## Jacob Gratten

List of Publications by Year in descending order

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LACOR CRATTEN

#	Article	IF	CITATIONS
1	Biological insights from 108 schizophrenia-associated genetic loci. Nature, 2014, 511, 421-427.	13.7	6,934
2	Identification of novel risk loci, causal insights, and heritable risk for Parkinson's disease: a meta-analysis of genome-wide association studies. Lancet Neurology, The, 2019, 18, 1091-1102.	4.9	1,414
3	Genome-wide association study identifies 74 loci associated with educational attainment. Nature, 2016, 533, 539-542.	13.7	1,204
4	Modeling Linkage Disequilibrium Increases Accuracy of Polygenic Risk Scores. American Journal of Human Genetics, 2015, 97, 576-592.	2.6	1,098
5	Genetic variants associated with subjective well-being, depressive symptoms, and neuroticism identified through genome-wide analyses. Nature Genetics, 2016, 48, 624-633.	9.4	870
6	Contribution of copy number variants to schizophrenia from a genome-wide study of 41,321 subjects. Nature Genetics, 2017, 49, 27-35.	9.4	838
7	Genomic Dissection of Bipolar Disorder and Schizophrenia, Including 28 Subphenotypes. Cell, 2018, 173, 1705-1715.e16.	13.5	623
8	Genome-wide association analyses of risk tolerance and risky behaviors in over 1 million individuals identify hundreds of loci and shared genetic influences. Nature Genetics, 2019, 51, 245-257.	9.4	536
9	Large-scale genomics unveils the genetic architecture of psychiatric disorders. Nature Neuroscience, 2014, 17, 782-790.	7.1	321
10	Life history trade-offs at a single locus maintain sexually selected genetic variation. Nature, 2013, 502, 93-95.	13.7	296
11	Parkinson's disease age at onset genomeâ€wide association study: Defining heritability, genetic loci, and αâ€synuclein mechanisms. Movement Disorders, 2019, 34, 866-875.	2.2	258
12	Polygenic prediction of educational attainment within and between families from genome-wide association analyses in 3 million individuals. Nature Genetics, 2022, 54, 437-449.	9.4	215
13	Improved precision of epigenetic clock estimates across tissues and its implication for biological ageing. Genome Medicine, 2019, 11, 54.	3.6	191
14	Gene mapping in the wild with SNPs: guidelines and future directions. Genetica, 2009, 136, 97-107.	0.5	181
15	Autism-related dietary preferences mediate autism-gut microbiome associations. Cell, 2021, 184, 5916-5931.e17.	13.5	172
16	Genetic pleiotropy in complex traits and diseases: implications for genomic medicine. Genome Medicine, 2016, 8, 78.	3.6	135
17	Estimation of Genetic Correlation via Linkage Disequilibrium Score Regression and Genomic Restricted Maximum Likelihood. American Journal of Human Genetics, 2018, 102, 1185-1194.	2.6	119
18	Compelling evidence that a single nucleotide substitution in TYRP1 is responsible for coat-colour polymorphism in a free-living population of Soay sheep. Proceedings of the Royal Society B: Biological Sciences, 2007, 274, 619-626.	1.2	116

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19	Genetic variants linked to education predict longevity. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 13366-13371.	3.3	110
20	Risk of psychiatric illness from advanced paternal age is not predominantly from de novo mutations. Nature Genetics, 2016, 48, 718-724.	9.4	98
21	A Localized Negative Genetic Correlation Constrains Microevolution of Coat Color in Wild Sheep. Science, 2008, 319, 318-320.	6.0	97
22	Imprint of assortative mating on the human genome. Nature Human Behaviour, 2018, 2, 948-954.	6.2	97
23	Cross-ethnic meta-analysis identifies association of the GPX3-TNIP1 locus with amyotrophic lateral sclerosis. Nature Communications, 2017, 8, 611.	5.8	93
24	Predicting gene targets from integrative analyses of summary data from GWAS and eQTL studies for 28 human complex traits. Genome Medicine, 2016, 8, 84.	3.6	91
25	Quantitative trait loci (QTL) mapping of resistance to strongyles and coccidia in the free-living Soay sheep (Ovis aries). International Journal for Parasitology, 2007, 37, 121-129.	1.3	87
26	Genome mapping in intensively studied wild vertebrate populations. Trends in Genetics, 2010, 26, 275-284.	2.9	85
27	Analysis of DNA methylation associates the cystine–glutamate antiporter SLC7A11 with risk of Parkinson's disease. Nature Communications, 2020, 11, 1238.	5.8	85
28	Interpreting the role of de novo protein-coding mutations in neuropsychiatric disease. Nature Genetics, 2013, 45, 234-238.	9.4	76
29	The emerging spectrum of allelic variation in schizophrenia: current evidence and strategies for the identification and functional characterization of common and rare variants. Molecular Psychiatry, 2013, 18, 38-52.	4.1	75
30	Heterogeneity of genetic architecture of body size traits in a freeâ€living population. Molecular Ecology, 2015, 24, 1810-1830.	2.0	72
31	Characterisation of the transcriptome of a wild great tit Parus major population by next generation sequencing. BMC Genomics, 2011, 12, 283.	1.2	67
32	Multiplex SNP‧CALE: a costâ€effective mediumâ€ŧhroughput single nucleotide polymorphism genotyping method. Molecular Ecology Resources, 2008, 8, 1230-1238.	2.2	65
33	Development of a Linkage Map and Mapping of Phenotypic Polymorphisms in a Free-Living Population of Soay Sheep (Ovis aries). Genetics, 2006, 173, 1521-1537.	1.2	57
34	Identification of purebredCrocodylus siamensis for reintroduction in Vietnam. The Journal of Experimental Zoology, 2002, 294, 373-381.	1.4	54
35	Introgression and the fate of domesticated genes in a wild mammal population. Molecular Ecology, 2013, 22, 4210-4221.	2.0	53
36	Evidence for Genetic Overlap Between Schizophrenia and Age at First Birth in Women. JAMA Psychiatry, 2016, 73, 497.	6.0	51

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37	Association of Schizophrenia Risk With Disordered Niacin Metabolism in an Indian Genome-wide Association Study. JAMA Psychiatry, 2019, 76, 1026.	6.0	51
38	Meta-analysis of genome-wide DNA methylation identifies shared associations across neurodegenerative disorders. Genome Biology, 2021, 22, 90.	3.8	49
39	MAPPING QUANTITATIVE TRAIT LOCI UNDERLYING FITNESS-RELATED TRAITS IN A FREE-LIVING SHEEP POPULATION. Evolution; International Journal of Organic Evolution, 2007, 61, 1403-1416.	1.1	48
40	Investigating the shared genetic architecture between multiple sclerosis and inflammatory bowel diseases. Nature Communications, 2021, 12, 5641.	5.8	46
41	The genetic basis of recessive self-colour pattern in a wild sheep population. Heredity, 2010, 104, 206-214.	1.2	43
42	Selection and microevolution of coat pattern are cryptic in a wild population of sheep. Molecular Ecology, 2012, 21, 2977-2990.	2.0	31
43	Significant out-of-sample classification from methylation profile scoring for amyotrophic lateral sclerosis. Npj Genomic Medicine, 2020, 5, 10.	1.7	25
44	Whole-exome sequencing in amyotrophic lateral sclerosis suggests NEK1 is a risk gene in Chinese. Genome Medicine, 2017, 9, 97.	3.6	23
45	Study protocol for the Australian autism biobank: an international resource to advance autism discovery research. BMC Pediatrics, 2018, 18, 284.	0.7	20
46	The genetic relationship between female reproductive traits and six psychiatric disorders. Scientific Reports, 2019, 9, 12041.	1.6	18
47	Molecular assessment of the genetic integrity, distinctiveness and phylogeographic context of the Saltwater crocodile (Crocodylus porosus) on Palau. Conservation Genetics, 2007, 8, 777-787.	0.8	17
48	Small non-coding RNA expression from anterior cingulate cortex in schizophrenia shows sex specific regulation. Schizophrenia Research, 2017, 183, 82-87.	1.1	17
49	Age at first birth in women is genetically associated with increased risk of schizophrenia. Scientific Reports, 2018, 8, 10168.	1.6	17
50	TECHNICAL ARTICLE: SNP-SCALE: SNP scoring by colour and length exclusion. Molecular Ecology Notes, 2007, 7, 377-388.	1.7	15
51	Whole exome sequencing and <scp>DNA</scp> methylation analysis in a clinical amyotrophic lateral sclerosis cohort. Molecular Genetics & Genomic Medicine, 2017, 5, 418-428.	0.6	14
52	Sizing up whole-genome sequencing studies of common diseases. Nature Genetics, 2018, 50, 635-637.	9.4	13
53	Gene networks associated with non-syndromic intellectual disability. Journal of Neurogenetics, 2018, 32, 6-14.	0.6	13
54	Rare variants are common in schizophrenia. Nature Neuroscience, 2016, 19, 1426-1428.	7.1	11

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55	Analysis of common genetic variation and rare CNVs in the Australian Autism Biobank. Molecular Autism, 2021, 12, 12.	2.6	11
56	Mapping quantitative trait loci in a wild population using linkage and linkage disequilibrium analyses. Genetical Research, 2010, 92, 273-281.	0.3	6
57	No evidence for warming climate theory of coat colour change in Soay sheep: a comment on Maloney et al Biology Letters, 2010, 6, 678-679.	1.0	6
58	Mapping and differential expression analysis from shortâ€read RNA‣eq data in model organisms. Quantitative Biology, 2016, 4, 22-35.	0.3	3
59	Trans-eQTLs identified in whole blood have limited influence on complex disease biology. European Journal of Human Genetics, 2018, 26, 1361-1368.	1.4	3
60	Comparing schizophrenia symptoms in the <scp>I</scp> ban of <scp>S</scp> arawak with other populations to elucidate clinical heterogeneity. Asia-Pacific Psychiatry, 2015, 7, 36-44.	1.2	2
61	Examining the Impact of Imputation Errors on Fine-Mapping Using DNA Methylation QTL as a Model Trait. Genetics, 2019, 212, 577-586.	1.2	2
62	Australian Parkinson's Genetics Study (APGS): pilot (n=1532). BMJ Open, 2022, 12, e052032.	0.8	1