Reviewer's report

Title: Whole-exome sequencing of a rare case of familial childhood acute lymphoblastic leukemia reveals putative predisposing mutations in Fanconi anemia genes.

Version: 4 Date: 14 June 2015

Reviewer: Maria Castella

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The changes introduced by the authors improved significantly the quality of the manuscript. The analysis of the exome data is now presented in a more complete and meaningful way. Although the findings presented can potentially be of interest to people working in the same field, the work remains speculative.

Level of interest: An article of limited interest

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