Oliver S P Davis

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	A Polygenic Approach to Understanding Resilience to Peer Victimisation. Behavior Genetics, 2022, 52, 1-12.	1.4	6
2	Shared genetic risk between eating disorder―and substanceâ€useâ€related phenotypes: Evidence from genomeâ€wide association studies. Addiction Biology, 2021, 26, e12880.	1.4	28
3	Positive wellbeing and resilience following adolescent victimisation: An exploration into protective factors across development. JCPP Advances, 2021, 1, e12024.	1.4	5
4	Peer victimisation during adolescence and its impact on wellbeing in adulthood: a prospective cohort study. BMC Public Health, 2021, 21, 148.	1.2	18
5	Understanding the potential and pitfalls of digital phenotypes to measure population mental health and wellbeing. Lancet, The, 2021, 398, S10.	6.3	1
6	Biotic analogies for self-organising cities. Environment and Planning B: Urban Analytics and City Science, 2020, 47, 268-286.	1.0	6
7	The association of DNA methylation with body mass index: distinguishing between predictors and biomarkers. Clinical Epigenetics, 2020, 12, 50.	1.8	36
8	Mapping Population Vulnerability and Community Support during COVID-19. International Journal of Population Data Science, 2020, 5, 1409.	0.1	2
9	Views on social media and its linkage to longitudinal data from two generations of a UK cohort study. Wellcome Open Research, 2020, 5, 44.	0.9	2
10	Views on social media and its linkage to longitudinal data from two generations of a UK cohort study. Wellcome Open Research, 2020, 5, 44.	0.9	2
11	Participant acceptability of digital footprint data collection strategies: an exemplar approach to participant engagement and involvement in the ALSPAC birth cohort study International Journal of Population Data Science, 2020, 5, 1728.	0.1	2
12	Genome-wide association study identifies eight risk loci and implicates metabo-psychiatric origins for anorexia nervosa. Nature Genetics, 2019, 51, 1207-1214.	9.4	641
13	Associations Between Attention-Deficit/Hyperactivity Disorder and Various Eating Disorders: A Swedish Nationwide Population Study Using Multiple Genetically Informative Approaches. Biological Psychiatry, 2019, 86, 577-586.	0.7	43
14	Identifying Critical Points of Trajectories of Depressive Symptoms from Childhood to Young Adulthood. Journal of Youth and Adolescence, 2019, 48, 815-827.	1.9	97
15	Schizophrenia liability shares common molecular genetic risk factors with sleep duration and nightmares in childhood. Wellcome Open Research, 2019, 4, 15.	0.9	5
16	Schizophrenia liability shares common molecular genetic risk factors with sleep duration and nightmares in childhood. Wellcome Open Research, 2019, 4, 15.	0.9	4
17	Investigation of common, low-frequency and rare genome-wide variation in anorexia nervosa. Molecular Psychiatry, 2018, 23, 1169-1180.	4.1	32
18	Evaluation of the causal effects between subjective wellbeing and cardiometabolic health: mendelian randomisation study. BMJ: British Medical Journal, 2018, 362, k3788.	2.4	59

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19	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	6.0	1,085
20	Evidence for three genetic loci involved in both anorexia nervosa risk and variation of body mass index. Molecular Psychiatry, 2017, 22, 192-201.	4.1	63
21	Significant Locus and Metabolic Genetic Correlations Revealed in Genome-Wide Association Study of Anorexia Nervosa. American Journal of Psychiatry, 2017, 174, 850-858.	4.0	410
22	Genetic and environmental correlations between subjective wellbeing and experience of life events in adolescence. European Child and Adolescent Psychiatry, 2017, 26, 1119-1127.	2.8	15
23	Social support and mental health in late adolescence are correlated for genetic, as well as environmental, reasons. Scientific Reports, 2017, 7, 13088.	1.6	27
24	Personalized Media: A Genetically Informative Investigation of Individual Differences in Online Media Use. PLoS ONE, 2017, 12, e0168895.	1.1	10
25	Moderators of wellbeing interventions: Why do some people respond more positively than others?. PLoS ONE, 2017, 12, e0187601.	1.1	24
26	Exploring the Genetic Etiology of Trust in Adolescents: Combined Twin and DNA Analyses. Twin Research and Human Genetics, 2016, 19, 638-646.	0.3	14
27	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
28	Heritability and genome-wide analyses of problematic peer relationships during childhood and adolescence. Human Genetics, 2015, 134, 539-551.	1.8	13
29	From observational to dynamic genetics. Frontiers in Genetics, 2014, 5, 6.	1.1	19
30	Genome-Wide Association Study of Receptive Language Ability of 12-Year-Olds. Journal of Speech, Language, and Hearing Research, 2014, 57, 96-105.	0.7	24
31	Common variation near ROBO2 is associated with expressive vocabulary in infancy. Nature Communications, 2014, 5, 4831.	5.8	82
32	Childhood intelligence is heritable, highly polygenic and associated with FNBP1L. Molecular Psychiatry, 2014, 19, 253-258.	4.1	241
33	Genetic predictors of antidepressant side effects: A grouped candidate gene approach in the Genome-Based Therapeutic Drugs for Depression (GENDEP) study. Journal of Psychopharmacology, 2014, 28, 142-150.	2.0	18
34	Genetic differences in cytochrome P450 enzymes and antidepressant treatment response. Journal of Psychopharmacology, 2014, 28, 133-141.	2.0	75
35	Why do spatial abilities predict mathematical performance?. Developmental Science, 2014, 17, 462-470.	1.3	67
36	A genome-wide association study of anorexia nervosa. Molecular Psychiatry, 2014, 19, 1085-1094.	4.1	282

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37	The correlation between reading and mathematics ability at age twelve has a substantial genetic component. Nature Communications, 2014, 5, 4204.	5.8	72
38	Relationship between obesity and the risk of clinically significant depression: Mendelian randomisation study. British Journal of Psychiatry, 2014, 205, 24-28.	1.7	62
39	DNA Evidence for Strong Genome-Wide Pleiotropy of Cognitive and Learning Abilities. Behavior Genetics, 2013, 43, 267-273.	1.4	91
40	Understanding the science-learning environment: A genetically sensitive approach. Learning and Individual Differences, 2013, 23, 145-150.	1.5	11
41	A common variant in Myosin-18B contributes to mathematical abilities in children with dyslexia and intraparietal sulcus variability in adults. Translational Psychiatry, 2013, 3, e229-e229.	2.4	28
42	Twins Early Development Study (TEDS): A Genetically Sensitive Investigation of Cognitive and Behavioral Development From Childhood to Young Adulthood. Twin Research and Human Genetics, 2013, 16, 117-125.	0.3	247
43	Common DNA Markers Can Account for More Than Half of the Genetic Influence on Cognitive Abilities. Psychological Science, 2013, 24, 562-568.	1.8	135
44	First Genome-Wide Association Study on Anxiety-Related Behaviours in Childhood. PLoS ONE, 2013, 8, e58676.	1.1	61
45	Genetics of Callous-Unemotional Behavior in Children. PLoS ONE, 2013, 8, e65789.	1.1	45
46	Common variants at 6q22 and 17q21 are associated with intracranial volume. Nature Genetics, 2012, 44, 539-544.	9.4	126
47	Common variants at 12q15 and 12q24 are associated with infant head circumference. Nature Genetics, 2012, 44, 532-538.	9.4	130
48	Chaotic Homes and Children's Disruptive Behavior. Psychological Science, 2012, 23, 643-650.	1.8	67
49	Genomeâ€wide association analysis of eating disorderâ€related symptoms, behaviors, and personality traits. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 803-811.	1.1	52
50	A genetic association study of DNA methylation levels in the DRD4 gene region finds associations with nearby SNPs. Behavioral and Brain Functions, 2012, 8, 31.	1.4	36
51	Visual analysis of geocoded twin data puts nature and nurture on the map. Molecular Psychiatry, 2012, 17, 867-874.	4.1	52
52	Socioeconomic Status (SES) and Children's Intelligence (IQ): In a UK-Representative Sample SES Moderates the Environmental, Not Genetic, Effect on IQ. PLoS ONE, 2012, 7, e30320.	1.1	200
53	Chaotic homes and school achievement: a twin study. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2011, 52, 1212-1220.	3.1	55
54	Visualizing genetic similarity at the symptom level: The example of learning disabilities. Behavioral and Brain Sciences, 2010, 33, 155-157.	0.4	0

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55	A Genome-Wide Association Study of Social and Non-Social Autistic-Like Traits in the General Population Using Pooled DNA, 500ÂK SNP Microarrays and Both Community and Diagnosed Autism Replication Samples. Behavior Genetics, 2010, 40, 31-45.	1.4	49
56	Response to comment by Stuart Macgregor. Behavior Genetics, 2010, 40, 48-48.	1.4	0
57	A Three-Stage Genome-Wide Association Study of General Cognitive Ability: Hunting the Small Effects. Behavior Genetics, 2010, 40, 759-767.	1.4	74
58	Genetics of Learning Abilities and Disabilities: Recent Developments from the UK and Possible Directions for Research in China. Behavior Genetics, 2010, 40, 297-305.	1.4	4
59	In search of genes associated with risk for psychopathic tendencies in children: a twoâ€stage genomeâ€wide association study of pooled DNA. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2010, 51, 780-788.	3.1	76
60	A genomeâ€wide association study identifies multiple loci associated with mathematics ability and disability. Genes, Brain and Behavior, 2010, 9, 234-247.	1.1	100
61	The heritability of general cognitive ability increases linearly from childhood to young adulthood. Molecular Psychiatry, 2010, 15, 1112-1120.	4.1	492
62	The nature (and nurture) of children's perceptions of family chaos. Learning and Individual Differences, 2010, 20, 549-553.	1.5	33
63	DNA methylation profiling using bisulfite-based epityping of pooled genomic DNA. Methods, 2010, 52, 255-258.	1.9	43
64	Learning abilities and disabilities: Generalist genes in early adolescence. Cognitive Neuropsychiatry, 2009, 14, 312-331.	0.7	77
65	The SNPMaP package for R: a framework for genome-wide association using DNA pooling on microarrays. Bioinformatics, 2009, 25, 281-283.	1.8	27
66	A Twin Study of the Genetics of High Cognitive Ability Selected from 11,000 Twin Pairs in Six Studies from Four Countries. Behavior Genetics, 2009, 39, 359-370.	1.4	54
67	Bisulfite-based epityping on pooled genomic DNA provides an accurate estimate of average group DNA methylation. Epigenetics and Chromatin, 2009, 2, 3.	1.8	60
68	The future of genetics in psychology and psychiatry: microarrays, genomeâ€wide association, and nonâ€coding RNA. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2009, 50, 63-71.	3.1	52
69	Common disorders are quantitative traits. Nature Reviews Genetics, 2009, 10, 872-878.	7.7	603
70	Dramatic Increase in Heritability of Cognitive Development from Early to Middle Childhood. Psychological Science, 2009, 20, 1301-1308.	1.8	77
71	Genomeâ€wide quantitative trait locus association scan of general cognitive ability using pooled DNA and 500K single nucleotide polymorphism microarrays. Genes, Brain and Behavior, 2008, 7, 435-446. 	1.1	127
72	Generalist genes and the Internet generation: etiology of learning abilities by web testing at age 10. Genes, Brain and Behavior, 2008, 7, 455-462.	1.1	37

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73	Testing replication of a 5-SNP set for general cognitive ability in six population samples. European Journal of Human Genetics, 2008, 16, 1388-1395.	1.4	8
74	Increasing Heritability of BMI and Stronger Associations With the FTO Gene Over Childhood. Obesity, 2008, 16, 2663-2668.	1.5	151
75	g in middle childhood: Moderate genetic and shared environmental influence using diverse measures of general cognitive ability at 7, 9 and 10Âyears in a large population sample of twins. Intelligence, 2008, 36, 68-80.	1.6	27
76	Internet Cognitive Testing of Large Samples Needed in Genetic Research. Twin Research and Human Genetics, 2007, 10, 554-563.	0.3	138
77	Mapping the genetic and environmental aetiology of autistic traits in Sweden and the United Kingdom. JCPP Advances, 0, , .	1.4	1