Reviewer's report

Title: An atypical form of AOA2 with myoclonus associated with mutations in SETX and AFG3L2

Version: 2 Date: 14 December 2014

Reviewer: Filippo M. Santorelli

Reviewer's report:

This work is important in the field and outlines general points in the era of NGS use in clinical genetics

Level of interest: An article of importance in its field

Quality of written English: Acceptable

Statistical review: No, the manuscript does not need to be seen by a statistician.

Declaration of competing interests:

I declare that I have no competing interests