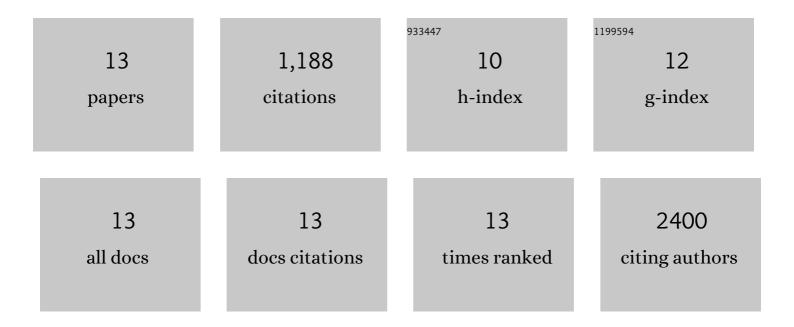
Mckinsey L Goodenberger

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Stankiewicz-Isidor syndrome: expanding the clinical and molecular phenotype. Genetics in Medicine, 2022, 24, 179-191.	2.4	9
2	Response to Commercial Genetic Testing and the Future of the Genetic Counseling Profession. Journal of Genetic Counseling, 2018, 27, 530-532.	1.6	1
3	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
4	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
5	Further Defining the Role of the Laboratory Genetic Counselor. Journal of Genetic Counseling, 2016, 25, 786-798.	1.6	29
6	PMS2 monoallelic mutation carriers: the known unknown. Genetics in Medicine, 2016, 18, 13-19.	2.4	51
7	The Utilization of Counseling Skills by the Laboratory Genetic Counselor. Journal of Genetic Counseling, 2015, 24, 6-17.	1.6	15
8	Response to ten Broeke and Nielsen. Genetics in Medicine, 2015, 17, 684-685.	2.4	0
9	Genetic testing utilization and the role of the laboratory genetic counselor. Clinica Chimica Acta, 2014, 427, 193-195.	1.1	32
10	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
11	Genetics of adult glioma. Cancer Genetics, 2012, 205, 613-621.	0.4	737
12	Rare Deletions at the Neurexin 3 Locus in Autism Spectrum Disorder. American Journal of Human Genetics, 2012, 90, 133-141.	6.2	182
13	Lynch Syndrome and MYH-Associated Polyposis. Journal of Clinical Gastroenterology, 2011, 45, 488-500.	2.2	62