Reviewer’s report

Title: Implementation of massive sequencing in the genetic diagnosis of hereditary cancer syndromes: diagnostic performance in the Hereditary Cancer Programme of the Valencia Community (FamCan-NGS)

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Reviewer: Magdalena Ratajska

Reviewer's report:

This is a well written and interesting analysis pointing out the challenges of molecular diagnostic in HCS patients.

Authors describe the advantages of using NGS over standard testing approach, and at the same time discuss potential difficulties while introducing high-throughput technologies to routine diagnostic laboratories.

I highly recommend this paper for publication.

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An article of importance in its field

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

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