Reviewer#1:

Dear Authors, I would like to thank you for submitting the case report entitled " A Rare Mutation in MKRN3 in Two Twin Sisters with Central Precocious Puberty: a Case Report". It describes twin sisters with central presenting with precocious puberty. This presentation was related to a specific genetic mutation that disrupts the hypothalamic-pituitary-gonadal axis. The mutated gene is located on chromosome 15q11.2. which codes the Makorin protein family. Case presentation, diagnostic work-up, genetic mutation, and treatment were described concisely and to the point. As a reviewer, I enjoyed reading the manuscript and find it acceptable for publishing. Many thanks again. Kind Regards, Sanem Guler Cimen

Reply: Thank you for your comments. I have revised the manuscript according to your request. Due to the limitation of the submission system of this journal, some subheadings cannot be corrected according to your instructions when generating the final document. The controversial wording has been corrected or deleted. Thank you again for your review, thank you.

Reviewer#2:

The finding is worth reporting. The case is well written except for the minor corrections highlighted in the review especially in the presentation of the history. overall, it is a good report.

Reply: Thank you for your comments.