Reviewer's report

Title: Inherited KIF21A and PAX6 gene Mutations in A Boy with Congenital Fibrosis of Extraocular Muscles and Aniridia

Version: 3 Date: 20 March 2013

Reviewer: Suma Shankar

Reviewer's report:

Major Compulsory Revisions

Conclusion statement regarding the importance of identifying PAX6 mutation.

Individuals with PAX6 mutations are not at risk for Wilm's tumor and hence will not require routine screening for Wilm's tumor as needed in WAGR syndrome or where the molecular diagnosis for aniridia is not known.

Level of interest: An article of limited interest

Quality of written English: Needs some language corrections before being published

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'