

# Doug Speed

## List of Publications by Year in descending order

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Version: 2024-02-01

37  
papers

8,458  
citations

361413  
20  
h-index

345221  
36  
g-index

53  
all docs

53  
docs citations

53  
times ranked

18279  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genome-wide association, prediction and heritability in bacteria with application to <i>Streptococcus pneumoniae</i> . <i>NAR Genomics and Bioinformatics</i> , 2022, 4, lqac011.	3.2	5
2	The Effects of Common Genetic Variation in 96 Genes Involved in Thyroid Hormone Regulation on TSH and FT4 Concentrations. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e2276-e2283.	3.6	6
3	SNP-based heritability and selection analyses: Improved models and new results. <i>BioEssays</i> , 2022, 44, e2100170.	2.5	14
4	P117. Predicting Genetic Risk for Depression and Anxiety Disorders. <i>Biological Psychiatry</i> , 2022, 91, S134-S135.	1.3	0
5	Graph pangenome captures missing heritability and empowers tomato breeding. <i>Nature</i> , 2022, 606, 527-534.	27.8	131
6	Bipolar disorder and cannabis use: A bidirectional two-sample Mendelian randomization study. <i>Addiction Biology</i> , 2021, 26, e13030.	2.6	9
7	Shared genetic basis between genetic generalized epilepsy and background electroencephalographic oscillations. <i>Epilepsia</i> , 2021, 62, 1518-1527.	5.1	5
8	Improved genetic prediction of complex traits from individual-level data or summary statistics. <i>Nature Communications</i> , 2021, 12, 4192.	12.8	76
9	Signatures of TSPAN8 variants associated with human metabolic regulation and diseases. <i>IScience</i> , 2021, 24, 102893.	4.1	5
10	Using common genetic variants to find drugs for common epilepsies. <i>Brain Communications</i> , 2021, 3, fcab287.	3.3	9
11	Evaluating and improving heritability models using summary statistics. <i>Nature Genetics</i> , 2020, 52, 458-462.	21.4	128
12	Genome-wide association study of the sensitivity to environmental stress and adversity neuroticism cluster. <i>Acta Psychiatrica Scandinavica</i> , 2020, 141, 476-478.	4.5	16
13	Estimating narrow-sense heritability using family data from admixed populations. <i>Heredity</i> , 2020, 124, 751-762.	2.6	6
14	Investigating the association between body fat and depression via Mendelian randomization. <i>Translational Psychiatry</i> , 2019, 9, 184.	4.8	90
15	Summary statistic analyses can mistake confounding bias for heritability. <i>Genetic Epidemiology</i> , 2019, 43, 930-940.	1.3	8
16	GWAS on longitudinal growth traits reveals different genetic factors influencing infant, child, and adult BMI. <i>Science Advances</i> , 2019, 5, eaaw3095.	10.3	86
17	Investigating the causal relationship between neuroticism and depression via Mendelian randomization. <i>Acta Psychiatrica Scandinavica</i> , 2019, 139, 395-397.	4.5	20
18	SumHer better estimates the SNP heritability of complex traits from summary statistics. <i>Nature Genetics</i> , 2019, 51, 277-284.	21.4	181

#	ARTICLE	IF	CITATIONS
19	Genetic correlations among psychiatric and immune-related phenotypes based on genome-wide association data. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2018, 177, 641-657.	1.7	158
20	Mendelian randomization study shows no causal relationship between circulating urate levels and Parkinson's disease. <i>Annals of Neurology</i> , 2018, 84, 191-199.	5.3	43
21	Transient structural variations have strong effects on quantitative traits and reproductive isolation in fission yeast. <i>Nature Communications</i> , 2017, 8, 14061.	12.8	472
22	Reevaluation of SNP heritability in complex human traits. <i>Nature Genetics</i> , 2017, 49, 986-992.	21.4	427
23	Genetic Complexity of Crohn's Disease in Two Large Ashkenazi Jewish Families. <i>Gastroenterology</i> , 2016, 151, 698-709.	1.3	54
24	Systems genetics identifies a convergent gene network for cognition and neurodevelopmental disease. <i>Nature Neuroscience</i> , 2016, 19, 223-232.	14.8	131
25	The genomic and phenotypic diversity of <i>Schizosaccharomyces pombe</i> . <i>Nature Genetics</i> , 2015, 47, 235-241.	21.4	174
26	Systems genetics identifies Sestrin 3 as a regulator of a proconvulsant gene network in human epileptic hippocampus. <i>Nature Communications</i> , 2015, 6, 6031.	12.8	158
27	Relatedness in the post-genomic era: is it still useful?. <i>Nature Reviews Genetics</i> , 2015, 16, 33-44.	16.3	228
28	Genetic Interactions with Sex Make a Relatively Small Contribution to the Heritability of Complex Traits in Mice. <i>PLoS ONE</i> , 2014, 9, e96450.	2.5	9
29	MultiBLUP: improved SNP-based prediction for complex traits. <i>Genome Research</i> , 2014, 24, 1550-1557.	5.5	258
30	A genome-wide association study and biological pathway analysis of epilepsy prognosis in a prospective cohort of newly treated epilepsy. <i>Human Molecular Genetics</i> , 2014, 23, 247-258.	2.9	33
31	Describing the genetic architecture of epilepsy through heritability analysis. <i>Brain</i> , 2014, 137, 2680-2689.	7.6	87
32	Long-stay patients with cancer on the intensive care unit: characteristics, risk factors, and clinical outcomes. <i>British Journal of Anaesthesia</i> , 2013, 111, 1026-1027.	3.4	3
33	Response to Lee et al.: SNP-Based Heritability Analysis with Dense Data. <i>American Journal of Human Genetics</i> , 2013, 93, 1155-1157.	6.2	17
34	Understanding complex traits: from farmers to pharma. <i>Genome Medicine</i> , 2012, 4, 59.	8.2	1
35	Improved Heritability Estimation from Genome-wide SNPs. <i>American Journal of Human Genetics</i> , 2012, 91, 1011-1021.	6.2	656
36	The genomic and transcriptomic architecture of 2,000 breast tumours reveals novel subgroups. <i>Nature</i> , 2012, 486, 346-352.	27.8	4,708

#	ARTICLE	IF	CITATIONS
37	Sparse Partitioning: Nonlinear regression with binary or tertiary predictors, with application to association studies. <i>Annals of Applied Statistics</i> , 2011, 5, .	1.1	3