

# Zeynep Baskurt

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2737055/publications.pdf>

Version: 2024-02-01

10  
papers

360  
citations

1307594

7  
h-index

1372567

10  
g-index

11  
all docs

11  
docs citations

11  
times ranked

734  
citing authors

#	ARTICLE	IF	CITATIONS
1	VikNGS: a C++ variant integration kit for next generation sequencing association analysis. <i>Bioinformatics</i> , 2020, 36, 1283-1285.	4.1	10
2	Genetic association and transcriptome integration identify contributing genes and tissues at cystic fibrosis modifier loci. <i>PLoS Genetics</i> , 2019, 15, e1008007.	3.5	56
3	Improving imputation in disease-relevant regions: lessons from cystic fibrosis. <i>Npj Genomic Medicine</i> , 2018, 3, 8.	3.8	9
4	Genetic association analysis with pedigrees: Direct inference using the composite likelihood ratio. <i>Genetic Epidemiology</i> , 2018, 42, 826-837.	1.3	2
5	Statistical Reasoning: Choosing and Checking the Ingredients, Inferences Based on a Measure of Statistical Evidence with Some Applications. <i>Entropy</i> , 2018, 20, 289.	2.2	14
6	Goodness of fit for the logistic regression model using relative belief. <i>Journal of Statistical Distributions and Applications</i> , 2017, 4, 17.	1.2	15
7	The Genetics of Reading Disability in an Often Excluded Sample: Novel Loci Suggested for Reading Disability in Rolandic Epilepsy. <i>PLoS ONE</i> , 2012, 7, e40696.	2.5	16
8	Using Parametric Multipoint Lods and Mods for Linkage Analysis Requires a Shift in Statistical Thinking. <i>Human Heredity</i> , 2011, 72, 264-275.	0.8	4
9	Centrotemporal sharp wave EEG trait in rolandic epilepsy maps to Elongator Protein Complex 4 (ELP4). <i>European Journal of Human Genetics</i> , 2009, 17, 1171-1181.	2.8	176
10	Evidence of shared genetic risk factors for migraine and rolandic epilepsy. <i>Epilepsia</i> , 2009, 50, 2428-2433.	5.1	58