Biallelic Loss of Function Variants in Myocardial Zonula Adherens Protein Gene (MYZAP) Cause a Severe Recessive Form of Dilated Cardiomyopathy

Juan Pablo Ochoa, MD, PhD; Laura Lalaguna, MSc; Jesús G. Mirelis, MD, PhD; Fernando Dominguez, MD, PhD; Esther Gonzalez-Lopez, MD, PhD; Clara Salas, MD; Gaston Roustan, MD; Kathryn A. McGurk, PhD; Sean L. Zheng, BM, BCh, MA, MRCP; Paul J.R. Barton, PhD; James S. Ware, MRCP, PhD; María Victoria Gómez-Gaviro, PhD; Enrique Lara-Pezzi, PhD; Pablo Garcia-Pavia, MD, PhD

Dilated cardiomyopathy (DCM) is the most frequent cause of heart failure in the young and the leading cause of transplantation. Almost half of the cases have a familial (hereditary) component, but even in familial cases, the diagnostic yield of genetic testing is lower than <40%.[1] *Myocardial zonula adherens protein* (*MYZAP*) gene encodes a protein widely expressed in cardiac tissue, being an emerging candidate to become a DCM-associated gene.

We evaluated a family from Spanish origin (European ancestry) in which 2 sisters were affected by a severe form of DCM with healthy unaffected parents. The study was approved by the Hospital Universitario Puerta de Hierro ethics committee and patients provided informed consent. Data are available from the corresponding author upon reasonable request.

The proband was a 35-year-old female with dyspnea (New York Heart Association II) and palpitations after her first pregnancy. Echocardiography showed mild dilatation of the left ventricle (LV) with systolic dysfunction (LV ejection fraction, 44%) and frequent ventricular ectopic beats (1000/24 h) were detected on Holter ECG. Cardiac magnetic resonance showed biventricular midventricular fibrosis predominantly located in the LV. Familial work-up was performed. Both the 62-year-old father and 60-year-old mother had normal cardiac studies. However, her asymptomatic 32-year-old sister showed LV dilatation with LV

ejection fraction of 35% and a similar pattern of late gadolinium enhancement on cardiac magnetic resonance (Figure [A]). Nonsustained VT episodes were detected, and an implantable cardioverter defibrillator was implanted. During follow-up she received an appropriate shock while sleeping due to sustained VT. None of the affected patients had extracardiac features. Initial genetic analysis of desmosomal genes and a next generation sequencing cardiomyopathy panel (126 genes) were negative.

Exome sequencing was performed in all family members. Assuming a recessive model, we identified *MYZAP* as the only gene with 2 rare candidate loss-of-function (LoF) variants in both siblings. The first was a nonsense variant (NM_001018100.5 c.349C>T) predicting a premature termination codon (p.Arg117*) in exon 4. The second, NM_001018100.5 c.933+1G>A, affected the first nucleotide of the splice donor site of intron 8, predicting the loss of the donor site. Each parent carried 1 variant (Figure [A]), confirming that they were present in compound heterozygosis (different alleles) in the affected daughters.

We hypothesized that only biallelic LoF variants in *MYZAP* would be associated with DCM. We performed immunostaining of skin samples with anti-MYZAP antibodies in the family to evaluate if there was protein expression as it had been described in pemphigus patients.[2] A similar distribution of the MYZAP staining pattern was observed

Key Words: cardiomyopathies ■ dilated cardiomyopathy ■ human genetics ■ tachycardia, ventricular ■ whole exome sequencing

Nonstandard Abbreviations and Acronyms

DCM	dilated cardiomyopathy
GFP	green fluorescent protein
LoF	loss of function
LV	left ventricle
MYZAP	myocardial zonula adherens protein

in all individuals, which was preferentially located in perivascular areas, but with a weaker expression in samples from carriers of biallelic variants (Figure B).

Cardiac muscle cell lines (HL-1) were transfected with GFP (green fluorescent protein)–tagged expression vectors for MYZAP: peGFP-C1-MYZAP-WT, peGFP-C1-MYZAP-p.Arg117*, and peGFP-C1-MYZAP-c.933+1G>A. Western blot analysis showed a significantly lower expression of MYZAP c.933+1G>A compared with
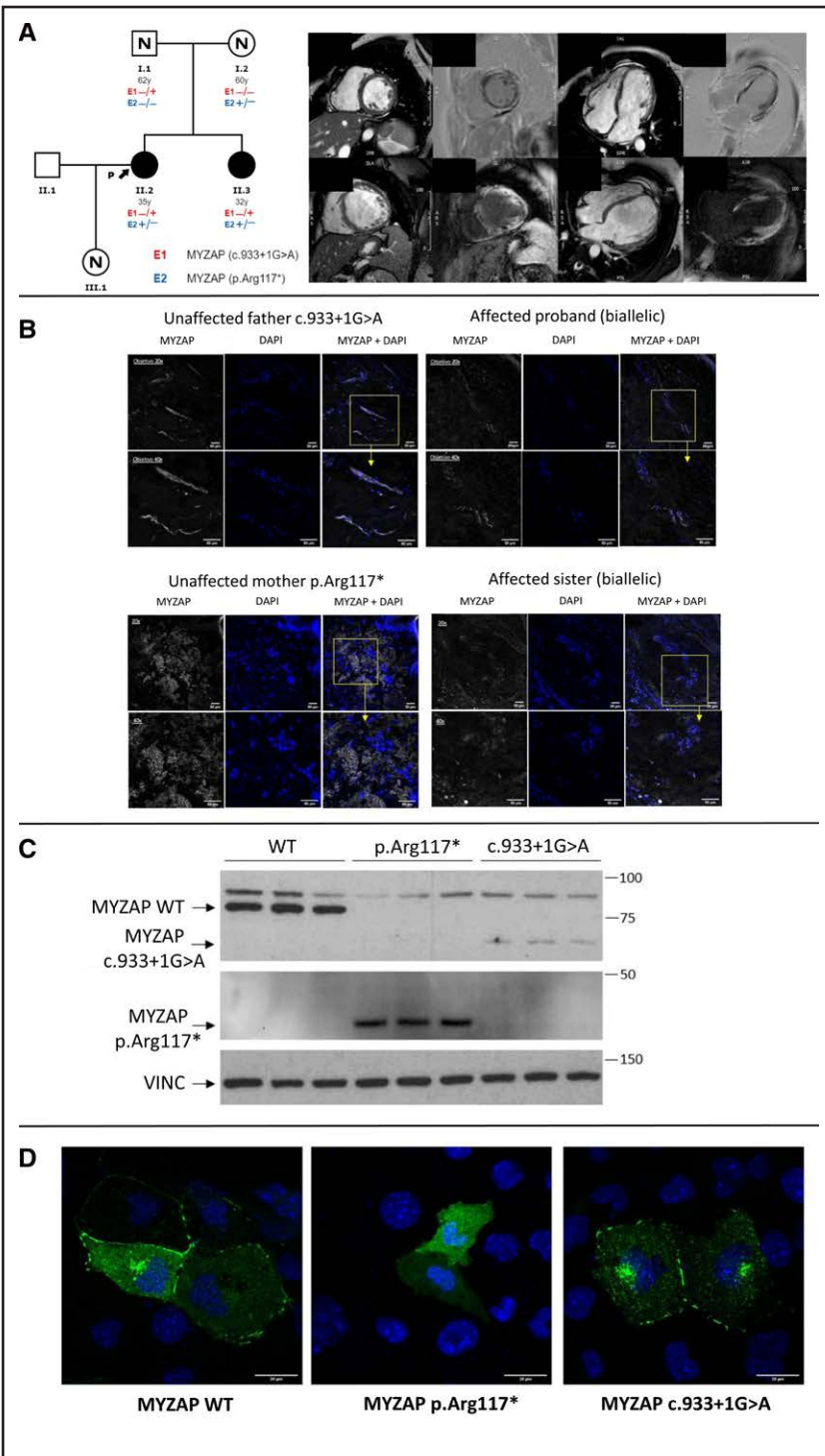

Figure. Clinical and functional evaluations performed in the family.
A, Family pedigree and cardiac magnetic resonance imaging of the index case (**upper**) showing LV dilatation (LVEDV=168 mL) and extensive areas of midventricular fibrosis predominantly in the left ventricle. Her affected sister (**lower**) had similar findings on magnetic resonance imaging (LVEDV=260 mL). **B**, MYZAP immunostaining in skin samples from affected siblings and unaffected parents. Skin biopsies were immunostained with anti-MYZAP antibody (white). Nuclei were stained with DAPI (blue). Images were acquired with a 20× objective. Magnifications of the selected areas (yellow squares) were acquired with a 40× objective. **C**, Western blot analysis of MYZAP in HL-1 cells transfected with MYZAP-WT, MYZAP-p.Arg117*, and MYZAP-c.933+1G>A. MYZAP-c.933+1G>A shows significantly lower expression level of MYZAP compared with MYZAP-WT and MYZAP-p.Arg117*. **D**, Immunofluorescence analysis of localization of MYZAP in transfected HL-1 cells showing that MYZAP-WT and MYZAP-c.933+1G>A are observed predominantly as perinuclear aggregations and in the cytoplasmatic membrane, while MYZAP-p.Arg117* delocalizes being distributed in the cytoplasm but not in any particular structure at cell level. Bar, 20 μm. DAPI indicates 4′,6-diamidino-2-phenylindole; LVEDV, left ventricle end diastolic volume; VINC, vinculin; and WT, wild-type.

MYZAP-WT and MYZAP-p.Arg117* (Figure [C]). Immunofluorescence to evaluate the subcellular protein localization showed that MYZAP-WT and MYZAP-c.933+1G were predominantly localized in the plasma membrane and as perinuclear aggregates, whereas MYZAP-p.Arg117* was distributed randomly throughout the cytoplasm, without reaching intercellular junctions (Figure [D]). These experiments confirm that neither of the 2 alleles are viable: one of them produces a protein that is degraded and the other produces a dysfunctional protein that is delocalized in the cytoplasm. These results, in which no functional protein was produced in affected carriers, support previous findings of functional studies in zebrafish[3] and knock-out mice[4] that develop DCM with decreased contractile function, and increased mortality, being MYZAP localized in the intercalated discs.

Finally, to evaluate the relevance of MYZAP variants in humans, we obtained data from 200 571 participants from the UK Biobank and found 29 distinct rare LoF variants in 98 individuals. There were no biallelic LoF carriers and none of the LoF heterozygous individuals were diagnosed with cardiomyopathy. Furthermore, none of 1446 patients with cardiomyopathies studied by whole genome sequencing in the 100 000 Genomes Project England had biallelic LoF variants in MYZAP. Cardiomyopathy was not more prevalent in carriers of heterozygous MYZAP LoF variants (2/26; 7.7%) compared with individuals without MYZAP variants (1420/15 552; 8.4%; χ^2 test: odds ratio, 0.91 [95% CI, 0.15–3.14]).

In this research, we describe and demonstrate the association between biallelic LoF variants in MYZAP and a severe form of DCM, characterized by profuse biventricular fibrosis and ventricular arrhythmias. This phenotype is coincident with 2 homozygous LoF cases recently reported.[5] Unlike the previous report, affected individuals in our family exhibited LoF in compound heterozygosity, providing further support to MYZAP as a recessive DCM-associated gene and suggesting that the relevance of MYZAP variants in DCM may be higher than previously anticipated. This could be particularly important in populations with high rates of consanguinity. Furthermore, by analyzing large population cohorts we show that although LoF variants in single heterozygosity in MYZAP are observed at low frequency in the general population, they are not associated with cardiac disease in the heterozygous state. Based on our findings, we propose MYZAP as a gene to be considered for evaluation in patients with DCM, particularly among gene-elusive patients with arrhythmogenic DCM whose parents do not show cardiac disease.

ARTICLE INFORMATION

Affiliations

Hospital Universitario Puerta de Hierro Majadahonda, IDIPHISA, Madrid, Spain (J.P.O., J.G.M., F.D., E.G.-L., C.S., G.R., P.G.-P.). CIBERCV, Instituto de Salud Carlos III, Madrid, Spain (J.P.O., F.D., E.G.-L., C.S., E.L.-P., P.G.-P.). Centro Nacional de Investigaciones Cardiovasculares, Madrid, Spain (J.P.O., L.L., F.D., E.G.-L., E.L.-P., P.G.-P.). National Heart and Lung Institute (K.A.M, S.L.Z., P.J.R.B., J.S.W.) and MRC London Institute of Medical Sciences (K.A.M., S.L.Z., P.J.R.B., J.S.W.), Imperial College London, United Kingdom. Royal Brompton and Harefield Hospitals, Guy's and St Thomas' Hospital NHS Trust, London, UK (S.L.Z., P.J.R.B., J.S.W.). Instituto de Investigación Sanitaria Gregorio Marañón, Madrid, Spain (M.V.G.-G.). Departamento de Bioingeniería e Ingeniería Aeroespacial, Universidad Carlos III de Madrid, Leganés, Spain (M.V.G.-G.). Centro de Investigación Biomédica en Red de Salud Mental, Madrid, Spain (M.V.G.-G.). Universidad Francisco de Vitoria, Pozuelo de Alarcón, Spain (P.G.-P.).

Disclosures

Dr Ochoa is an employee of Health in Code. Dr Ware has consulted for MyoKardia Inc, Foresite Labs, and Pfizer. The other authors reports no conflicts.

REFERENCES

1. Escobar-Lopez L, Ochoa JP, Mirelis JG, Espinosa MA, Navarro M, Gallego-Delgado M, Barriales-Villa R, Robles-Mezcua A, Basurte-Elorz MT, Gutiérrez García-Moreno L, et al. Association of genetic variants with outcomes in patients with nonischemic dilated cardiomyopathy. J Am Coll Cardiol. 2021;78:1682–1699. doi: 10.1016/j.jacc.2021.08.039

2. Abreu Velez AM, Yi H, Warfvinge G, Howard MS. Autoantibodies to full body vascular cell junctions colocalize with MYZAP, ARVCF, desmoplakins I and II and p0071 in endemic pemphigus in Colombia, South America. Int J Dermatol. 2018;57:291–298. doi: 10.1111/ijd.13827

3. Frank D, Rangrez AY, Poyanmehr R, Seeger TS, Kuhn C, Eden M, Stiebeling K, Bernt A, Grund C, Franke WW, et al. Mice with cardiac-restricted overexpression of Myozap are sensitized to biomechanical stress and develop a protein-aggregate-associated cardiomyopathy. J Mol Cell Cardiol. 2014;72:196–207. doi: 10.1016/j.yjmcc.2014.03.016

4. Rangrez AY, Eden M, Poyanmehr R, Kuhn C, Stiebeling K, Dierck F, Bernt A, Lüllmann-Rauch R, Weiler H, Kirchof P, et al. Myozap deficiency promotes adverse cardiac remodeling via differential regulation of mitogen-activated protein kinase/serum-response factor and β-catenin/GSK-3β protein signaling. J Biol Chem. 2016;291:4128–4143. doi: 10.1074/jbc.M115.689620

5. Maver A, Zigman T, Rangrez AY, Coric M, Homolak J, Saric D, Skific I, Udovicic M, Zekusic M, Saleem U, et al. A biallelic loss-of-function variant in MYZAP is associated with a recessive form of severe dilated cardiomyopathy. Cold Spring Harb Mol Case Stud. 2022;8:a006221. doi: 10.1101/mcs.a006221