Reviewer’s report

Title: Inherited FGFR2 mutation in a Chinese patient with Crouzon syndrome and luxation of bulbus oculi provoked by trauma: A case report

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Reviewer: N Al-Namnam

Reviewer's report:

In the manuscript 'BOPH-D-19-00517' entitled "Luxation of the bulbus oculi in Crouzon syndrome: a case report' by ji yang; tao tao; hai liu; zhulin hu presented luxation of bulbus oculi which is not new symptoms to the Crouzon syndrome cases.

the authors presented the genetic diagnosis of crouzon syndrome in general rather than discuss the any new gene mutation that result of this phenotype that carry this symptoms (Luxation of the bulbus oculi) with other phenotype of Crouzon syndrome who does not has this symptoms. This will add a significant value to this field.

Background section, line 20: Crouzon syndrome (CS) is one of the causes of fibroblast growth factor receptor 2 mutations related to craniosynstosis, It should be" results of" not "causes of".

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

Yes

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

Yes
Are the conclusions drawn adequately supported by the data shown?
If not, please explain in your comments to the authors.

Yes

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:

Acceptable

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