Reviewer's report

Title: A unique phenotype in a patient with a rare triplication of the 22q11.2 region, and new clinical insights of the 22q11.2 deletion and microduplication syndromes: a report of four cases

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Reviewer: Shi-Hui Yu

Reviewer's report:

Major concerns:
Both 22.11.2 deletion and duplications have been systemically investigated previously. The three cases (patients 1-3) in this report present nothing new, both genetically and phenotypically.
The 22q11.2 triplication in Patient 4 is interesting. An aCGH image showing the triplicated region is necessary.
Necessary description of material and method used in this report is required.

Minor concerns:
There are numbers of grammertical errors in this manuscript.

Level of interest: An article of limited interest

Quality of written English: Needs some language corrections before being published

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