A nonâ€sense <scp>MCM9</scp> mutation in a f insufficiency

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Citation Report

#	Article	IF	Citations
1	Premature Ovarian Insufficiency: New Perspectives on Genetic Cause and Phenotypic Spectrum. Endocrine Reviews, 2016, 37, 609-635.	20.1	170
2	Pathogenic germline MCM9 variants are rare in Australian Lynch-like syndrome patients. Cancer Genetics, 2016, 209, 497-500.	0.4	8
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16	Rare variants in FANCA induce premature ovarian insufficiency. Human Genetics, 2019, 138, 1227-1236.	3.8	56
17	Contribution to colonic polyposis of recently proposed predisposing genes and assessment of the prevalence of NTHL1 â€•and MSH3 â€associated polyposes. Human Mutation, 2019, 40, 1910-1923.	2.5	24
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20	Association study of the three functional polymorphisms (TAS2R46G>A, OR4C16G>A, and) Tj ETQq1 1 0.784314 rgBT /Overloc	1.4	21
21	Gene variants identified by whole-exome sequencing in 33 French women with premature ovarian insufficiency. Journal of Assisted Reproduction and Genetics, 2019, 36, 39-45.	2.5	35
22	Genetics and Genomics of Primary Ovarian Insufficiency. , 2019, , 427-445.		3
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