

Consensus Report

Proposed Diagnostic Criteria for the Brugada Syndrome

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Introduction

A syndrome characterized by ST segment elevation in right precordial leads (V₁ to V₃), unrelated to ischemia, electrolyte disturbances or obvious structural heart disease, was reported as early as 1953^[1] but first described as a distinct clinical entity associated with a high risk of sudden cardiac death in 1992^[2] (see^[3,4] for review). The Brugada syndrome is a familial disease displaying an autosomal dominant mode of transmission with incomplete penetrance and an incidence ranging between 5 and 66 per 10 000. In regions of Southeast Asia where it is endemic, the clinical presentation of Brugada syndrome is distinguished by a male predominance (8:1 ratio of males:females) and the appearance of arrhythmic events at an average age of 40 years (range: 1 to 77 years)^[2,5]. Although a number of candidate genes are considered plausible, thus far the syndrome has been linked only to mutations in *SCN5A*, the gene encoding for the α subunit of the sodium channel^[6].

A number of ambiguities exist concerning the diagnosis of Brugada syndrome. The electrocardiographic signature of the syndrome is dynamic and often concealed, but can be unmasked by potent sodium channel blockers such as flecainide, ajmaline and

procainamide^[7], although the specificity of this effect for uncovering patients at risk for sudden death has been an issue of concern. A recent report by Remme *et al.*^[8] has shown that the number of idiopathic ventricular fibrillation patients diagnosed as having Brugada syndrome is a sensitive function of the diagnostic criteria applied. What are the proper diagnostic criteria to be used in identifying Brugada syndrome? A definitive answer to this question has been out of reach and is the reason for the establishment of a special Arrhythmia Working Group of the European Society of Cardiology that met on 31 August–1 September 2000. This report is the consensus document emanating from that meeting. The diagnostic criteria described herein are based on the currently available clinical data and state-of-the-art understanding of the molecular and cellular mechanisms underlying Brugada syndrome. The proposed criteria must be considered a work-in-progress that will be fine-tuned as confirmatory data from future molecular studies and prospective trials become available.

Electrocardiographic characteristics

ECG abnormalities constitute the hallmark of Brugada syndrome. They include repolarization as well as depolarization abnormalities in the absence of identifiable structural cardiac abnormalities or other conditions or agents known to lead to ST-segment elevation in the right precordial leads (Table 1). Three types of repolarization patterns are recognized (Fig. 1). Type 1, described in the 1992 paper^[2], is characterized by a prominent coved ST-segment elevation, displaying

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Table 1 ST segment abnormalities in leads V_1 – V_3

	Type 1	Type 2	Type 3
J-wave amplitude	≥ 2 mm	≥ 2 mm	≥ 2 mm
T-wave	Negative	Positive or biphasic	Positive
ST-T configuration	Coved type	Saddle back	Saddle back
ST segment (terminal portion)	Gradually descending	Elevated ≥ 1 mm	Elevated <1 mm

1 mm=0.1 mV, the terminal portion of the ST-segment refers to the latter half of the ST-segment.

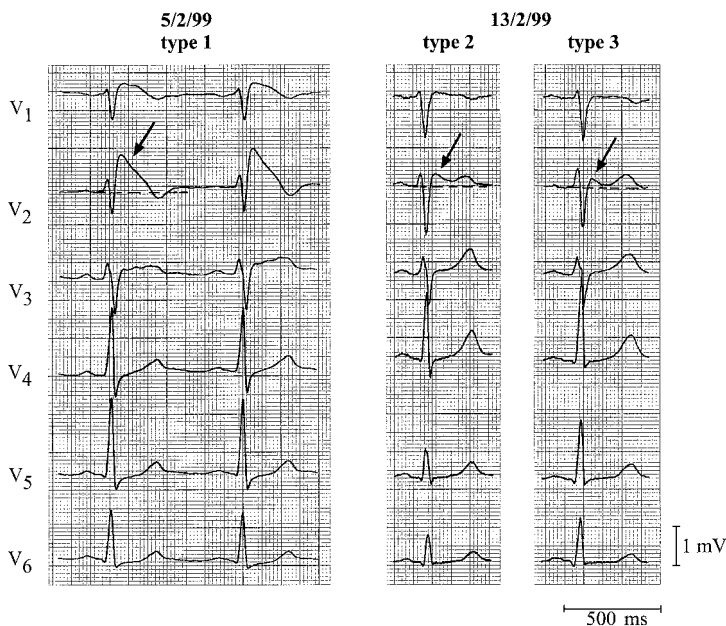


Figure 1 Precordial leads of a resuscitated patient with Brugada syndrome. Note the dynamic ECG changes in the course of a couple of days. All three patterns are shown. Arrows denote the J-wave (see text for definition). The left panel shows a clear type 1 ECG. Between 7 February 1999 and 13 February 1999 types 2 and 3 appear. Calibrations are given.

a J-wave amplitude or elevated ST segment elevation ≥ 2 mm or 0.2 mV at its peak followed by a negative T-wave, with little or no isoelectric separation. Type 2 also has a high take-off ST-segment elevation but in this case the J wave amplitude (≥ 2 mm) gives rise to a gradually descending ST-segment elevation (remaining ≥ 1 mm above the baseline) followed by a positive or biphasic T-wave resulting in a saddle back configuration. Type 3 is a right precordial ST-segment elevation comprising a <1 mm saddle back or coved configuration (or both). It should be stressed that delineation of the J wave is sometimes tricky (i.e. second ECG in Fig. 1) and that these descriptions are based on the correct placement of precordial leads, although characteristic ECG features obtained with alternative placement of right precordial leads in a superior intercostal space in individuals with high clinical suspicion (aborted sudden cardiac death (SCD) victims, family members of patients with Brugada syndrome, see below) may also disclose the presence of arrhythmic substrate^[9]. In select cases

one may even consider rightward displacement. However, the r' deflection in leads V_3R , V_4R etc should be interpreted with caution. Characteristic ECG morphologies recorded in the first few hours after resuscitation or immediately after DC shock cannot be taken as diagnostic of Brugada syndrome. As shown in Fig. 1, the ST-segment is dynamic. Different patterns may be observed sequentially in the same patient or following the introduction of specific drugs (see below).

The QT-interval is often within normal limits (in the absence of anti-arrhythmic drug therapy), but may be prolonged. In the initial series described by Brugada and Brugada (1992), three out of six males had a $QT_c \geq 440$ ms^[2]. Also in male Thai patients with RBBB and ST-elevation, the mean QT_c was slightly longer than normal^[10]. Families with (drug induced) ST elevation and QT prolongation have also been described^[11,12].

Conduction disorders vary from non-specific to specific for any given part of the conduction system (in drug-free state). To the former category belongs the

often-encountered pronounced broad S in leads I, II and III, giving rise to left or extreme axis deviation. A left axis may also indicate a left anterior hemiblock. A true RBBB may be seen with or without right or left axis deviation. The high take-off ST-segment previously discussed may mimic an RBBB pattern, but the absence of S-waves in the left-lateral leads precludes the true presence of right ventricular conduction delay.

The PR interval is often increased (≥ 200 ms) and presumably reflects the presence of HV-conduction delay (≥ 55 ms). In a recent survey the latter was found to be present in 20/21 Brugada patients^[4], and was usually in the range of 65 ms, but could be as long as 110 ms. However, Eckardt *et al.* reported a mean HV interval of 49 ± 12 ms in 35 patients, with only six patients over 60 ms^[13].

Particular problems exist in the pediatric population because of the lack of control data, the different chest morphology, and the age-dependent predominance of right ventricular forces. However, typical ECG patterns have been observed in small infants, where eventual lethal arrhythmias might resemble sudden infant death syndrome^[5,14]. Hence, suspect symptoms with typical electrocardiographic features and/or a family history for sudden cardiac death, even at young age, should alert pediatricians to the possibility of Brugada syndrome.

Drug challenge

Intravenous administration of certain drugs may modify the ECG pattern. Ajmaline ($1 \text{ mg} \cdot \text{kg}^{-1}$ body weight; $10 \text{ mg} \cdot \text{min}^{-1}$), flecainide ($2 \text{ mg} \cdot \text{kg}^{-1}$, max 150 mg ; in 10 min) and procainamide ($10 \text{ mg} \cdot \text{kg}^{-1}$; $100 \text{ mg} \cdot \text{min}^{-1}$) exaggerate the ST-segment elevation or unmask it when it is initially absent. Sensitivity and specificity (with genetic data as the gold standard) for intravenous drug challenges are disputed^[7,12]. However, there is consensus that in the case of procainamide, sensitivity is relatively low. Reproducibility of the test has not been established and a recent study suggests that it might be less than 100%^[15].

Drug challenge should be performed while the patient is continuously monitored (12-lead ECG and blood pressure) and with defibrillator and ACLS facilities close at hand. Accurate lead position and correct venous access should be ascertained. Drug administration should be stopped when the test is positive (see below) and/or when ventricular arrhythmias, including ventricular premature complexes, are evident or when significant QRS widening ($\geq 30\%$) is observed. In the case of a negative baseline ECG, a J-wave amplitude of $>2 \text{ mm}$ absolute amplitude in lead V_1 and/or V_2 and/or V_3 with or without RBBB is considered positive. In patients with type 1 ECGs, drug testing is not of additional diagnostic value. In patients with types 2 and 3 ECGs, the test is recommended for the purpose of clarifying the diagnosis. Conversion of types 2 or 3 ECG to type 1 is considered positive (Fig. 2). An increase in the J-wave amplitude of more than 2 mm without the

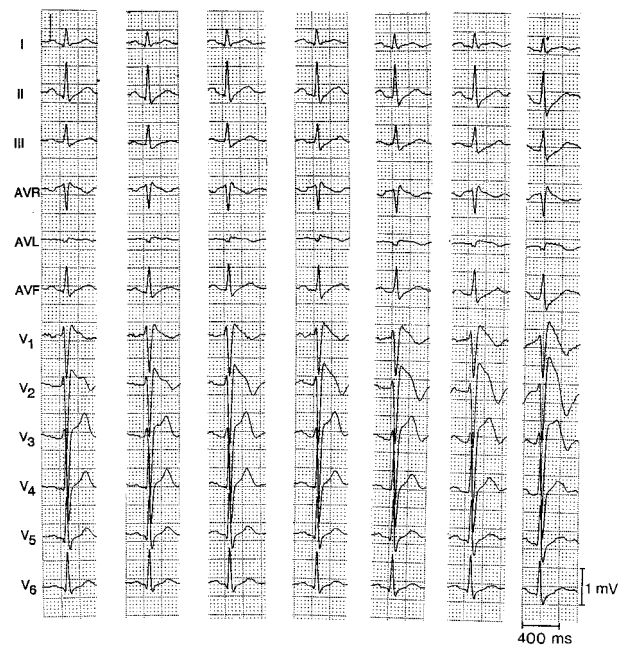


Figure 2 ECGs from a 35-year-old male who has been successfully resuscitated. From left to right two control ECGs, and 1, 2, 3, 4 and 5 min during i.v. infusion of 50 mg ajmaline. The ECG is of 'saddle back' type (type 2) in the left panels and becomes coved in the right panels (type 1). Calibrations are standard.

development of a type 1 configuration is also considered significant, but in our experience is rarely observed. Conversion of type 3 ECG into type 2 is considered inconclusive.

Monitoring is recommended until the ECG has normalized (plasma half-lives of the different drugs are: flecainide 20 h, procainamide 3–4 h, ajmaline inactivated within a few minutes). Serious ventricular arrhythmias, including VF, may occur during the test. Immediate discontinuation of the drug is required and isoproterenol infusion might be needed to treat the arrhythmias ($1\text{--}3 \mu\text{g} \cdot \text{min}^{-1}$ isoproterenol^[14]; P. Brugada, personal observations).

It seems intuitive that the more sodium channel block is needed to elicit the Brugada phenotype, the less likely the patient is to be at risk under baseline conditions. Indeed, recently it was shown that asymptomatic patients, with an abnormal ECG only upon drug challenge, have a benign prognosis^[16]. Canine ventricular wedge studies suggest that a flecainide induced Brugada phenotype does not necessarily indicate the presence of an arrhythmic substrate; it does denote the ability of sodium channel block to create the conditions under which the arrhythmic substrate may readily develop^[17].

Clinical presentation

All too often syncope or sudden cardiac death is the only symptom in patients with Brugada syndrome. In some cases, sudden death is the first symptom of the disease.

Table 2 (a) Abnormalities that can lead to ST-segment elevation in the right precordial leads

Right or left bundle branch block, left ventricular hypertrophy^[28]
 Acute myocardial ischemia or infarction^[29]
 Acute myocarditis^[30]
 Right ventricular ischemia or infarction^[31]
 Dissecting aortic aneurysm^[32]
 Acute pulmonary thromboemboli^[33]
 Various central and autonomic nervous system abnormalities^[34,35]
 Heterocyclic antidepressant overdose^[36]
 Duchenne muscular dystrophy^[37]
 Friedreich's ataxia^[38]
 Thiamine deficiency^[39,40]
 Hypercalcemia^[41]
 Hyperkalemia^[42]
 Cocaine intoxication^[43,44]
 Mediastinal tumor compressing RVOT^[45]
 Arrhythmogenic right ventricular dysplasia/cardiomyopathy^[24,25]
 LQTS, type 3^[11,12].

Table 2(b) Other conditions that can lead to ST-segment elevation in the right precordial leads

Early repolarization syndrome
 Other normal variants (particularly in men)

Whereas these two conditions, which can lead to ST-segment elevation are more likely to give rise to type 2 and 3 ECGs, most conditions mentioned in this table (a + b) can give rise to a type 1 ECG.

Monitoring of such patients has revealed rapid polymorphic VTs as the underlying cause. VT usually starts with a short coupling interval. Self-terminating episodes typically lead to repeated episodes of syncope. Indeed, 80% of patients with documented VF have a history of syncope^[15]. Clinical reports indicate that sudden death in Brugada patients most commonly occurs during sleep, in particular during the early morning hours^[9,18].

It has been suggested that there is a higher than normal incidence of supraventricular tachyarrhythmias, including atrial and AV reentrant tachycardia in the Brugada population^[13]. Rarely monomorphic VT is observed^[19,20]. Family history is of paramount importance and is often positive for sudden cardiac death at a young age.

The mean age at which symptoms first appear in affected individuals (males and females) is in the third to fourth decade. However, among the first patients described were symptomatic twins, 1 year of age, and more recent reports describe families with symptomatic children^[5,14]. A patient in which first symptoms appeared at an age of 77, has also been described^[2].

Electrophysiological studies

Electrophysiological studies (EPS) may be helpful in risk stratification and in some cases in establishing the diagnosis. A complete EPS is recommended in all symptomatic patients. In VF survivors, EPS may be of little or no diagnostic value, but may be helpful in providing further insight into the predictive value of available

diagnostic tools. In the absence of data on sensitivity or specificity of any EPS protocol, we suggest a protocol using two stimulation sites (RVA, RVOT), at least three cycle lengths (600, 430, 330 ms), 1, 2 and 3 extrastimuli and a minimal coupling interval of 200 ms. It is at present unknown whether shortening of the coupling interval to refractoriness, which might be very short in patients with Brugada syndrome, has additive value. Inducibility early in the protocol is common but non-inducible cases have been described. About half of the patients are inducible from the outflow tract. Isoproterenol does not enhance the likelihood of inducibility and in theory should have the opposite effect. The diagnostic value of repeating EPS after class I drug treatment (or challenge) is not fully established. Similarly, the value of epicardial stimulation^[21] as well as the value of right ventricular mapping with monophasic action potentials is not known^[22]. Asymptomatic patients with a positive family history for SCD should be investigated in a similar way. The necessity for EPS is questionable in patients displaying the Brugada ECG, but who are asymptomatic and have a negative family history. In fact, accuracy to predict outcome is debated. Positive predictive value varies from 50%^[15] to 37%^[23] and negative predictive value varies from 46%^[15] to 97%^[16,23].

Differential diagnosis

The list of factors contributing to ST segment elevation is long (Table 2). Clinical diagnostic evaluation should be directed to excluding each of these causes.

Table 3 Differential diagnosis between ARVC and Brugada syndrome

Clinical characteristics	ARVC	Brugada syndrome
Age at presentation	25–35	35–40
Sex (male/female)	3:1	8:1
Distribution	Worldwide*	Worldwide *
Inheritance	AD(AR)	AD
Chromosomes	1,2,3,10,14 (17)	3
Gene	hRYR2, plakoglobin	SCN5A
Symptoms	Palpitations Syncope Cardiac arrest	Syncope Cardiac arrest
Circumstances	Effort	Rest
Imaging	Morpho-functional RV (and LV) abnormalities	Normal
Pathology	Fibrofatty replacement	Normal
ECG repolarization	Inverted T-waves in precordial leads	High take-off ST-segment V ₁ –V ₃
ECG depolarization	Epsilon-waves QRS-prolongation	RBBB/LAD
AV conduction	Normal	50% abnormal PR/HV
Atrial arrhythmias	Late (secondary)	Early (primary 10–25%)
ECG changes	Fixed (mostly)	Variable
Ventricular arrhythmias	Monomorphic VT/VF	Polymorphic VT/VF
Mechanism of arrhythmias	Scar-related	Phase 2
Drug effect Class I	↓	↑
Drug effect Class II	↓	↑
Drug effect Class III	↓	-/↑
Drug effect Class IV	-/↓	-
Beta-stimulation	↑	↓
Natural history	Sudden death Heart failure	Sudden death

Arrows denote changes in ST segment elevation (↑, increased; ↓, decreased; -/, small change, if any).

*ARVC (arrhythmogenic right ventricular cardiomyopathy) is frequently found in North-East Italy and Brugada syndrome may be particularly prevalent in South-East Asia.

Discrimination between Brugada syndrome and arrhythmogenic right ventricular cardiomyopathy (ARVC) may be particularly difficult since ARVC may at times mimic Brugada syndrome and structural abnormalities may only be found at time of autopsy^[24,25]. Before the diagnosis of Brugada syndrome is made a serious attempt should be taken to exclude ARVC. Drug challenge with sodium channel blockers may be useful in discriminating between these two entities^[7]. Intuitively one would expect a rightward shift of the epsilon wave, if present, in ARVC patients. Table 3 lists characteristics of the two diseases that may be useful in making a differential diagnosis.

Brugada syndrome should also be discriminated from the early repolarization syndrome (with an eventual elevated J-wave amplitude in the left precordial leads) and from normal degrees of right precordial ST elevation in men, which may mimic a type 2 or 3 Brugada ECG pattern^[26]. Once again, a drug challenge might provide the clue for a proper diagnosis.

Molecular genetics

Genetic linkage analysis may be of great help in arriving at a definitive diagnosis. However, there are several limitations. First, it may take weeks to months

before a mutation in *SCN5A*, the only gene thus far causally linked to Brugada syndrome, is identified. Second, there are few diagnostic laboratories capable of screening for the disease; and, third, only a small fraction of patients can be successfully genotyped at this time (estimates vary between 10 and 30%). With the identification of other genes, this fraction will increase in future years. Finally, the identification of a new mutation in a sporadic case is equivocal at best. In such cases, data concerning the function of the abnormally encoded protein may be helpful in substantiating the physiologic relevance of the gene mutation and in estimating the level of clinical risk. With regard to screening of relatives, a 'proven' mutation in the proband could enable better follow-up and possibly more effective treatment of family members who may still be asymptomatic.

SCN5A mutations linked to Brugada syndrome have been found throughout the gene^[26]. A common denominator in all of these is a functional reduction in the availability of the sodium channel current. Mechanisms thus far include failure of the channel to express, accelerated or premature inactivation, shifts of steady state inactivation curves to more negative values and increased the likelihood of the channel to enter an intermediate inactivated state (with slow recovery kinetics) (for review see ^[27]).

The diagnosis of Brugada syndrome

Brugada syndrome should be strongly considered in the following cases:

(1) Appearance of a type 1 ST segment elevation (coved type, Table 1) in more than one right precordial lead (V_1 – V_3), in the presence or absence of a sodium channel blocker, and one of the following: documented ventricular fibrillation, self terminating polymorphic ventricular tachycardia, a family history of SCD (<45 years), coved type ECGs in family members, electrophysiological inducibility, syncope or nocturnal agonal respiration. There should be no other factor(s) accounting for the ECG abnormality.

The appearance of the ECG features, without these clinical symptoms, is referred to as an idiopathic Brugada ECG pattern (not Brugada syndrome).

(2) Appearance of type 2 ST-segment elevation (saddle-back type) in more than one right precordial lead under baseline conditions with conversion to type 1 following challenge with a sodium channel blocker is considered equivalent to case 1 above. Drug-induced ST-segment elevation to a value greater than 2 mm should raise the possibility of Brugada syndrome, when one or more clinical criteria are present (see case 1 above). Based on our limited knowledge at present, a patient with a negative drug test (i.e. no change in the ST-segment in response to a sodium channel blocker) is unlikely to have Brugada syndrome; drug induced ST-elevation to <2 mm is considered inconclusive.

(3) Appearance of type 3 ST segment elevation in more than one lead under baseline conditions with conversion to type 1 following challenge with a sodium channel blocker is considered equivalent to case 1 above and should be screened accordingly. Drug-induced conversion of type 3 to type 2 ST segment elevation is considered inconclusive.

Patients who do not fully qualify the proposed criteria (e.g. type 1 ECG with a J-wave amplitude of only 1 mm), but with one or more clinical criteria as defined above, should be considered seriously. Most often a drug challenge will disclose the diagnosis Brugada syndrome. In addition, in these and other cases mentioned, EPS might be useful.

Limitations

This consensus meeting started to develop a diagnostic scheme similar to “point systems” available for other arrhythmogenic disorders, like the long QT syndrome. However, it was abandoned when we came to the realization that such a quantitative approach would require more data than currently available. In particular, we lack data on the spectra of ECG patterns that may be associated with an increased risk of death. As such we would like to stress once again that the proposed criteria are based on currently available data and that it is a work-in-progress awaiting confirmatory molecular,

patho-anatomical and prospective clinical data. In particular, discrimination from ARVC should receive due attention.

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