Reviewer’s report

Title: Type II diabetes and impaired glucose tolerance due to severe hyperinsulinism in patients with 1p36 deletion syndrome and Prader-Willi like phenotype

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Reviewer: Fiorella Gurrieri

Reviewer’s report:

The report is of interest because the knowledge of increased diabetes and obesity susceptibility in those patients is important for clinical management.

Minor essential revisions:
the manuscript needs robust editing and, possibly, should be shortened as it is too long for a case report.
A methylation test for Prader Willy syndrome may be useful to exclude additional genetic causes for this phenotype.
In the methods (page 5 and 7) it seems that array-CGH was done with two different approaches, but it is probably not the case, so please check.

Level of interest: An article whose findings are important to those with closely related research interests

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'