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.

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Reviewer: Tugba Tumer

Reviewer's report:

The manuscript by Spinella et al., is clearly structured, well focused and gives a comprehensive overview over the topic. However, since they reported the case of a single nonsyndromic pre-B childhood ALL family with two non-twinned siblings diagnosed with ALL, I suggest to publish this article as a case report rather than a research article.

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.