Reviewer's report

Title: A rare missense variant in APC interrupts splicing and causes AFAP in two Danish families

Version: 0 Date: 17 Jan 2020

Reviewer: Andrzej Plawski

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Very interesting material but it may be coincidence of the both diseases in my opinion authors did not proof the role of the mutations of APC in pathogenesis of Caroli Disease In my opinion most interesting is formation new cryptic acceptor splice site. c.289 is located in the middle of exon and it is very important information for all studding the genetic bases of hereditary disease.

Level of interest
Please indicate how interesting you found the manuscript:

An exceptional article

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

Acceptable

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