Reviewer's report

Title: Lack of significant association between mutations of KCNJ10 or FOXI1 and SLC26A4 mutations in Pendred syndrome/Enlarged Vestibular Aqueducts

Version: 2 Date: 24 July 2013

Reviewer: Pu Dai

Reviewer's report:

Discretionary Revisions
1. Results from hearing loss family members will contribute to the judgement of the pathogenity of variants.

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.