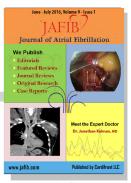


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# Influence Of Novel Electrocardiographic Features Of Provocable Brugada ECG In Arrhythmogenic Cardiomyopathy And Its Exclusion By Lead AVR

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#### Abstract

In 19 patients (14 females, mean age 49.1 ± 11.3 years) with typical arrhythmogenic cardiomyopathy and provocable type I Brugada ECG pattern by ajmaline administration were analysed by novel electrocardiographic features as having "true" or "false" Brugada syndrome. Three patients turned out as having false Brugada syndrome, the diagnosis is pure arrhythmogenic cardiomyopathy. In 16 patients, however, true Brugada syndrome could be provoked. In these patients the diagnosis was arrhythmogenic cardiomyopathy associated by provocable Brugada syndrome.

## Introduction

In a large collective of 385 patients (212 males, mean age 46.3  $\pm$  11.1 years) with typical arrhythmogenic cardiomyopathy lead aVR was analysed. A morphology of large Q wave of 3mm or more, a small R wave of 2mm or less, and T-wave inversion turned out to be the best predictor of arrhythmogenic cardiomyopathy.

In 1498 healthy probands (859 males in an age range of 18 - 85 years) the same morphologic parameters were analysed. Similar results were obtained in 284 probands (18.9%). Specificity and positive predictive value were low, but negative predictive value was nearly 100%.

An association between arrhythmogenic cardiomyopathy and Brugada syndrome seems to be a matter of fact.<sup>1,2,3</sup>

A continuum between these both diseases has been described.<sup>4</sup> Causal gene mutations have been confirmed in plakophilin-2,<sup>5</sup> desmoglein-2<sup>6</sup> and desmoplakin.<sup>7</sup>

To differentiate true or false provocable Brugada syndrome novel electrocardiographic features have been presented <sup>8</sup> as follows:

- concave (coved) ST-segment morphology with negative symmetrical T-waves

- QRS-ST at least 2mm high in lead V1

## Key Words:

ECG, AVR, Cardiomyopathy.

Disclosures: None.

Corresponding Author: Stefan Peters Chief Cardiologist St.Elisabeth Hospital gGmbH Salzgitter Liebenhaller Str. 20 38259 Salzgitter Germany. - ST-segment morphology shows progressive decline

- the ratio between the peak height of QRS-ST after 80ms is greater than  $1\,$ 

- the duration of the QRS in leads V1 and V2 is greater than in the middle and left precordial leads

- type-1 Brugada syndrome ECG may be seen in a single lead, V1 or V2, but never exclusively in V3

We analysed 19 patients (14 females, mean age  $49.1 \pm 11.3$  years) with typical diagnosis of arrhythmogenic cardiomyopathy and provocable Brugada syndrome by ajmaline administration.

In three patients without novel electrocardiographic criteria we could rule out true provocable type I Brugada –ECG pattern. These patients ends up in the diagnosis of pure arrhythmogenic cardiomyopathy.

In 16 patients with novel electrocardiographic criteria demonstrated true provocable type I Brugada ECG pattern. These patients ends up in a combination of arrhythmogenic cardiomyopathy and Brugada syndrome supporting the continuum between these two cardiac entities.

In order to diagnose or to exclude arrhythmogenic cardiomyopathy we like to focus the interest to lead aVR. Lead aVR is the only lead which points directly to the right ventricle.

In a large collective of 385 patients (212 males, mean age 46.3  $\pm$  11.1 years) with typical arrhythmogenic cardiomyopathy the morphology of lead aVR was analysed.

In 97% of cases large Q wave of 3mm or more, small R wave of 2mm or less and T-wave inversion were found. In a control collective of the University Hospital of Glasgow, U.K. (Prof. Peter Macfarlane, Cardiology and Electrocardiography) of 1498 probands (859 males in an age range of 18 – 85 years) the same morphologic parameters were analysed. Similar results were obtained in 284 healthy probands

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## (18.9%).

Specificity and positive predictive value were low, but negative predictive value was nearly 100% (9).

Lead aVR is an excellent tool to exclude arrhythmogenic cardiomyopathy as demonstrated in different publications.<sup>10,11</sup>

#### Conclusions

In conclusion, there are new tools to confirm provocable true Brugada syndrome in arrhythmogenic cardiomyopathy to reveal a continuum between these two entities and to definitely exclude arrhythmogenic cardiomyopathy by electrocardiographic means.

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