

Kelsey E Grinde

List of Publications by Year in descending order

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Version: 2024-02-01

12
papers

281
citations

1307594

7
h-index

1281871

11
g-index

12
all docs

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docs citations

12
times ranked

1110
citing authors

#	ARTICLE	IF	CITATIONS
1	Circular RNA Profiles in Viremia and ART Suppression Predict Competing circRNAâ€“miRNAâ€“mRNA Networks Exclusive to HIV-1 Viremic Patients. <i>Viruses</i> , 2022, 14, 683.	3.3	3
2	Whole genome sequence analyses of eGFR in 23,732 people representing multiple ancestries in the NHLBI trans-omics for precision medicine (TOPMed) consortium. <i>EBioMedicine</i> , 2021, 63, 103157.	6.1	14
3	Coagulation factor VIII: Relationship to cardiovascular disease risk and whole genome sequence and epigenomeâ€“wide analysis in African Americans. <i>Journal of Thrombosis and Haemostasis</i> , 2020, 18, 1335-1347.	3.8	17
4	A fully adjusted twoâ€“stage procedure for rankâ€“normalization in genetic association studies. <i>Genetic Epidemiology</i> , 2019, 43, 263-275.	1.3	60
5	Genome-wide Significance Thresholds for Admixture Mapping Studies. <i>American Journal of Human Genetics</i> , 2019, 104, 454-465.	6.2	25
6	Generalizing polygenic risk scores from Europeans to Hispanics/Latinos. <i>Genetic Epidemiology</i> , 2019, 43, 50-62.	1.3	89
7	Mo1146 ANESTHESIA PROFESSIONAL-DELIVERED SEDATION IS ASSOCIATED WITH SIMILAR OUTCOMES COMPARED TO NURSE ADMINISTERED SEDATION IN PATIENTS WITH ACUTE UPPER GASTROINTESTINAL BLEEDING. <i>Gastrointestinal Endoscopy</i> , 2018, 87, AB418-AB419.	1.0	0
8	Illustrating, Quantifying, and Correcting for Bias in Post-hoc Analysis of Gene-Based Rare Variant Tests of Association. <i>Frontiers in Genetics</i> , 2017, 8, 117.	2.3	4
9	Local Ancestry Inference in a Large US-Based Hispanic/Latino Study: Hispanic Community Health Study/Study of Latinos (HCHS/SOL). <i>G3: Genes, Genomes, Genetics</i> , 2016, 6, 1525-1534.	1.8	51
10	A general method for combining different family-based rare-variant tests of association to improve power and robustness of a wide range of genetic architectures. <i>BMC Proceedings</i> , 2016, 10, 165-170.	1.6	4
11	A multistep approach to single nucleotide polymorphismâ€“set analysis: an evaluation of power and type I error of gene-based tests of association after pathway-based association tests. <i>BMC Proceedings</i> , 2016, 10, 349-355.	1.6	2
12	A general approach for combining diverse rare variant association tests provides improved robustness across a wider range of genetic architectures. <i>European Journal of Human Genetics</i> , 2016, 24, 767-773.	2.8	12