Reviewer’s report

Title: Dissecting TSC2-mutated renal and hepatic angiomyolipomas in an individual with ARID1B-associated intellectual disability

Version: 1 Date: 11 Jan 2019

Reviewer: Shuli Liang

Reviewer's report:

This manuscript was well written, and reported a patient with NDD and renal and hepatic AMLs. Exome and panel sequencing fail to find difference in the ARID1B variant in blood and AMLs, however the TSC2 gene and SIPA1L3 gene mutations were found in AMLs but not blood. The rare clinical case and post-zygotic variant were adequately analyzed. I think the manuscript can be accepted now.

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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