syncope in Patients with Right Ventricle Outflow Tract Premature Beats and no Apparent Structural Cardiopathy

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Objective: Study the prevalence of neurocardiogenic etiology in patients with unexplained syncope and ventricular premature beats, with right ventricle outflow tract morphology (RVOT) and no apparent structural cardiopathy.

Methods: Ninety patients (66 women, mean age 40.2 ± 16.95 years of age) with monomorphic premature beats originated at RVOT were evaluated prospectively. Fifty-four (54) patients reported syncopes or near-syncopes associated to palpitations or not; twenty-seven (27) presented palpitations with no pre-syncope or syncope, and nine (9) were asymptomatic. All patients were submitted to echocardiogram, high resolution ECG and cardiac MRI to rule out structural cardiopathy, to exertion test to rule out adrenergic dependent ventricular tachycardia, and ECG prolonged outpatient monitoring (Holter and symptomatic events monitor) to correlate symptoms and ventricular arrhythmias. Investigation on the susceptibility to neurocardiogenic syncope was evaluated by Tilt Table Test (TTT). Groups were compared regarding gender, age, premature beats frequency and complexity, at exertion or not, TTT results and clinical course.

Results: In the syncope and pre-syncope groups, TTT was positive for 38% of cases, and in groups with palpitations and asymptomatics, it was positive for 11% (p = 0.0257). After recommendations and treatment of neurocardiogenic syncope, 85% of syncope and pre-syncope patients and positive TTT was asymptomatic along the 40-month follow-up. Two patients with syncope and negative TTT presented spontaneous, sustained ventricular tachycardia during clinic course.

Conclusion: The prevalence of neurocardiogenic syncope in patients with idiopathic RVOT premature beats is high. Patients with recurrent, unexplained syncope and idiopathic VE must be kept under investigation.

Key words: Ventricular premature beats, neurocardiogenic syncope, ventricular arrhythmias, ventricular tachycardia.
to rule out right ventricle arrhythmogenic dysplasia through echocardiography (ECHO), cardiac MRI, radioisotopic ventriculography, and signal average ECG. To investigate the correlation between arrhythmia and symptoms, patients were submitted to 24-hour Holter monitoring, to loop recorder, and ergometric test.

All patients were submitted to Tilt Table Test (TTT) to evaluate their susceptibility to neurocardiogenic syncope. TTT was carried out in the morning, following prolonged baseline protocol, as previously described. Patients who had a negative prolonged baseline TTT were submitted to a pharmacological test with sublingual 1.25 mg of isosorbide dinitrate administration. General diet and behavior recommendations were given to all patients after TTT whenever there was a positive response. If symptoms persisted, pharmacological treatment was started after initial recommendations. Patients were evaluated every three months at the outpatient unit. Additional evaluation was carried out whenever syncope occurred. All patients agreed to participate in the research protocol, which was approved by the Institutional Scientific and Ethics Committee.

Statistical analysis - Sample size was designed to detect differences higher than 20% in number of patients with positive results for TTT among groups of symptomatic and asymptomatic patients, with significance level at 5% and test power at 80%. The groups of patients were compared as to quantitative variables by using variance analysis for data with normal distribution; Kruskal-Wallis test was used for data whenever the assumption of normality was not met. Pearson χ², chi-square test and Fisher exact test were used for qualitative variables in group comparisons. Multiple comparisons were carried out based on chi-square partitions.

Results

Patients were divided into four groups, following symptomatology: I – patients with syncopes (with or without palpitations): n = 34; 24 females: mean age 37.4 ± 15.6 years of age (8 to 77, median = 39 years of age); II – patients with near-syncope (with or without palpitations): n = 20; 15 females: mean age 43.7 ± 13.4 years of age (14 to 65, median = 47 years of age); II – patients with palpitations but no pre-syncope or syncope manifestations: n = 27; 22 females, mean age 42.8 ± 15.5 (14 to 73, median = 43 years of age); IV – asymptomatic patients: n = 9; 5 females, mean age: 37 ± 23 (6 to 79, median = 27 years of age);

The comparison between groups in regard to clinical variations: age (0.3901), gender (0.1881) and number of ventricular PB under 24-hour Holter (0.2126) did not show significant difference. When the presence or absence of NSVT was investigated – at the 12-lead ECG, 24-hour Holter or ergometric test – no significant difference was found between groups (p = 0.3613). In the syncope group, two patients presented increase in ventricular PB on exertion; one of them presented paroxysmatic atrial fibrillation at peak of effort, which disappeared after exertion period was interrupted, with no clinical manifestation. Most patients presented decrease in number or absence of changes in ventricular PB behavior under physical stress. In the group with palpitations a higher number of PB under 24-hour Holter was observed as compared to the asymptomatics, and to the syncope and near-syncope group, although with no statistic difference (Figure 2).

TTT test results - The groups reported differences regarding TTT positivity (Table I). While grouping patients with syncopes and near-syncope and comparing them with groups with palpitations and asymptomatic, significant difference was observed regarding results for the TTT (Figure 3).

Thirty-eight percent of patients in the syncope and near-syncope group presented positive TTT for neurocardiogenic etiology whereas only 11% of patients with palpitations and asymptomatic presented the same result (p = 0.0257). Patients with positive TTT were younger than those with negative TTT (34.28 ± 16.01 versus 42.63 ± 15.63 years of age, respectively, with p = 0.0271).

Clinical course - Patients were followed for an average period of 37.7 ± 20.6 months (median = 40 months). Asymptomatic individuals were kept under no medication, and none reported clinical manifestation during follow-up.

Patients with palpitations and correlation between symptoms and PB received beta-blockers or anti-arrhythmic drugs. Clinical control was reached in sixteen patients (59.3%). Eleven patients (40.7%) presented persistence of symptoms, and were submitted to radiofrequency ablation (Figure 4).

From the 21 patients with syncope and near-syncope and positive TTT, 8 (38%) presented remission of symptoms with general recommendations only. From the 13 (62%) patients who also received pharmacological treatment, only 3 presented recurrence of symptoms during the follow-up. Therefore,
85.7% of patients with positive TTT turned asymptomatic, and 14.3% presented improvement of symptoms during follow-up. None of those patients presented sustained ventricular tachycardia (SVT). However, 2 patients with syncpe and negative TTT presented SVT in their clinical course. SVT was well tolerated and interrupted at the emergency unit. No death was reported.

**Discussion**

The present study demonstrates that patients with RVOT premature beats and normal heart who present unexplained, recurrent syncope must have vasovagal etiology investigated – and the prevalence is high (41%). Such information is relevant, since patients with recurring syncope and PB may present ventricular tachycardias – sustained and not sustained – as the cause of syncope, with distinctive prognostic and treatment. In such cases, two clinical conditions potentially associated with the risk of sudden death must be investigated and ruled out – arrhythmogenic ventricular displasia and catecholaminergic ventricular tachycardia – conditions under which the presence of syncope acts as a risk marker for the presence of sustained tachycardia and sudden death. In the present study, echocardiogram and cardiac MRI were carried out systematically, and used as exclusion criteria or patients with suspicion of arrhythmogenic ventricular displasia. Radioisotopic ventriculography was also carried out for cases posing doubts; patients with enlarged RV or decreased ventricular function were excluded. Patients with catecholaminergic tachycardia typically present syncope on exertion and this was an exclusion criterion for patient recruiting. Additionally, the presence of polymorphic PB on exertion was also an exclusion criterion. Therefore, this was a highly selected group. Negative results from the exams for the identification of those clinical syndromes are tranquilizing for patients with outflow PB and recurring syncope. However, the lack of a conclusive diagnosis for syncope etiology makes patients and families feel insecure, as the assisting doctor as well, who end up empirically prescribing anti-arrhythmic drugs.

The tilt table test - the method used to confirm vasovagal syncope diagnosis - is quite specific, although its positivity index does not exceed 60%, according to literature. Some patients who have persisted in their unexplained syncope condition may have had the vasovagal etiology as well, although TTT low sensitivity has not allowed their identification. Under such condition, diagnosis can only be defined through data on the syncope during prolonged ECG monitoring by implantable or external events monitoring. An electrophysiologic study may be an additional method to elucidate some cases. In the present study, 2 cases of documented SVT during their course occurred in patients whose unexplained syncope persisted. That finding suggests that although the heart may be structurally normal, the investigation follow-up using electrophysiologic studies should be recommended for patients with frequent recurrences; whenever negative, prolonged monitoring of events must be considered.

Most patients in the present study showed to be kept asymptomatic during clinical follow-up. The action mechanism of beta blockers in symptom control is not clearly defined. Krittayaphong et al. have studied patients with RVOT PB and have observed that atenolol helped to improve clinical symptoms and decrease arrhythmia frequency in the 24-hour Holter monitoring. But patients treated with placebo also showed improvement of symptoms, although no variations were reported for ventricular arrhythmia in the Holter monitoring. Such observation reiterates the idea that the symptoms reported by those patients are not fully related to ventricular arrhythmia, but rather, may be associated to autonomic tonus or psychogenic disorders. Hayashi et al. have demonstrated that the predominance of sympathetic tonus plays a relevant role in triggering repetitive VE or SVT in patients with RVOT arrhythmia without structural cardiopathy, and that propranolol not only can reduce PB density but also its complexity.
In our materials, palpitations were the most recurring symptoms. Even the 2 patients with documented spontaneous, sustained VT during clinical course, VT was correlated to palpitations rather than with syncope or pre-syncope.

Conclusions

The prevalence of neurocardiogenic etiology in patients with syncope and RVOT PB, with no structural cardiopathy is high and must always be investigated to identify patients with favorable and benign clinical course. For patients who are kept in their recurring, unexplained syncope condition invasive investigation (electrophysiologic studies) must be considered, as well as prolonged ECG outpatient unit monitoring.

References


