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

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Corrigendum

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Corrigenda
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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.

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

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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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