## Mckinsey L Goodenberger

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2184230/publications.pdf

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933447 1199594 13 1,188 10 12 citations h-index g-index papers 13 13 13 2400 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Genetics of adult glioma. Cancer Genetics, 2012, 205, 613-621.	0.4	737
2	Rare Deletions at the Neurexin 3 Locus in Autism Spectrum Disorder. American Journal of Human Genetics, 2012, 90, 133-141.	6.2	182
3	Lynch Syndrome and MYH-Associated Polyposis. Journal of Clinical Gastroenterology, 2011, 45, 488-500.	2.2	62
4	PMS2 monoallelic mutation carriers: the known unknown. Genetics in Medicine, 2016, 18, 13-19.	2.4	51
5	Deletions in 16q24.2 are associated with autism spectrum disorder, intellectual disability and congenital renal malformation. Journal of Medical Genetics, 2013, 50, 163-173.	3.2	36
6	Genetic testing utilization and the role of the laboratory genetic counselor. Clinica Chimica Acta, 2014, 427, 193-195.	1.1	32
7	Further Defining the Role of the Laboratory Genetic Counselor. Journal of Genetic Counseling, 2016, 25, 786-798.	1.6	29
8	Copy number variant discrepancy resolution using the ClinGen dosage sensitivity map results in updated clinical interpretations in ClinVar. Human Mutation, 2018, 39, 1650-1659.	2.5	23
9	The Utilization of Counseling Skills by the Laboratory Genetic Counselor. Journal of Genetic Counseling, 2015, 24, 6-17.	1.6	15
10	Pathologic findings in breast, fallopian tube, and ovary specimens in non-BRCA hereditary breast and/or ovarian cancer syndromes: a study of 18 patients with deleterious germline mutations in RAD51C, BARD1, BRIP1, PALB2, MUTYH, or CHEK2. Human Pathology, 2017, 70, 14-26.	2.0	11
11	Stankiewicz-Isidor syndrome: expanding the clinical and molecular phenotype. Genetics in Medicine, 2022, 24, 179-191.	2.4	9
12	Response to Commercial Genetic Testing and the Future of the Genetic Counseling Profession. Journal of Genetic Counseling, 2018, 27, 530-532.	1.6	1
13	Response to ten Broeke and Nielsen. Genetics in Medicine, 2015, 17, 684-685.	2.4	0