Reviewer’s report


Version: 1 Date: 23 Apr 2018

Reviewer: Leticia Manuel Apolinar

This article establishes a case of early diabetes that is distinguished by a number of atypical clinical features, also in this study was identified a rare mutation at a highly conserved site in exon 8 of HNF1β

Mutations in HNF1β are commonly associated with both monogenic diabetes and kidney disease.

The article is acceptable for publication.

Are the methods appropriate and well described?
If not, please specify what is required in your comments to the authors.
Yes

Does the work include the necessary controls?
If not, please specify which controls are required in your comments to the authors.
Yes

Are the conclusions drawn adequately supported by the data shown?
If not, please explain in your comments to the authors.
Yes

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Please indicate the quality of language in the manuscript:

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