Reviewer's report

Title: Absence of mutations in NR2E1 and SNX3 in five patients with MMEP (microcephaly, microphthalmia, ectrodactyly, and prognathism) and related phenotypes

Version: 2 Date: 29 May 2007
Reviewer: Diane W Cox

Reviewer's report:

General
All concerns have been answered.

Major Compulsory Revisions (that the author must respond to before a decision on publication can be reached)
no further changes required

Minor Essential Revisions (such as missing labels on figures, or the wrong use of a term, which the author can be trusted to correct)

Discretionary Revisions (which the author can choose to ignore)

What next?: Accept without revision

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