Reviewer’s report

Title: Intrafamilial phenotypic heterogeneity in a Taiwanese family with MAPT p.R5H mutation: a case report and literature review

Version: 0 Date: 09 Jul 2017

Reviewer: Hiroaki Adachi

Reviewer’s report:

The manuscript by Lin et al examined the pathogenic mutations of the MAPT gene in a patient with frontotemporal degeneration. The authors found a c.14G>A (p.R5H) mutation in a Taiwanese FTD family. However, the symptoms and signs are different among the patients in the family. And his siblings do not showed any neurological symptoms. Collectively, this paper includes no novel finding so far.

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?
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No

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I am able to assess the statistics

Quality of written English
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Acceptable

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