Brugada Syndrome: Risk Stratification And Management

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Introduction

The Brugada syndrome (BrS) is an arrhythmogenic disease associated with an increased risk of ventricular fibrillation and sudden cardiac death. The risk stratification and management of BrS patients, particularly of asymptomatic ones, still remains challenging. A previous history of aborted sudden cardiac death or arrhythmic syncope in the presence of spontaneous type 1 ECG pattern of BrS phenotype appear to be the most reliable predictors of future arrhythmic events. Several other ECG parameters have been proposed for risk stratification. Among these ECG markers, QRS-fragmentation appears very promising. Although the value of electrophysiological study still remains controversial, it appears to add important information on risk stratification, particularly when incorporated in multiparametric scores in combination with other known risk factors. The present review article provides an update on the pathophysiology, risk stratification and management of patients with BrS.

Risk Stratification Of Individuals With Brugada Syndrome

It is widely accepted that symptomatic BrS patients are at increased risk for future events. However, in a post-mortem study, the majority of SCDs related to BrS occurred in asymptomatic individuals (72%). Based on the Second Expert Consensus Conference on BrS, 68% of this population would have been categorized as low risk. Asymptomatic BrS patients display an annual event rate of arrhythmic events between 0.5 and 1%. Emerging evidence clearly underscore our inability to stratify patients with BrS. Several markers has been proposed for risk stratification, but the majority of them have not been tested in a prospective manner. The present study focus on current risk stratification markers as well as on the management of high risk BrS patients.

Key Words:
Brugada Syndrome, Risk Stratification, Electrophysiological Study, Sudden Cardiac Death.
over time, assuming an annual event rate of 1%, a 10% event rate at 10-year follow-up in otherwise healthy patients is extremely high. Proper risk stratification of BrS patients, particularly of asymptomatic ones, is therefore of paramount importance. Several clinical, echocardiographic, electrocardiographic and electrophysiological markers have been proposed as risk stratifiers (figure 1).

Clinical Markers

A history of aborted SCD has been consistently associated with the highest risk of future arrhythmic events in all studies, and thus has a major prognostic impact.\(^\text{10-12,14-20}\) In a recent meta-analysis, the incidence of arrhythmic events (sustained ventricular arrhythmia or appropriate ICD therapy or SCD) was 13.5% per year in patients with a history of SCD, 3.2% per year in patients with syncope and 1% per year in asymptomatic patients.\(^\text{21}\) The arrhythmic risk among patients with a history of aborted SCD is 35% at 4 years,\(^\text{10,15}\) 44% at 7 years,\(^\text{12}\) and 48% at 10 years.\(^\text{21}\) A history of syncope has been associated with an increased incidence of future arrhythmic events in several studies, including a meta-analysis.\(^\text{10,11,14,16-21}\) However, Priori’s group have initially demonstrated that the association of syncope and spontaneous ST-segment elevation has the best predictive value to identify individuals at high risk, and not a history of syncope as a single risk factor.\(^\text{24}\) Kamakura et al. showed that patients presenting with aborted SCD had a grim prognosis, while those presenting with syncope or no symptoms had an excellent prognosis irrespective of their ECG pattern.\(^\text{15}\) Conte et al. from P. Brugada’s group have recently demonstrated that patients with a history of syncope display a similar clinical course with asymptomatic ones. In particular, 11% of patients with syncope and 13% of asymptomatic subjects received appropriate shocks during a long term follow-up period of 83.8 ± 57.3 months.\(^\text{12}\) This inconsistency possibly reflects our difficulty to differentiate arrhythmic from neurally-mediated syncope. A high incidence of neurally-mediated susceptibility in asymptomatic individuals with BrS ECG pattern has been previously shown.\(^\text{25}\) In Conte's study, after ICD placement, 21 patients (11.9%) experienced episodes of syncope. Of them, 5 patients had neurally-mediated syncope. In 8 patients with recurrent syncope after ICD implantation, the rate of ventricular pacing was <1%, and no ventricular arrhythmias were detected.\(^\text{26}\) This possibly explains why some patients with a previous history of undetermined syncope display an excellent prognosis.

The majority of large studies on BrS have demonstrated that a family history of SCD is not predictive of future arrhythmic events.\(^\text{10,12,24,26}\) In the largest registry, a family history of SCD was not predictive of arrhythmic events in either symptomatic (3.3% vs. 3.0%) or asymptomatic patients (0.5% vs. 0.6%).\(^\text{16}\) On the contrary, Kamakura et al. have demonstrated that a family history of SCD occurring at age <45 years is an independent risk factor of poor prognosis irrespective of ECG type. Delise et al. have shown that family history of SCD may be of prognostic significance only in combination with other risk factors.\(^\text{17}\)

In a previous meta-analysis accumulating data on 1.545 patients, male gender has been associated with a malignant clinical course.\(^\text{20}\) In FINGER registry, male gender tended to be associated with a shorter time to first event, but this difference did not reach statistical significance (mean event rate per year, 3.0% for men versus 0.9% for women).\(^\text{18}\) Similarly, in Conte’s study, males displayed a near 3-fold higher risk for appropriate shocks during follow-up.\(^\text{12}\) Data from S. Priori’s group showed that there is a non-significant excess of events in males (13%) as opposed to females (9%).\(^\text{24}\) These data suggest that females with BrS ECG pattern should not be regarded as a low risk group.

The prevalence of atrial fibrillation in BrS patients is higher than in the general population of the same age. In a large study of 560 BrS patients, 48 (99%) had atrial fibrillation/flutter.\(^\text{18}\) In 176 BrS patients with an ICD, 18% of patients developed paroxysmal AF during a long-term follow-up period of 83.8 ± 57.3 months.\(^\text{12}\) A higher incidence of atrial tachyarrhythmias (fibrillation/flutter) (24%) has been demonstrated in our BrS series.\(^\text{27}\) Spontaneous atrial fibrillation has been associated with higher incidence of syncopal episodes (60.0% vs. 22.2%) and documented VF (40.0% vs. 14.3%). In patients with documented VF, higher incidence of spontaneous atrial fibrillation (30.8% vs. 10.0%, \(p < 0.05\)), atrial fibrillation induction (53.8% vs. 20.0%), and prolonged interatrial conduction time was observed.\(^\text{28}\) Siera et al. have recently reported that asymptomatic BrS subjects with history of sinus node dysfunction display an 8-fold increased risk for future arrhythmic events.\(^\text{29}\)

Genetic Markers

A genetic defect on the SCN5A gene has not been associated with a higher risk of future arrhythmic events in several studies, suggesting that genetic analysis is a useful diagnostic parameter but it is not helpful for risk stratification.\(^\text{10-12,24,26}\)

Echocardiographic Markers

A reduced right ventricular ejection fraction and an increased right ventricular end-diastolic volume were independently associated with history of syncope or SCD at the time of diagnosis.\(^\text{30}\) Using tissue velocity imaging, Van Malderen et al. have demonstrated that a previous history of malignant events is associated with prolonged right ventricular ejection delay.\(^\text{31}\)

Electrocardiographic Markers

A spontaneous type 1 ECG pattern of BrS has been consistently associated with a worse outcome in large studies,\(^\text{10-12}\) and thus should be considered as a malignant marker. A meta-analysis showed that individuals with spontaneous type 1 ECG features exhibit a 3- to 4-fold increased risk of events compared to those with a drug-induced ECG pattern.\(^\text{20}\) We recently demonstrated that asymptomatic subjects with spontaneous type 1 ECG pattern of BrS exhibit a 3.5 higher risk of future arrhythmic events.\(^\text{32}\) It is therefore

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**Figure 1:** Clinical, electrocardiographic and electrophysiological markers proposed for risk stratification in BrS
very important to establish or not the presence of spontaneous type 1 ECG pattern in BrS patients. In patients with drug-induced type 1, spontaneous type 1 BrS can be more frequently detected with twelve lead Holter monitoring compared with conventional follow-up with periodic ECGs. Twelve lead Holter recording might avoid 20% of the pharmacological challenges with sodium channel blockers, which are not without risks, and should thus be considered as the first screening test, particularly in children or in the presence of borderline diagnostic basal ECG.43

Apart from the spontaneous type 1 ECG pattern, several other ECG parameters have been proposed for risk stratification of subjects with BrS phenotype. Among these ECG markers, QRS-fragmentation in twelve-lead ECG appears very promising. Morita et al. have initially demonstrated that QRS-fragmentation is more commonly seen in BrS patients with VF (85%) and syncpe (50%) compared to asymptomatic ones (34%).44 The PRELUDE study confirmed these findings and showed that QRS-fragmentation is an independent predictor of future arrhythmias.45 Tokioka et al. have recently demonstrated that the presence of QRS-fragmentation lead to a 5-fold increase of arrhythmic events.46 A prospective study using signal-averaged ECG suggested that positive late potentials may have predictive value of malignant arrhythmic events in BrS.47 Ajiro et al. have showed that symptomatic subjects with BrS display significantly lower RMS40, longer LAS40, and longer filtered QRS duration compared to the asymptomatic ones.48 The positive predictive value, negative predictive value, and predictive accuracy of late potentials were 92.0%, 78.9%, and 86.4%, respectively. Daily fluctuations in ECG and SAECG characteristics could be useful for distinguishing between high- and low-risk patients with BrS.49

First degree atrioventricular block has been independently associated with SCD or appropriate ICD therapies.50 A prolonged QRS duration in leads II, V2 and V651,52 as well as a prolonged QTc interval >460 ms in lead V241 have been associated with life-threatening arrhythmic events in BrS. The Tpeak-Tend interval, a marker of transmural dispersion of repolarization, has been linked to malignant ventricular arrhythmias in various clinical settings including the BrS. Castro Hevia et al. were the first to link increased Tpeak-Tend interval as a risk factor in patients with BS.53 Maury et al. has been recently demonstrated that the Tpeak-Tend interval from lead V1 to lead V4, the maximum value of the Tpeak-Tend interval, and the Tpeak-Tend interval dispersion in all precordial leads were significantly higher in symptomatic patients (aborted SCD, appropriate ICD therapy, syncope) than in asymptomatic patients. In multivariate analysis, a max Tpeak-Tend of >100 ms was independently related to arrhythmic events.54 We have previously shown that the Tpeak-Tend interval and Tpeak-Tend interval/QT ratio were associated with VT/VF inducibility in BrS.42 A more negative T-wave in lead V1 has been also associated with poor prognosis.43 Finally, the appearance of T-wave alternans after pilsicainide administration was predictive for spontaneous VF.44

Mastrur et al. performed a systematic review including 166 BrS patients undergoing exercise testing.45 ST-segment augmentation was observed in 95 of 166 (57%) BrS patients. The ST augmentation occurred during early recovery after exercise in 93 Brugada patients, whereas 2 patients developed ST augmentation during the effort phase of exercise. Exercise unmasked the BrS ECG pattern in 5 patients. Three patients developed ventricular arrhythmias with exercise: 2 developed ventricular tachycardia, and 1 developed multiple ventricular extrasystoles. All 3 arrhythmias occurred during early recovery after exercise testing and resolved spontaneously. Makimoto et al. showed that ST-segment augmentation at early recovery was specific in BrS patients, and was significantly associated with a higher cardiac event rate, notably for patients with previous episode of syncope or for asymptomatic patients.46 On the contrary, Amin et al. failed to show significant differences in the ECG variables and their changes during exercise between symptomatic (prior syncope) and asymptomatic (no prior syncope) BrS patients.47

An infero-lateral early repolarization pattern has been shown to be predictive of arrhythmic events.57,58 Tokioka et al. have recently shown that the combination of QRS-fragmentation and early repolarization pattern enables the identification of high risk patients.21 In a previous study, including 290 individuals with BrS, an early repolarization pattern manifested as notched or slurred J-point elevation mainly in lateral leads was observed in 35 subjects (12%). However, in this study, the presence of early repolarization pattern was not associated with arrhythmic events during follow-up.59 Finally, the aVR sign, defined as R wave ≥0.3 mV or R/q ≥0.75 in lead aVR, has been associated arrhythmic events during follow-up.60

Electrophysiological Markers

Conflicting evidence exists on the prognostic value of electrophysiological study (EPS) in asymptomatic BS subjects. Previous studies have demonstrated an excellent negative predictive value of EPS.17,52 On the contrary, in the PRELUDE registry, a negative EPS was not associated with a low risk of an arrhythmic event.11 Disagreement also exists regarding the positive predictive value of EPS. Data from the Brugada’s series have shown that VT/VF inducibility is predictive for future events.14,52,53 However, data from other studies do not support the use of EPS in risk stratification.10,12,18,19

Important data on the prognostic significance of EPS in BrS are coming from Pedro Brugada’s group.29 In their series, patients with VF inducibility presented a hazard ratio for events of 8.3. Event free survival for the non-inducible group was 99.0% at 1 year and 96.8% at 5, 10 and 15 years. Among the inducible patients it was 89.0% at 1 year, 78.4% at 5 years and 75.0% at 10 and 15 years. Among asymptomatic patients, those without EPS inducibility had an event free survival of 100.0% at 1 year, and 99.2% at 5, 10 and 15 years. Inducible subject’s event free survival was 90.6% at 1 year and 79.5% at 5, 10 and 15 years. EPS inducibility was also significant for the asymptomatic subjects. Sensitivity of EPS for predicting arrhythmic events was 64.0% and specificity was 86.6%. Positive predictive value was 21.6% and negative predictive value was 97.7%. If restricted to asymptomatic patients, these values increased to a sensitivity of 75.0% and a specificity of 91.3% and predictive values to 18.2% and 98.3% respectively.

Based on recent meta-analyses, EPS inducibility appears to have a prognostic role in risk stratification of BrS. Faucher et al. performed a meta-analysis of 13 studies evaluating the prognostic role of EPS in BrS patients according to clinical presentation.21 In the whole population of BrS patients, VF inducibility was associated with a non significant higher risk of arrhythmic events during follow-up. However, induction of sustained ventricular arrhythmia was significantly and homogeneously associated with an increased risk of arrhythmic events during follow-up in patients with syncope (odds ratio of 3.30). Similarly, the asymptomatic patients with inducible
VF had an increased risk of arrhythmic events during follow-up (odds ratio of 4.62) with homogeneous results across the different studies. In Stroubek et al. meta-analysis, VF induction was associated with cardiac events during follow-up with a hazard ratio of 2.66, with the greatest risk observed among those induced with single or double extrastimuli. We have recently conducted a meta-analysis of 12 studies comprising 1,104 asymptomatic subjects with BrS who underwent EPS. During follow-up, arrhythmic events occurred in 3.3% of cases. Inducible ventricular arrhythmias at EPS were predictive of future arrhythmic events with an odds ratio of 3.5.32

VF inducibility by programmed electrical stimulation, abnormal restitution properties, and ventricular effective refractory period <200 ms. Finally, EPS may establish the presence of sinus node dysfunction, clarify the cause of syncope or treat supraventricular arrhythmias that can mislead the diagnosis or eventually lead to inappropriate ICD therapies.

Multiparametric Risk Stratification Scores

The relationship of these markers and the usefulness of their combination have not been sufficiently examined. Brugada et al. have shown that patients with a spontaneously abnormal ECG, a previous history of syncope, and inducible sustained ventricular arrhythmias had a probability of 27.2% of suffering arrhythmic events during follow-up. Similarly, Priori et al. have demonstrated that the combined presence of spontaneous type 1 ECG pattern and the history of syncope identifies subjects at risk of cardiac arrest. In the same line, Delise et al. have recently proposed that subjects at highest risk are those with spontaneous type 1 ECG pattern and at least two additional risk factors (syncope, family history of SCD, or positive EPS). Okamura et al. have shown that syncope, spontaneous type 1 ECG pattern, and inducible ventricular arrhythmias at EPS are important risk factors and the combination of these risks well stratify the risk of later arrhythmic events. When dividing patients according to the number of these 3 risk factors present, patients with 2 or 3 risk factors experienced arrhythmic events more frequently than those with 0 or 1 risk factor.55

Management Of Patients With Brugada Syndrome

Based on 2015 ESC Guidelines for the management of patients with ventricular arrhythmias and the prevention of SCD, ICD implantation is recommended in BrS patients with aborted SCD or with documented spontaneous sustained ventricular arrhythmias (Class I, LOE C). ICD implantation should be considered in patients with a spontaneous diagnostic type 1 pattern and history of syncope (Class IIa, LOE C). ICD implantation in BrS patients with inducible VF at EPS receives a weak recommendation in the new 2015 ESC Guidelines (Class IIb, LOE C). However, based on recent long term follow-up data as well as on two meta-analyses, asymptomatic individuals with spontaneous type 1 ECG and inducible VF at EPS are possibly at high risk. Nevertheless, the decision to implant an ICD in asymptomatic patients should be made weighing the potential individual risk for future arrhythmic events against risk of complications and quality of life. In a recent study, a significant number of young BrS patients experience device-related complications (15.9%). Complications consisted of fracture of the ventricular electrode and lead dislocation, and less commonly device infection and pulse generator migration. In this series, 18.7% of patients suffered inappropriate shocks. Quinidine may be considered as an alternative therapy in patients who denied ICD as well as for treatment of supraventricular arrhythmias. Low doses of quinidine are effective to prevent the recurrence of VF, including arrhythmic storm, in subjects with BrS with an ICD. Furthermore, quinidine effectively prevents VT/VF induction in patients with BrS at EPS. Belhassen et al. have recently reported the long-term outcomes of BrS patients with initially inducible VF at EPS who received quinidine (n=54), disopyramide (n=2) or both (n=4) and underwent a second EPS procedure. Fifty four patients (90%) were responders to 1 anti-arrhythmic drugs and became non-inducible. No arrhythmic events occurred during class 1A anti-arrhythmic drugs therapy in any of EPS-drug responders and in patients with no baseline inducible VF during a very long follow-up period of 113.3±71.5 months.

Isoprenaline infusion is effective in the management of repeated ICD shocks and arrhythmic storms. Cilostazol and milrinone that boost calcium channel current and quinidine, bepridil and the Chinese herb extract Wenxin Keli that inhibit the transient outward current may be used to suppress the triggers for VF in BrS. Nademanee et al. initially showed that catheter ablation of fractionated electrograms in the epicardial right ventricular outflow rendered VF non-inducible and normalization of the BrS ECG pattern. Long-term outcomes were excellent, with no recurrent VF in all patients off medication. Similarly, in a recent study, following catheter ablation and elimination of the functional substrate in right ventricular outflow tract, all patients became non-inducible during programmed electrical stimulation using up to 3 extrastimuli, while repeated flecainide infusion failed to unmask the diagnostic BrS ECG pattern. Although these findings have to be confirmed in future studies, they provide new important information regarding the therapeutic management of BrS patients.

Conclusions

Risk stratification of BrS patients represents a great challenge for the treating physician. Despite the lack of evidence, it is of major importance to make the best risk stratification using every available tool/modality that has been shown to display any prognostic significance. The diagnostic yield may be therefore increased if we use the current tools properly. Although single risk factors display limited prognostic value, multiparametric scores appear to improve risk stratification.

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