Case Report

A Novel SCN5A Mutation in a Patient with Coexistence of Brugada Syndrome Traits and Ischaemic Heart Disease

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Received 25 June 2009; Accepted 11 September 2009

Recommended by Dierk Thomas

Brugada syndrome (BrS) is a primary electrical heart disease, which can lead to sudden cardiac death. In older patients with BrS, the disease may coexist with ischaemic heart disease (IHD) and recent studies support a synergistic proarrhythmic effect of the two disease entities. We report a case that illustrates this. The index patient was a middle-aged patient with BrS traits, IHD, and aborted sudden cardiac death. Mutation analysis discovered a novel mutation P468L in the NaV1.5 sodium channel. Surprisingly, voltage-clamp experiments on the wild-type and mutant NaV1.5 channels expressed in HEK cells revealed no functional effect of the mutation. In a patient like ours, the distinction between IHD and BrS as the cause of an aborted sudden cardiac death is hard to establish and mounting evidence shows that coexistence of the two may have a synergistic proarrhythmic effect.

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1. Introduction

The Brugada syndrome (BrS) is an inherited primary electrical heart disease. It is characterized by characteristic ST segment elevations in the right precordial leads of the ECG in patients with structurally normal hearts and a high frequency of malignant ventricular arrhythmias [1]. Patients with BrS are predominantly male, and the onset of symptoms is usually around the age of 40 years. The disease is believed to be rather common in South East Asia [2], but is rare in western countries [3]. Arrhythmic events are most common at rest or during sleep [4]. In 15–20% of BrS patients, the syndrome is associated with mutations in the SCN5A gene encoding the cardiac NaV1.5 sodium channel [5–7]. The inheritance is autosomal dominant with varying penetrance. In middle aged and elderly patients the phenotype of BrS and ischaemic heart disease (IHD) may coexists creating a mixed phenotype and hence possible diagnostic difficulties. Furthermore studies support a synergistic effect between BrS and IHD in the development of arrhythmic events [8, 9]. Here we describe a patient with BrS traits, IHD, and a novel missense mutation in the SCN5A gene with an apparently normal function.

2. Methods

2.1. Molecular-Genetic Analysis. DNA sequencing of the translated exons with flanking intron sequences of five genes encoding KCNQ1, KCNH2, KCNE1, KCNE2, and SCN5A was performed [10].

2.2. Functional and Biochemical Analysis of NaV1.5 Channels. HEK293 cells were transiently cotransfected with 0.3 μg WT
or P468L Na\textsubscript{v}1.5 constructs and 0.2 μg of pcDNA3-EGFP. Human Na\textsubscript{v}1.5 in pcDNA3 was a gift from Dr. H. Abriel (Lausanne University).

### 3. Case Presentation

The patient was a 61-year-old Caucasian male with a history of angina pectoris and four prior elective coronary angiographies (CAG) in 1996, 1998, 2002, and 2003; all of these showing atheromatosis but no significant stenosis. He had a long history of cigarette smoking and hypertension.

In December 2006, the patient experienced a syncope followed by chest pain and was transported to the nearest emergency department. Upon arrival he developed ventricular fibrillation, but after two direct current counter shocks sinus rhythm was restored. An electrocardiogram (ECG) taken immediately after resuscitation showed (Figure 1(a)) ST segment elevations in lead V\textsubscript{1} and V\textsubscript{2} with a type-1 BrS configuration in V\textsubscript{1}. Shortly, hereafter the patient was transferred from the local community hospital to a regional hospital to undergo an acute CAG. The CAG demonstrated a 90% stenosis on the circumflex coronary artery, which was treated with Percutaneous Coronary Intervention (PCI). TIMI flow both before and after stenting were 3. Blood samples taken at time of admission showed normal biochemistry, especially normal troponin T and creatinine kinases. Transthoracic echocardiography revealed borderline hypertrophy of the left ventricle but was otherwise normal.

Subsequently the patient underwent an invasive electrophysiological study (EPS) during which ventricular fibrillation was induced. Because of the EPS findings and clinical signs of BrS in conjunction with an episode of ventricular fibrillation, the patient received an implantable cardioverter-defibrillator (ICD). The CAG demonstrated a 90% stenosis on the circumflex coronary artery, which was treated with Percutaneous Coronary Intervention (PCI). TIMI flow both before and after stenting were 3. Blood samples taken at time of admission showed normal biochemistry, especially normal troponin T and creatinine kinases. Transthoracic echocardiography revealed borderline hypertrophy of the left ventricle but was otherwise normal.

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The genetic analysis revealed that the patient was heterozygous for a novel missense mutation P468L (c.1403C > T) in exon 11 of the SCN5A gene. The mutation was not identified in a group of 630 unrelated subjects referred for genetic testing. The patient did not have the H558R polymorphism.

3.1. Genetic Analysis. The genetic analysis revealed that the patient was heterozygous for a novel missense mutation P468L (c.1403C > T) in exon 11 of the SCN5A gene. The mutation was not identified in a group of 630 unrelated subjects referred for genetic testing. The patient did not have the H558R polymorphism.

3.2. Family History, Clinical Evaluation, and Genetic Testing of Family Members. The father (I:2) died at age 62 from lung cancer (Figure 2). The mother (I:1) died suddenly at age 64 but no cause of death was established. One brother (II:2) died at the age of 28, he had a heart valve operation in the years before his death, but no further information was available. Another brother (II:3) died at age 59. He had a history of ischaemic heart disease, but again no cause of death was known.

The patient had four children, three daughters and one son, in the age 30–42 years. All daughters carried the mutation, but not the son. None have had any known cardiac diseases or symptoms. The clinical examination revealed an incomplete right bundle branch block bearing resemblance to a Type 3 BrS configuration in the 42 year old daughter (III:4) ECG and a flecainide challenge was carried out but came out negative. ECGs from the last three children were normal.

3.3. Electrophysiological Characterization of Na\textsubscript{v}1.5 P468L. The peak current density did not reveal any difference between wild type and P468L mutant channels. To study the onset of fast inactivation, individual current traces...
were fitted using a biexponential function and the resulting time constants (τ) were plotted as a function of voltage. No significant differences in the τfast or τslow values of mutant channel compared to wild type were found. To further investigate the phenotype of Nav1.5 P468L, steady-state activation and inactivation were analyzed, as well as the voltage- and time-dependent release from inactivation (Table 1). Again, there was no difference between wild type and P468L channels.

### 4. Discussion

Cardiac enzymes after the cardiac arrest were normal, hence acute myocardial infarction (AMI) may be excluded, but transient ischemia cannot. The patients never exhibited diagnostic ECG changes for BrS, as a type-1 configuration was only seen in one lead and not the required two. In the clinical setting the patient was treated as having BrS though.

The patient experienced two syncopies years prior to the cardiac arrest. The cause of the syncopies could be vasovagal, but self-limiting episodes of ventricular tachycardia cannot be ruled out.

Genetic screening revealed a novel mutation in SCN5A. The mutation was found in 3 out of 4 of the progeny, but none had a BrS phenotype. The mutation not being present in 630 controls speaks in favor of the P468L substitution being the underlying cause of the BrS traits in the patient, but the lack of genotype/phenotype cosegregation in the family weakens the argument. The electrophysiological investigations of the mutation did not reveal any functional effect of the P468L substitution, and the substitution could be an innocent bystander instead of disease causing. However, the mutated sodium channels expressed in HEK-293 cells may behave differently from in vivo due to lack of sodium channel interacting proteins, different trafficking or existence of a different set of polymorphisms.

Coexistence of BrS traits and IHD in a patient with ASCD can pose a problem in the clinical setting since the latter represents a potentially reversible cause of heart arrest and the former does not. Future ischemic events can in many cases be prevented by appropriate interventions whereas with the diagnosis of BrS and an episode of ASCD an ICD implantation is mandated [5].

As to the pathogenesis of the ASCD in this patient, the distinction between IHD and BrS as the substrate of sudden cardiac death is muddled by a number of reports showing that BrS increases the risk for arrhythmias during ischemia [8, 9] although not all studies support this conclusion [11].

The suggested mechanism for this accentuation of the risk of arrhythmias is the difference in the ion current balance (I_{Na,I}/I_{Na,I}/I_{Ca,L}) in epicardial versus endocardial cells of especially the right ventricular outflow tract. A balance that, according to this theory, is disturbed in BrS as well as during ischemia and the two in combination may have a synergistic proarrhythmic effect [12]. Furthermore, evidence is mounting that polymorphisms in SCN5A exons as well as in promoter regions has an impact on SCD susceptibility, albeit not specifically in the setting of IHD, but one could speculate that ischemia as well as these genetic changes both reduce the “antifibrillatory reserve” in a synergistic manner [13].

Currently many BrS patients are middle aged [14], and consequently we will in the near future see a large cohort of BrS patients coming into an age where IHD becomes prevalent and thus be facing the problems reported in this study. Further research into the modulating effect of BrS on the risk of SCD in the setting of IHD is warranted, as this could potentially lead to changes in risk stratification for BrS patients with high risk of IHD, as well as a better understanding of the arrhythmia inducing mechanisms underlying both.

In conclusion, we reported a case of aborted sudden cardiac death in a patient with coexistence of Brugada Syndrome traits, ischaemic heart disease, and a novel SCN5A mutation. The distinction between IHD and BrS as the cause of an aborted sudden cardiac death is hard to delineate, and is muddled by mounting evidence showing that coexistence of the two may have a synergistic proarrhythmic effect.

### Acknowledgments

This work was supported by the Foundation of 17-12-1981, the John and Birthe Meyer Foundation, and the Danish National Research Foundation. We would like to thank the patient and his family members who participated in this study.

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