Repellative syncopal episodes in a child with documented ventricular tachycardia, early repolarization pattern in leads I and aVL, Brugada syndrome, and fever

Humberto Rodriguez-Reyes MD, FACC, FHRS | Andrés Pérez-Riera MD | Brenda Lucia López MD | Maria del Refugio Salinas MD | Luz Ma Mayela Muñoz MD | Cesar Ivan Laguna MD | Nikus Kjell MD

1Sociedad Cardiovascular y Arritmias (SOCAyA), Aguascalientes, México
2Laboratório de Metodologia de Pesquisa e Escrita Científica, Faculdade de Medicina do ABC, São Paulo, Brazil
3Heart Center, Tampere University Hospital and Faculty of Medicine and Health Technology, Tampere University, Tampere, Finland

Abstract
We present a small child with febrile peaks and syncopal episodes secondary to ventricular tachycardia, in whom it was eventually possible to demonstrate the Brugada Syndrome with a special presentation in the ECG; early repolarization pattern in lead I and aVL and Brugada pattern during fever in V1-V2. This is, to our knowledge, the first case with this special ECG presentation in a small child.

KEYWORDS
Brugada syndrome, early repolarization, fever, syncope, ventricular tachycardia

INTRODUCTION
Brugada syndrome (BrS) is an inherited arrhythmic disorder that may result in sudden cardiac death (SCD), and it is estimated to account for 4% of all SCDs and for 20% of all SCDs in patients with an apparently normal heart. The first arrhythmic event (AE) typically occurs in male patients aged 40 to 50 years, but rarely during childhood. According to the Survey on Arrhythmic Events in Brugada Syndrome...
(SABRUS) data, in only 4.3% the first AE occurs in the pediatric population (Milman et al., 2017). The Brugada syndrome ECG is much more prevalent in patients with fever than in afebrile controls (Michowitz et al., 2018).

We present a small child with febrile peaks and syncopal episodes, in whom it was eventually possible to demonstrate the Brugada syndrome with a special presentation in the ECG: early repolarization pattern in leads I and aVL and Brugada pattern during fever in V1-V2.

2 | CASE REPORT

Our case is a 2-year 6 months old boy, weight 14 kg with a family history of diabetes mellitus and hypertension in paternal grandparents and uncles. A cousin had arrhythmias of unknown origin, but without syncope or other alarming symptoms. There was no history of sudden death in the first-degree relatives.

At 1 year of age, he presented with a sudden episode (3 hr) of flaccidity, diaphoresis, cyanosis, polypnea, with spontaneous

FIGURE 2  Nodal rhythm with high precordial leads (marked as leads V1-V6 below), note J-wave elevation in I, aVL and high V2
recovery. In the emergency medical service, he was diagnosed with an upper respiratory tract infection. In a short period of time, he was diagnosed with a sob spasm, and repeated episodes were observed on at least five occasions within 1 year, and they were accompanied by cyanosis and diaphoresis.

Twelve hours before the current admission, he presented a new episode of flaccidity, diaphoresis, cyanosis, polypnea, seizures, and prolonged hypothermia (approximately 3 hr) with spontaneous improvement. In the following morning, the parents noticed prolonged symptoms (several hours) of diaphoresis, flaccidity, and polypnea. The attending physician directed him to the intensive care unit, where the ECG-1 (Figure 1) was performed. It showed ventricular tachycardia. Propranolol 30 mg orally was administered. After 30 min, during the placement of a central catheter the ECG-2 (Figure 2) was performed, and it showed reversal of the tachyarrhythmia observed in ECG-1. The nodal rhythm lasted for 8 hr, probably due to the effect of the beta-blocker. After that, sinus rhythm ensued (Figure 3). Rx thorax and echocardiogram were considered normal. In the afternoon, a 12 lead 24-hr Holter was performed during a febrile peak of 38 degrees Celsius (Figure 4).

**FIGURE 3** Sinus rhythm with high precordial leads (below), note J-wave elevation in aVL and high V2
3 | DISCUSSION

BrS is an inherited arrhythmic disorder that may result in SCD. The first AE of malignant ventricular arrhythmias are documented either at patient presentation because of aborted sudden cardiac arrest (SCA) or after prophylactic implantable cardioverter-defibrillator (ICD) implantation. The BrS ECG is much more prevalent in patients with fever than in afebrile controls with a high
prevalence in children aged 0–5 years (Adler et al., 2013). Although the risk of fever-related AE in BrS markedly varies according to age group, sex, and ethnicity, with a high prevalence of fever and AEs in the Caucasian population, where there is a high proportion of patients with an SCNSA mutation (77%) (Michowitz et al., 2018). The principal mechanism of AEs during fever is the worsening of the biophysical properties of the defective ion channels at higher temperatures, leading to further loss of function of the sodium channel current (Dumaine et al., 1999).

The Brugada type 1 pattern and the early repolarization pattern (ERP) are two ECG patterns characterized by the existence of J waves. Some experts have argued that based on their similarities in terms of demographics, ECG signatures, and response to neuro-modulation, ERP and BrS represent parts of a continuous spectrum of phenotypic expression, and classification under the umbrella term of "J-wave syndrome" was proposed (Antzelevitch & Yan, 2010). Inferolateral ERP occurs relatively frequently spontaneously in BrS, and these patients have a more severe phenotype (Kawata et al., ). Class 1 antiarrhythmic drug (AAD) administration provokes an inferolateral coved BrS pattern in 4.6% of patients (Sarkosy et al., 2009). Our case showed a rare presentation of J-wave elevation in leads I and aVL in association with the Br pattern in V1, V2, and in high V2.

In young patients, the presence of the spontaneous Brugada type 1 ECG pattern and symptoms at diagnosis are predictors of life-threatening arrhythmias (LTA). The management of BrS should be age-specific, and prevention of SCD may involve genetic testing and aggressive use of antipyretics and quinidine, with risk-specific consideration for ICD implantation (Andorin et al., 2016). Provocative drug testing during childhood had a low yield for predicting future AEs and was associated with serious complications in some patients. In addition, Conte et al (Conte et al., 2014) found that in 23% of children (aged 11.4 ± 2.5 years) with a negative ajmaline challenge test, BrS will be unmasked when the test is repeated several years later (age, 20.9 ± 4.2 years). Therefore, in asymptomatic patients with periodical normal ECG, adopting a cutoff of 16 years for sodium channel blocker challenge would be reasonable and in agreement with the cutoff of 15 years suggested by Andorin et al (Andorin et al., 2016). Quinidine can be used safely until adult age and posture are achieved. Quinidine also proved to be effective during relatively short follow-up and can be proposed as a valid alternative or as a bridge to ICD implantation (Probst et al., 2007). Concomitant epicardial right ventricular outflow tract ablation and ICD implantation with epicardial leads can be a safe, feasible, and effective approach for symptomatic BrS patients (de Asmundis et al., 2018). ICD therapy is an effective strategy in young patients with symptomatic BrS, treating potentially lethal arrhythmias in >25% of patients during follow-up. Appropriate shocks were significantly associated with previously aborted SCD and spontaneous type I ECGs. However, 20% of the patients experienced inappropriate shocks. Of these, 29% were secondary to supraventricular tachycardia, 42% to lead fracture and 29% to T-wave oversensing (Gonzalez Corcia et al., 2018).

4 | CONCLUSIONS

We present a young boy, who probably had ventricular tachycardia associated with fever since 1 year of age. The ECG presentation was special with ERP in leads I and aVL and BrS-type spontaneous ST elevation during fever in V1 and V2. This is, to our knowledge, the first case with this special ECG presentation in a small child.

ORCID

Humberto Rodríguez-Reyes [8] 0000-0003-3164-8515
Andrés Pérez-Riera [9] 0000-0003-4948-538X

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