

# MPDU1 mutations underlie a novel human congenital disorder of glycosylation, designated type If

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## Corrigendum

Original citation: *J. Clin. Invest.*108:1687–1695 (2003). doi:10.1172/JCI13419. Citation for this corrigendum: *J. Clin. Invest.*111:925 (2003). doi:10.1172/JCI13419C1. Barbara Schenk,<sup>1</sup> Timo Imbach,<sup>2</sup> Christian G. Frank,<sup>1</sup> Claudia E. Grubenmann,<sup>2</sup> Gerald V. Raymond,<sup>3</sup> Haggit Hurvitz,<sup>4</sup> Isabelle Korn-Lubetzki,<sup>4</sup> Shoshana Revel-Vik,<sup>4</sup> Annick Raas-Rotschild,<sup>5</sup> Anthony S.Luder,<sup>6</sup> Jaak Jaeken,<sup>7</sup> Eric G. Berger,<sup>2</sup> Gert Matthijs,<sup>8</sup> Thierry Hennet,<sup>2</sup> and Markus Aebi<sup>1</sup> <sup>1</sup>Institute of Microbiology, Swiss Federal Institute of Technology, Zurich, Switzerland <sup>2</sup>Institute of Physiology, University of Zurich, Switzerland <sup>3</sup>Kennedy Krieger Institute, Baltimore, Maryland, USA <sup>4</sup>Department of Pediatrics, Bikur Cholim Hospital, Jerusalem, Israel <sup>5</sup>Genetic Clinic, Hadassah University Hospital, Jerusalem, Israel <sup>6</sup>Department of Pediatrics, Sieff Hospital, Safed, Israel, and Faculty of Medicine, Technion, Haifa, Israel <sup>7</sup>Department of Pediatrics, University Hospital, Leuven, Belgium <sup>8</sup>Center for Human Genetics, Catholic University, Leuven, Belgium

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**Idiopathic restrictive cardiomyopathy is part of the clinical expression of cardiac troponin I mutations**

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The authors wish to correct errors that appeared in the Methods section and throughout the paper. The correct sentences are below. The authors regret the errors.

Mutation analysis of *TNNI3* by direct sequencing identified a 87A→G nucleotide substitution of exon 8 resulting in an Asp190Gly amino acid substitution that segregated with the disease in the family (maximal two-point lode score: 4.8).

Direct sequencing of *TNNI3* identified a 93G→A nucleotide substitution of exon 8, which resulted in an Arg192His amino acid substitution.

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During the preparation of this manuscript for publication, errors were introduced into the author list. The corrected author list and affiliations appear below. The authors regret these errors.

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