Reviewer’s report

Title: Whole-exome sequencing identifies homozygous mutation in TTI2 in a child with primary microcephaly: a case report

Version: 1 Date: 05 Feb 2020

Reviewer: Ni-Chung Lee

Reviewer’s report:

All the points had been addressed, though the author did not understand the request of listing the % of patients having a specific phenotype for table 1, so that reader can understand more about the clinical phenotype. It will be great if author can do so.

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.

Unable to assess

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

Yes

Are you able to assess any statistics in the manuscript or would you recommend an additional statistical review?
If an additional statistical review is recommended, please specify what aspects require further assessment in your comments to the editors.

Not relevant to this manuscript

Quality of written English
Please indicate the quality of language in the manuscript:

Acceptable

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