Reviewer's report

Title: Whole-exome sequencing identifies MYO15A mutations as a cause of autosomal recessive nonsyndromic hearing loss in Korean families

Version: 2 Date: 28 June 2013

Reviewer: Mustafa Tekin

Reviewer's report:

Revised manuscript has improved and is suitable for publication

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