Reviewer's report

Title: A Novel Frame-shift deletion in FANCF Gene Causing Autosomal Recessive Fanconi Anemia: a case report

Version: 0 Date: 17 Jan 2019

Reviewer: Babu Rao Vundinti

Reviewer's report:

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1. The manuscript entitled "A Novel Frame-shift deletion in FANCF Gene Causing Autosomal Recessive Fanconi Anemia: a case report submitted by Zareifar and colleagues. The manuscript is poorly written.
2. It is a single case report with novel frameshift mutation. The case report is lacking genotype-phenotype correlation.
3. The chromosomal breakage frequency need be given properly.
4. Discussion is lacking the mutations reported in FANCF gene and its clinical correlation.
5. Authors are not discussed about the clinical severity of the novel mutations compared to literature reports.
6. The parental status of genetic mutations not given in the manuscript.
7. The management importance genetic mutations need to be clearly written.

Are the methods appropriate and well described?
If not, please specify what is required in your comments to the authors.

No

Does the work include the necessary controls?
If not, please specify which controls are required in your comments to the authors.

No

Are the conclusions drawn adequately supported by the data shown?
If not, please explain in your comments to the authors.
Are you able to assess any statistics in the manuscript or would you recommend an additional statistical review?
If an additional statistical review is recommended, please specify what aspects require further assessment in your comments to the editors.

Not relevant to this manuscript

Quality of written English
Please indicate the quality of language in the manuscript:

Not suitable for publication unless extensively edited

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