Reviewer’s report

Title: A case report of a mild form of multiple acyl-CoA dehydrogenase deficiency due to compound heterozygous mutations in the ETFA gene

Version: 0 Date: 18 Nov 2019

Reviewer: Brian Meyer

Reviewer's report:

This is a comprehensive and well written case report which merits publication based upon the somewhat milder phenotype associated with novel compound heterozygous mutations. The authors clearly explain the differential clinical and genetic diagnosis and present the case in a manner that is both of interest and value to future laboratory or clinical based researchers and physicians. Some minor English language edits are necessary prior to publication.

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

Yes

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Yes

Are the conclusions drawn adequately supported by the data shown?
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Yes

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Not relevant to this manuscript

Quality of written English
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Needs some language corrections before being published
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