Normalization of Naxos plakoglobin levels restores cardiac function in mice

Zhiwei Zhang, … , Xinmin Zhou, Ju Chen


Arrhythmogenic cardiomyopathy (AC) is associated with mutations in genes encoding intercalated disc proteins and ultimately results in sudden cardiac death. A subset of patients with AC have the autosomal recessive cardiocutaneous disorder Naxos disease, which is caused by a 2-base pair deletion in the plakoglobin-encoding gene _JUP_ that results in a truncated protein with reduced expression. In mice, cardiomyocyte-specific plakoglobin deficiency recapitulates many aspects of human AC, and overexpression of the truncated Naxos-associated plakoglobin also results in an AC-like phenotype; therefore, it is unclear whether Naxos disease results from loss or gain of function consequent to the plakoglobin mutation. Here, we generated 2 knockin mouse models in which endogenous _Jup_ was engineered to express the Naxos-associated form of plakoglobin. In one model, Naxos plakoglobin bypassed the nonsense-mediated mRNA decay pathway, resulting in normal levels of the truncated plakoglobin. Moreover, restoration of Naxos plakoglobin to WT levels resulted in normal heart function. Together, these data indicate that a gain of function in the truncated form of the protein does not underlie the clinical phenotype of patients with Naxos disease and instead suggest that insufficiency of the truncated Naxos plakoglobin accounts for disease manifestation. Moreover, these results suggest that increasing levels of truncated or WT plakoglobin has potential as a therapeutic approach to Naxos disease.

Find the latest version:

https://jci.me/80335/pdf
Normalization of Naxos plakoglobin levels restores cardiac function in mice

Zhiwei Zhang,1,2 Matthew J. Stroud,1 Jianlin Zhang,1 Xi Fang,1 Kunfu Ouyang,3 Kensuke Kimura,1 Yongxin Mu,1 Nancy D. Dalton,1 Yusu Gu,1 William H. Bradford,1 Kirk L. Peterson,1 Hongqiang Cheng,2 Xinmin Zhou,2 and Ju Chen1

1School of Medicine, UCSD, La Jolla, California, USA. 2Department of Cardiothoracic Surgery, The Second Xiangya Hospital, Central South University, Changsha, Hunan, China. 3School of Chemical Biology and Biotechnology, Peking University, Shenzhen, China. cDepartment of Pathology and Pathophysiology, Program in Molecular Cell Biology, Zhejiang University School of Medicine, Hangzhou, China.

Abstract

Arrhythmogenic cardiomyopathy (AC) is associated with mutations in genes encoding intercalated disc proteins and ultimately results in sudden cardiac death. A subset of patients with AC have the autosomal recessive cardiocutaneous disorder Naxos disease, which is caused by a 2-base pair deletion in the plakoglobin-encoding gene JUP that results in a truncated protein with reduced expression. In mice, cardiomyocyte-specific plakoglobin deficiency recapitulates many aspects of human AC, and overexpression of the truncated Naxos-associated plakoglobin also results in an AC-like phenotype; therefore, it is unclear whether Naxos disease results from loss or gain of function consequent to the plakoglobin mutation. Here, we generated 2 knockin mouse models in which endogenous Jup was engineered to express the Naxos-associated form of plakoglobin. In one model, Naxos plakoglobin bypassed the nonsense-mediated mRNA decay pathway, resulting in normal levels of the truncated plakoglobin. Moreover, restoration of Naxos plakoglobin to WT levels resulted in normal heart function. Together, these data indicate that a gain of function in the truncated form of the protein does not underlie the clinical phenotype of patients with Naxos disease and instead suggest that insufficiency of the truncated Naxos plakoglobin accounts for disease manifestation. Moreover, these results suggest that increasing levels of truncated or WT plakoglobin has potential as a therapeutic approach to Naxos disease.

Introduction

Arrhythmogenic cardiomyopathy (AC) (1–3) is observed in Naxos disease, which is caused by a 2-base pair deletion in the JUP gene encoding the intercalated disc (ICD) protein plakoglobin (1). This mutation causes a frameshift, resulting in a premature stop codon, and expression of a truncated plakoglobin lacking 56 residues from the C terminus. Naxos disease is autosomal recessive, suggesting that truncated plakoglobin is unable to function as its full-length counterpart. Studies to model Naxos disease have yielded different conclusions.

To investigate whether loss of plakoglobin in cardiomyocytes recapitulates Naxos cardiac disease, several mouse models have been generated. Ablation of plakoglobin in adult cardiomyocytes recapitulated many features of Naxos disease, including progressive cardiomyocyte loss, excessive inflammation, and fibrosis. Furthermore, desmosomal ultrastructure and cardiac function were affected in these mice (4). Ablation of plakoglobin in fetal cardiomyocytes resulted in mice with a more severe Naxos-like cardiac phenotype, with cardiac dysfunction, fibrosis, ventricular thinning, and spontaneous ventricular arrhythmias (5). These mouse models, and the recessive nature of Naxos disease, suggest that the disorder is governed by a loss of function (LOF) of plakoglobin.

In addition to LOF mouse models, gain-of-function (GOF) transgenic mice and zebrafish overexpressing the truncated plakoglobin have been generated (6, 7). In GOF models, many characteristic signs of Naxos cardiomyopathy are also observed, including fibrofatty replacement of cardiomyocytes. Overexpressed mutant plakoglobin, in vivo in cardiomyocytes (6, 7) or in cultured neonatal rat ventricular myocytes, localized to the nucleus (6, 7) and interfered with β-catenin signaling (6, 7). Currently, there are no data from cardiac tissue from patients with Naxos disease that address the extent to which the mutant plakoglobin protein is reduced or its subcellular localization changed.

To gain insight into mechanisms underlying Naxos disease, we used mouse genetics to generate 2 mouse models in which the Jup gene was modified to recapitulate the Naxos mutation. Mice that bypassed nonsense-mediated decay to express truncated plakoglobin at levels comparable to those of full-length WT protein did not display any phenotype characteristic of Naxos disease. Our data demonstrated that GOF of the mutant protein does not underlie the clinical phenotype of patients with Naxos disease, leaving the alternate explanation that LOF of Naxos protein accounts for disease manifestation. An important implication of our findings is that a viable therapeutic approach to Naxos disease will be to increase levels of truncated or WT plakoglobin protein.

Results and Discussion

A 2-base pair deletion in both alleles of the JUP gene encoding plakoglobin causes Naxos disease in humans (1). To investigate this disease in mice and to exactly mimic the mutation found in
human patients with Naxos disease, we generated a mouse line called original Naxos (OriNax), in which 2 base pairs were deleted to result in a frameshift and premature stop codon (Supplemental Figure 1, A–C; supplemental material available online with this article; doi:10.1172/JCI80335DS1).

OriNax mice were born at expected Mendelian ratios (24.2% OriNax mice compared with 25% expected from 190 pups from heterozygote crosses) but died on postnatal day 1 (PD1) (Figure 1A). Plakoglobin mRNA levels were significantly downregulated in OriNax mice compared with those in WT littermates (Figure 1B). This was because the frameshift mutation introduced in exon 11 of the jup gene results in a premature termination codon in exon 12 that is recognized by a translation-coupled surveillance pathway termed nonsense-mediated RNA decay. This pathway leads to the specific degradation of plakoglobin mRNA (8). In accordance with the lower mRNA levels, plakoglobin protein levels were markedly downregulated in OriNax mice compared with those in WT littermates (Figure 1C).

Masson’s Trichrome staining revealed no loss of cardiomyocytes, fibrosis, thinning of the ventricles, and fatty deposits in PD1 hearts from OriNax mutants when compared with littermate controls (Figure 1D). Consistent with the absence of fatty deposits, we observed no changes in the levels of proadipogenic genes, such as lipoprotein lipase or adiponectin (Figure 1E). Furthermore, we observed no changes in heart weight/body weight ratios (Figure 1F) or collagen 1A or proinflammatory genes Tgfb1 or Nfkβ (Figure 1E). Global plakoglobin knockout mice die from cardiac defects at around embryonic day 13 (9, 10). Despite OriNax mice having low levels of plakoglobin, they appear to have a sufficient amount for normal cardiac development. However, the skin of OriNax mice was more fragile and wrinkled than that of WT controls. Furthermore, Masson’s Trichrome staining of skin revealed that the epidermis of OriNax mice was thinner and looser than that of their WT littermates (Supplemental Figure 1D).

Overall, OriNax mice died soon after birth and did not display characteristic hallmarks of AC. The lack of cardiac phenotype is not too surprising when understood in the context of the progression of Naxos disease, in which cardiac phenotype is age dependent (2). OriNax mutant perinatal lethality has also been observed in 50% of mice that are hypomorphic for WT plakoglobin, expressing approximately 40% of normal levels of WT plakoglobin (14). Perinatal lethality has not been reported for patients with Naxos disease. Reasons for this discrepancy are not clear but may reflect
cyte loss in FuseNax mice (Figure 2A). Cardiac remodeling is often accompanied by activation of the fetal gene program (15). Consistent with our histological data, we found no changes in the fetal gene program (Figure 2B). Furthermore, we observed no differences in cardiac function, heart weight/body weight ratio, chamber size, or wall thickness between FuseNax mice and their WT littermates at 47 weeks of age and older (Figure 2, C–F, and data not shown). In contrast to OriNax mice, no skin or hair abnormalities were observed in FuseNax mice (data not shown).

Plakoglobin plays a fundamental role at the ICD; therefore, we sought to understand whether expression of truncated plakoglobin would affect other ICD proteins. Both subcellular localization and expression levels of β-catenin and connexin 43 appeared normal in myocardium of FuseNax mice when compared with WT littermates (Supplemental Figure 3, A and B) and expressed the truncated plakoglobin at the same level as full-length plakoglobin (see below and Supplemental Figure 3C).

Pathological cardiac remodeling and cardiac dysfunction were reported in transgenic mice and zebrafish overexpressing truncated plakoglobin, resulting in premature death (6, 7). FuseNax mice were born at expected Mendelian ratios, surviving with no overt morphological phenotype.

Masson’s Trichrome staining of heart sections revealed no gross morphological defects, changes in fibrosis, or cardiomyocyte loss in FuseNax mice (Figure 2A). Cardiac remodeling is often accompanied by activation of the fetal gene program (15). Consistent with our histological data, we found no changes in the fetal gene program (Figure 2B). Furthermore, we observed no differences in cardiac function, heart weight/body weight ratio, chamber size, or wall thickness between FuseNax mice and their WT littermates at 47 weeks of age and older (Figure 2, C–F, and data not shown). In contrast to OriNax mice, no skin or hair abnormalities were observed in FuseNax mice (data not shown).

Plakoglobin plays a fundamental role at the ICD; therefore, we sought to understand whether expression of truncated plakoglobin would affect other ICD proteins. Both subcellular localization and expression levels of β-catenin and connexin 43 appeared normal in myocardium of FuseNax mice when compared with WT littermates (Figure 3, A–D). Furthermore, the majority of ICD proteins were unaffected in FuseNax mice (Figure 3, C–E). However, levels of DSC2, a desmosomal protein that interacts with plakoglobin (16), were reduced to 40% of those of WT mice, as measured by densitometry of the Western blot (Figure 3E).
It has been shown that an overexpressed truncated Naxos plakoglobin protein translocates to the nucleus and interferes with β-catenin signaling (6, 7). To examine whether this nuclear localization occurred in mice expressing truncated plakoglobin under the endogenous promoter, we performed experiments using 2 antibodies generated against different epitopes at the N terminus of plakoglobin. We found no evidence of mutant protein localizing to the nucleus, using either immunofluorescence or biochemical subcellular fractionation approaches (Figure 3, F and G, and Supplemental Figure 3D). Furthermore, genes downstream of β-catenin were unaffected in FuseNax mice (Figure 3H).

Given that FuseNax mice showed no impairment of cardiac contractile function at 11 months of age, we wanted to investigate whether they would exhibit impairment in cardiac contractile function at 11 months of age.
tile function in response to β-adrenergic stimulation. Accordingly, 9-week-old mice were treated with increasing amounts of dobutamine and subjected to hemodynamic analyses. Results demonstrated that FuseNax mice responded similarly to WT littersmates, with no impairment in contractile function in response to β-adrenergic stimulation (Supplemental Figure 4, A–F).

To investigate whether FuseNax mice displayed arrhythmias, we performed telemetry recordings over a 48-hour period. We found no changes in heart rate between FuseNax and WT littersmates (Supplemental Figure 4, G and H). Furthermore, QRS complex duration and PR interval were unaffected (Supplemental Figure 4I). In addition, no ventricular tachycardia was detected.

Previous approaches to understand molecular mechanisms underlying AC in Naxos disease have involved conditional knock-out approaches and transgenic mouse and zebrafish models in which plakoglobin is overexpressed (6, 7). Here, we specifically generated 2 knockin mouse models in which the Jup gene encoding plakoglobin was modified to mimic the Naxos mutation. Our initial OriNax model exhibited perinatal lethality, as observed by Ruiz et al. in a mouse model with significantly reduced levels of WT plakoglobin (9). In FuseNax mice, in which the truncated Naxos plakoglobin is expressed at levels equivalent to the WT protein, we found no impairment of cardiac function, cardiac hypertrophy, or chamber dilation and no evidence for increased fibrosis, fibrofatty replacement, or myofibrillar disarray. These observations demonstrated that WT levels of truncated Naxos plakoglobin in FuseNax mice were sufficient to maintain cardiac function.

It should be pointed out, however, that although hearts appeared to be functionally normal, DSC2 protein levels were consistently reduced in FuseNax hearts, implying that the C terminus of WT plakoglobin may stabilize DSC2 at the ICD, and the truncated protein may not possess the full function of WT plakoglobin.

Overall, our results showed that restoration of truncated Naxos plakoglobin to WT levels resulted in normal cardiac function. These data demonstrate that GOF of mutant Naxos protein does not underlie the clinical phenotype of patients with Naxos disease, leaving the alternate explanation that insufficiency of Naxos protein accounts for disease manifestation. Importantly, our results suggest that increasing levels of truncated plakoglobin or WT plakoglobin protein would be beneficial for treatment of patients with Naxos disease. One approach to this would be to use antisense technology to specifically block nonsense-mediated decay of mutant plakoglobin mRNA, enabling expression of the truncated protein at increased levels (17).

**Methods**

**Statistics.** Data are presented as mean ± SEM unless indicated otherwise. We used 2-tailed Student’s t test for comparisons among groups. Analysis was performed using Microsoft Excel software. P values of less than 0.05 were considered significant.

**Study approval.** All animal procedures were approved by the UCSD Animal Care and Use Committee. UCSD has an Animal Welfare Assurance (A3033-01) on file with the Office of Laboratory Animal Welfare and is fully accredited by AAALAC International.

See the Supplemental Methods for additional details; see Supplemental Table 1 for details regarding antibodies and Supplemental Table 2 for primers used for qRT-PCR.

**Acknowledgments**

M.J. Stroud was supported by an American Heart Association postdoctoral fellowship (13POST17060120). J. Chen was funded by grants from National Heart, Lung, and Blood Institute and is the American Heart Association Endowed Chair. Microscopy work was performed at the UCSD Neuroscience Microscopy Shared Facility and was supported by the NIH (grant P30 NS047101).

Address correspondence to: Ju Chen, UCSD School of Medicine, 9500 Gilman Drive, La Jolla, California 92093-0613, USA. Phone: 858.822.4276; E-mail: juchen@ucsd.edu.