Reviewer's report

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

Version: 1 Date: 23 May 2013

Reviewer: Fiona FM Macdonald

Reviewer's report:

This paper provides further information on the role, or in this case lack of a role, of two genes, KCNJ10 and FOXI1, in Pendred's syndrome in a large series of patients. It is clearly written, methods are well described and results sound but concise given the outcomes.

Level of interest: An article whose findings are important to those with closely related research interests

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