Mutations in Desmoglein-2 Gene Are Associated With Arrhythmogenic Right Ventricular Cardiomyopathy

Kalliopi Pilichou, BSc; Andrea Nava, MD; Cristina Basso, MD, PhD; Giorgia Beffagna, BSc, PhD; Barbara Bauce, MD, PhD; Alessandra Lorenzon, BSc; Gianfranco Frigo, MD, PhD; Andrea Vettori, BSc, PhD; Marialuisa Valente, MD; Jeffrey Towbin, MD; Gaetano Thiene, MD; Gian Antonio Danieli, BSc; Alessandra Rampazzo, BSc, PhD

Background—Arrhythmogenic right ventricular cardiomyopathy (ARVC) is an inherited cardiomyopathy characterized by progressive myocardial atrophy with fibrofatty replacement. The recent identification of causative mutations in plakoglobin, desmoplakin (DSP), and plakophilin-2 (PKP2) genes led to the hypothesis that ARVC is due to desmosomal defects. Therefore, desmoglein-2 (DSG2), the only desmoglein isoform expressed in cardiac myocytes, was screened in subjects with ARVC.

Methods and Results—In a series of 80 unrelated ARVC probands, 26 carried a mutation in DSP (16%), PKP2 (14%), and transforming growth factor-β3 (2.5%) genes; the remaining 54 were screened for DSG2 mutations by denaturing high-performance liquid chromatography and direct sequencing. Nine heterozygous DSG2 mutations (5 missense, 2 insertion-deletions, 1 nonsense, and 1 splice site mutation) were detected in 8 probands (10%). All probands fulfilled task force criteria for ARVC. An endomyocardial biopsy was obtained in 5, showing extensive loss of myocytes with fibrofatty tissue replacement. In 3 patients, electron microscopy investigation was performed, showing intercalated disc paleness, decreased desmosome number, and intercellular gap widening.

Conclusions—This is the first investigation demonstrating DSG2 gene mutations in a significant number of ARVC-unrelated probands. Cardiac phenotype is characterized clinically by typical ARVC features with frequent left ventricular involvement and morphologically by fibrofatty myocardial replacement and desmosomal remodeling. The presence of mutations in desmosomal encoding genes in 40% of cases confirms that many forms of ARVC are due to alterations in the desmosome complex. (Circulation. 2006;113:1171-1179.)

Key Words: arrhythmogenic right ventricular dysplasia ■ cell adhesion molecules ■ genetics ■ pathology ■ sudden death

Arrhythmogenic right ventricular cardiomyopathy (ARVC) is a myocardial disease characterized by progressive myocyte loss and fibrous and fatty tissue replacement of the right ventricular free wall, which is the substrate for reentrant arrhythmias and sudden death. Familial occurrence is common. Five disease genes have been identified thus far that encode desmosomal proteins, except for the cardiac ryanodine receptor 2 (RyR2) gene reported in arrhythmogenic right ventricular cardiomyopathy type 2 (ARVD2) and the transforming growth factor-β3 (TGFβ3) gene in arrhythmogenic right ventricular cardiomyopathy type 1 (ARVD1). A deletion in plakoglobin has been proven to cause a recessive form of ARVC associated with palmoplantar keratosis and woolly hair, ie, Naxos disease. More recently, mutations of desmoplakin (DSP) and plakophilin-2 (PKP2) genes have been found in ARVC, in the absence of skin and hair abnormalities. Of note, a homozygous desmoplakin mutation has also been reported in Carvajal syndrome, which is characterized by hair and skin disorders associated with cardiac disease. Desmosomal adhesion is mediated by calcium-dependent cell adhesion glycoproteins (cadherins) of the desmocollin (DSC) and desmoglein (DSG) types, which interact later...
ally and transcellularly. Both desmosomal cadherins types are encoded by a small multigene family on chromosome 18q12.1, each of which consists of cell type–restricted members. Unlike other cadherins that are restricted to stratified epithelial tissues, DSG2 and DSC2 are expressed in all desmosome-possessing tissues, including myocardium and epithelia.

We report here for the first time mutations of the DSG2 gene associated with ARVC and clinicopathological data of the 8 probands in whom such mutations were detected in the course of a systematic mutation screening.

**Methods**

**Clinical Evaluation**

The study was approved by the University of Padua Medical School–Azienda Ospedaliera ethical committee. Informed consent was obtained from all participating individuals. Clinical evaluation consisted of a detailed personal/family history, physical examination, 12-lead ECG, 2-dimensional echocardiogram, signal-averaged ECG (SAECG), and stress test ECG, performed according to previously reported methods. Invasive studies including angiography and right ventricular endomyocardial biopsy were performed in selected cases when deemed necessary for the diagnosis. In 80 probands of Italian descent, a clinical diagnosis of ARVC was made on the basis of the established European Society of Cardiology/International Society and Federation of Cardiology Task Force major and minor criteria.

**Mutation Screening**

The coding region of DSP, PKP2, and TGFβ3 genes was screened for mutations in all study subjects. Because none of the subjects had effort-induced polymorphic ventricular arrhythmias or gross skin and hair abnormalities, we chose not to screen RyR2 or plakoglobin. Thirteen probands (16%) carried a DSP mutation, 11 (14%) a PKP2 mutation, and 2 (2.5%) a TGFβ3 mutation. Fifty-four probands (33 males and 21 females) negative for mutations of these genes were considered for further investigation.

On the basis of previous findings on desmosomal genes involved in ARVC and on DSG2 expression in myocardial tissue, we hypothesized that mutations in human DSG2 could account for ARVC. The 54 ARVC probands were screened for DSG2 mutations by denaturing high-performance liquid chromatography (DHPLC) and direct sequencing. Polymerase chain reaction (PCR) primers flanking each exon of the human DSG2 gene were designed by PRIMER3. PCR amplifications were performed in a final volume of 25 μL, containing 50 ng of genomic DNA, 1 μmol/L MgCl2 (Applied Biosystems), 400 nmol/L each primer (Sigma Genosys), 100 μmol/L deoxynucleotide triphosphates (Invitrogen), and 0.8 U of Taq DNA polymerase (Taq Gold, Applied Biosystems). Cycling conditions (denaturation at 94°C for 30 sec, annealing at 60°C for 30 seconds, and extension at 72°C for 45 seconds) were repeated for 35 cycles. PCR primers and DHPLC conditions are reported in Table 1. DHPLC analysis was performed with the use of WAVE Nucleic Acid Fragment Analysis System 3500HT with DNASep HT cartridge (Transgenomic Ltd NE). Temperatures for sample analysis were selected with the use of WAVEMAKER software. The gradient mobile phase consisted of buffer A (0.1 mol/L triethyl ammonium

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TABLE 1. PCR Primers and Conditions for DHPLC Analysis of DSG2 Gene
TABLE 2. Clinical, Pathological, and Genetic Findings in DSG2 Mutation Unrelated Carriers

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<th>Family History</th>
<th>12-Lead ECG</th>
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<th>Arrhythmias</th>
<th>RV Size/Function</th>
<th>Amino Acid Change</th>
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<td>Negative T Waves</td>
<td>Negative GS in RV</td>
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<td>LV Involvement</td>
<td>Diagnostic Criteria</td>
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<td>Leads aVF</td>
<td>SAEDG</td>
<td>Nor</td>
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<td>42</td>
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<td>2</td>
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<td>V1 through V3</td>
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<td>M</td>
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<td>V1 through V3</td>
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<td>5</td>
<td>F</td>
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<td>-</td>
<td>-</td>
<td>V1 through V3</td>
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<td>(172)</td>
<td></td>
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<tr>
<td>6</td>
<td>F</td>
<td>55</td>
<td>-</td>
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<td>V1 through V3</td>
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LBBB indicates left bundle-branch block; LV, left ventricular; NP, not performed; NSVT, nonsustained ventricular tachycardia; RV, right ventricular; SVT, sustained ventricular tachycardia; and VF, ventricular fibrillation.

Mutations in Desmoglein-2

Nine DSG2 mutations have been identified in 8 (5 males and 3 females; mean age at diagnosis/symptom onset, 38±20 years; range, 11 to 63 years) of 80 ARVC index cases (10%).
Overall, DSP, PKP2, TGFβ3, and DSG2 account for 42.5% of genotyped ARVC probands.

Of the 9 DSG2 mutations, 5 were missense, 2 insertion-deletions, 1 a nonsense, and 1 a splice site mutation (Figure 1). None of the detected nucleotide changes was found in 560 control chromosomes. Identified mutations altering restriction sites were confirmed by restriction digest. The PCR products were submitted to digestion with the following restriction enzymes: HpyCH4V for the Tyr87Cys, PflmI for Gly100Arg, HpyAV for Glu331Lys, AlwNI for Gln557Ter, and BsmAI for 1253_1257insATGA. Missense mutations Y87C, G100R, N266S, K294E, and E331K occurred in residues that are highly conserved among species. Mutations Y87C, G100R, and N266S, which are also conserved in all the desmogleins, and K294E and E331K are located in the extracellular cadherin (EC) domains. Cadherin domains are important for homophilic intercellular associations and form Ca^{2+}-dependent rodlike structures. All the inherent amino acid changes may destabilize the rod structure and influence the homophilic binding. Moreover, mutation N266S is localized in a putative calcium binding site (DXNDN).

The 4-bp insertion in exon 9 (1253_1257insATGA) causes the addition of an amino acid residue before a premature stop signal is introduced (E418fsX419); the predicted truncated DSG2 molecule would lack transmembrane and cytoplasmic components.

The single base pair deletion in exon 14 (2036delG) results in a frameshift mutation that would leave the first 678 amino acids of the DSG2 protein intact, followed by addition of 2 novel amino acids (G678fsX681). The predicted truncated protein should maintain the transmembrane domain, but it would lack the intracellular cadherin-typical segment, thus precluding normal interaction with plakoglobin.

Mutation 1672C→T generates a termination codon (Q558X); therefore, the mutant should encode a short DSG2 molecule lacking transmembrane and cytoplasmic domains.

Mutation 1881 to 2A→G affects the acceptor splice site of intron 12. Sequencing analysis of the aberrant DSG2 transcript obtained from lymphocyte RNA of the proband showed that this mutation activates an alternative cryptic splice site in exon 13, located 38 bp downstream from the authentic 3′ splice acceptor site. This aberrant spliced mRNA contains a 38-bp deletion (Figure 2) and should code for a truncated protein of 646 amino acids in length, missing the cytoplasmic domain.

In 1 patient (patient 5), 2 different mutations (E331K and 1881 to 2A→G) were detected. Family members were available for molecular analysis (Figure 3, family 172).
Subject II,1 carried both mutations, subjects I,2 and III,2 carried the missense mutation, and subject III,1 carried the splice site mutation, thus demonstrating that the 2 mutations are in trans orientation. Although both mutations are potentially pathogenic, the possibility that the missense mutation is a rare polymorphism cannot be ruled out.

Clinical Findings

Clinical Presentation

The first symptom consisted of sustained ventricular tachycardia (VT) in 3 patients, palpitations in 3, and chest pain with increased serum markers of myocardial necrosis (creatine phosphokinase/MB 363/45 U/L and troponin I 11.9 μg/L) in 1 who had angiographically normal coronary arteries. One patient was asymptomatic and was examined because ECG abnormalities were detected at preparticipation screening for sport activity. Skin and hair were grossly normal at physical examination in all. Mean age at symptom onset/diagnosis was 26 ± 13 years in males versus 58 ± 4 years in females (P = 0.001).

ECG/SAECCG Findings

All patients showed ECG abnormalities (Table 2) (Figure 4A), consisting of complete right bundle-branch block in 1, incomplete right bundle-branch block in 3, negative T waves in the precordial leads in 7, negative T waves in inferior leads in 3, epsilon wave in 3, low voltages of QRS in 1, and ST segment elevation in 1. Moreover, 3 patients showed a PQ prolongation (PQ > 200 ms).

Late potentials were detected in 7 subjects and were evidenced with 25-, 40-, and 80-Hz filters in 4 cases and with 40- and 80-Hz filters in 3 cases.

Ventricular Arrhythmias

Ventricular arrhythmias were recorded in all probands and ranged from sustained VT with left bundle-branch block morphology (3 cases) to nonsustained VT (3 cases) and isolated monomorphic PVCs (2 cases).

Echocardiographic Findings

Abnormal echocardiographic findings were present in all probands, with right ventricular kinetic abnormalities involving only 1 region in 1 patient and ≥ 2 regions in the remaining 7 (Figure 4B). A left ventricular involvement was present in 4 patients, with normal left ventricular ejection fraction in 1 (57%) and mildly decreased left ventricular ejection fraction in 3 (ranging from 45% to 48%); left ventricular kinetic abnormalities were localized in 1 and diffuse in 3 patients.

Familial Study

A family history of premature sudden death due to suspected ARVC was present in 1 proband and of clinically proven ARVC (based on diagnostic criteria) in 3 probands. Family members of 2 index cases (patient 5, family 172 and patient 6, family 169) were available for clinical and genetic testing (Figure 3). In family 172, subject II,1, who
was found to carry both mutations, showed negative T waves in V1 through V4 on 12-lead ECG, right ventricular enlargement and kinetic alterations on 2D echocardiogram, and premature ventricular beats (>1000/24 h) with left bundle-branch block morphology. Clinical investigation in subjects I,2 and III,2 was negative, whereas it was not feasible in subject III,1.

In family 169, the DSG2 mutation N266S was found in a daughter (subject II,1) and a son (subject II,4) of the index case. Clinical findings were negative in the former, whereas subject II,4 showed negative T waves in V1 through V2 on 12-lead ECG, right ventricular dilation together with kinetic abnormalities on 2D echo, and nonsustained ventricular arrhythmias.

All family members not carrying DSG2 mutations were negative at clinical investigation.

**Light Microscopy and Electron Microscopy Findings**

In the 5 patients who underwent endomyocardial biopsy (patients 1, 2, 5, 8, 9), histomorphometric evaluation revealed a mean area of residual myocardium 47±8%, fibrous tissue 24±11%, and fatty tissue 20±13%. Moreover, dysmetric and dysmorphic nuclei and prominent cytoplasmic vacuolization were evidenced in all (Figure 4C and 4D).

In 3 patients (patients 1, 2, 8), electron microscopy investigation was also performed. At the ultrastructural level, we found a decreased desmosome number per 10 μm unity length of intercalated disc (3.1±0.4 versus 5.5±3 in controls) and an increased desmosome gap (31.7±17.8 versus 21.7±3.4 nm in controls). Moreover, abnormal small junctions composed of series of repeating couplings, abnormally located desmosomes, and pale internal plaques were visible compared with controls (Figure 5).

**Discussion**

We identified DSG2 as a novel disease gene involved in ARVC, a genetically determined myocardial disease recognized as a common cause of sudden death among young adults. No association of DSG2 mutations with inherited human diseases has been reported thus far. Desmosomes consist of 2 desmosomal-specific cadherin family members, DSGs and DSCs, as well as a collection of cytoplasmic plaque proteins including plakoglobin, DSP, and PKPs. All cadherins have tripartite functional domains: (1)
a calcium-inducible, extracellular amino terminal domain, important for homophilic intercellular associations, with 4 domains (EC1 to EC4), followed by an extracellular anchor domain (EA); (2) a single transmembrane domain; and (3) a cytoplasmic domain anchoring the cytoskeleton, an essential process for cell adhesion.24,25

Among the 9 mutations herein reported, 7 are located in the EC amino terminal domain, which directly participates in the adhesive interaction. Crystallographic and biochemical studies of classic cadherin ectodomains suggest that they form homophilic dimers in cis and in trans orientations.26 Calcium binding to the EC domains is important for stabilizing their structure and function. Whereas trans-dimerization is likely required to form the adhesive interface in adjacent cells, it has been proposed that cis-dimerization is important for promoting adhesion via molecular clustering.26 It is possible that even a single amino acid change could result in differences in molecular affinity and possibly in the abolition of the adhesive capacity of cadherins. In vitro functional studies are needed to evaluate this.

Effect of the DSG2 Mutations at the Cellular Level

Unlike PKP2 and plakoglobin, no data are available on cardiac abnormalities in the DSG2 knockout mouse because loss of DSG2 resulted in early embryonic lethality.27 In vitro antisense experiments against DSC2,28 expression of truncated DSG3 and DSG1 proteins,29,30 and knockout mice of the DSG3 gene31 all resulted in a decreased number of desmosomes, associated with increased asymmetry and detachment. These electron microscopy observations are in agreement with those obtained in 3 DSG2 mutations carriers herein reported, which show a decreased number of desmosomes as well as a widening of intercalated disc gap. On the basis of these ultrastructural findings and of the histological demonstration of myocardial atrophy in patients who underwent endomyocardial biopsy, we can postulate that in desmosomal cardiomyopathies (ie, Naxos syndrome, Carvajal syndrome, and ARVC),8,10–12,14–16 impaired cell-to-cell adhesion leads to myocyte detachment, cell death, and fibrofatty repair.17,32 Moreover, because TGFβ3 could modulate expression of genes encoding desmosomal proteins in different cell types,33 the same pathogenic mechanism could be advocated in ARVD1 patients.12

From the clinical point of view, in our series of unrelated DSG2 mutation carriers, a typical form of ARVC was documented with left ventricular involvement in almost half of the cases. Genotyping of available family members of 2 index cases allowed us to demonstrate the transmission of the disease allele in all affected relatives. However, prospective studies of a larger number of family members are needed to evaluate the penetrance of DSG2 mutations and the entire clinical spectrum of the disease from the concealed to the overt forms. It is noteworthy that both sexes are represented equally, although the mean age at symptom onset/diagnosis was significantly higher in female than in male mutation carriers. These data confirm previous observations from clinical series and could be interpreted either as a gender-related penetrance or as the consequence of additional acquired and/or genetic factors involved in disease expression.13

None of the 8 probands carrying the pathogenic mutations showed gross skin/hair abnormalities. All DSG isoforms are expressed in the epidermal tissues, whereas DSG2 is the only one expressed in the myocardium.32 We may hypothesize that compensation by other DSG isoforms might take place in the epidermis but not in the myocardium of DSG2 mutation carriers, thus accounting for the cardiac-specific phenotype.

Clinical diagnosis of ARVC is challenging, and the currently available diagnostic criteria, although acknowledged to be specific, are lacking in sensitivity. Because mutation detection is feasible in a significant proportion of patients, genetic screening could assume a pivotal role in the clinical evaluation of familial forms. In this setting, the benefits of detection of disease-gene mutations would include identification of asymptomatic carriers among family members and interpretation of borderline clinical phenotypes.

In conclusion, this is the first demonstration that mutations in DSG2 gene are associated with ARVC. On the basis of our data showing that 40% of ARVC probands carry a mutation in desmosomal proteins encoding genes,
we confirm that many forms of ARVC are due to alterations in the desmosome complex.

Acknowledgments
This study was supported by the Ministry of Health, MIUR, and Telethon grant GGP05261, Rome; National Institutes of Health grant U04HL 65652; ARVC/D Project, QLG1-CT-2000-01091, Fifth Framework Programme European Commission, Bruxelles, and Fondazione Cassa di Risparmio, Padova e Rovigo. Dr Bause is a recipient of a research temporary position pursuant to the ARVC/D Project, European Commission. The authors are deeply indebted to Paola Marcon for her help in collecting ARVC families and to Mila Della Barbera, BSc, PhD, for her help in electron microscopy investigation.

Disclosures
None.

References
The recent identification of causative mutations in plakoglobin, desmoplakin, and plakophilin-2 genes in arrhythmogenic right ventricular cardiomyopathy (ARVC) led to the hypothesis that many forms of ARVC are due to alterations in the desmosome complex. In this article, the authors demonstrate for the first time that mutations in desmoglein-2, the only desmoglein isoform reported thus far as expressed in cardiac myocytes, are associated with ARVC. With this new discovery, mutation detection by genetic screening is now achievable in a significant proportion of probands affected by ARVC. In the clinical evaluation of family members, genetic screening is assuming a pivotal role, allowing the early identification of asymptomatic carriers and interpretation of borderline clinical phenotypes. On the other hand, the identification of a growing number of asymptomatic carriers with mild or subclinical disease expression leads to new problems in clinical management. When one takes into account that sudden death is not so rarely the first manifestation of ARVC, risk stratification remains a clinical challenge, and noninvasive markers able to predict the risk of life-threatening ventricular arrhythmias in mutation carriers should be prospectively evaluated in international registries.
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_Circulation_. 2006;113:1171-1179; originally published online February 27, 2006;
doi: 10.1161/CIRCULATIONAHA.105.583674

_Circulation_ is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
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Print ISSN: 0009-7322. Online ISSN: 1524-4539

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