Desmond D Campbell

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/7700880/publications.pdf

Version: 2024-02-01

29 papers 2,368 citations

16 h-index 28 g-index

30 all docs 30 docs citations

30 times ranked

5653 citing authors

#	Article	IF	CITATIONS
1	Effects of depression on employment and social outcomes: a Mendelian randomisation study. Journal of Epidemiology and Community Health, 2022, 76, 563-571.	3.7	17
2	Mental health and health behaviours before and during the initial phase of the COVID-19 lockdown: longitudinal analyses of the UK Household Longitudinal Study. Journal of Epidemiology and Community Health, 2021, 75, jech-2020-215060.	3.7	323
3	Effects of increased body mass index on employment status: a Mendelian randomisation study. International Journal of Obesity, 2021, 45, 1790-1801.	3.4	4
4	Clinical burden, risk factor impact and outcomes following myocardial infarction and stroke: A 25-year individual patient level linkage study. Lancet Regional Health - Europe, The, 2021, 7, 100141.	5. 6	18
5	The causal effects of health conditions and risk factors on social and socioeconomic outcomes: Mendelian randomization in UK Biobank. International Journal of Epidemiology, 2020, 49, 1661-1681.	1.9	33
6	Gene Variants at Loci Related to Blood Pressure Account for Variation in Response to Antihypertensive Drugs Between Black and White Individuals. Hypertension, 2019, 74, 614-622.	2.7	14
7	Multifactorial disease risk calculator: Risk prediction for multifactorial disease pedigrees. Genetic Epidemiology, 2018, 42, 130-133.	1.3	6
8	Salt stress in the renal tubules is linked to TAL-specific expression of uromodulin and an upregulation of heat shock genes. Physiological Genomics, 2018, 50, 964-972.	2.3	7
9	Interplay between Schizophrenia Polygenic Risk Score and Childhood Adversity in First-Presentation Psychotic Disorder: A Pilot Study. PLoS ONE, 2016, 11, e0163319.	2.5	52
10	Genetic study of congenital bile-duct dilatation identifies de novo and inherited variants in functionally related genes. BMC Medical Genomics, 2016, 9, 75.	1.5	11
11	Two subtypes of intervertebral disc degeneration distinguished by large-scale population-based study. Spine Journal, 2016, 16, 1079-1089.	1.3	51
12	Sacral agenesis: a pilot whole exome sequencing and copy number study. BMC Medical Genetics, 2016, 17, 98.	2.1	15
13	Local True Discovery Rate Weighted Polygenic Scores Using GWAS Summary Data. Behavior Genetics, 2016, 46, 573-582.	2.1	15
14	A Single Nucleotide Variant in HNF- $1\hat{l}^2$ is Associated with Maturity-Onset Diabetes of the Young in a Large Chinese Family. Iranian Journal of Public Health, 2016, 45, 170-8.	0.5	1
15	Cross-Disorder Genome-Wide Analyses Suggest a Complex Genetic Relationship Between Tourette's Syndrome and OCD. American Journal of Psychiatry, 2015, 172, 82-93.	7.2	117
16	Cost effective assay choice for rare disease study designs. Orphanet Journal of Rare Diseases, 2015, 10, 10.	2.7	3
17	Identifying Gene-Environment Interactions in Schizophrenia: Contemporary Challenges for Integrated, Large-scale Investigations. Schizophrenia Bulletin, 2014, 40, 729-736.	4.3	229
18	Error in sample size formula. Journal of Human Reproductive Sciences, 2014, 7, 155.	0.9	0

#	Article	IF	CITATION
19	The emerging molecular architecture of schizophrenia, polygenic risk scores and the clinical implications for GxE research. Social Psychiatry and Psychiatric Epidemiology, 2014, 49, 169-182.	3.1	68
20	Cognitive impairment among children at-risk for schizophrenia. Journal of Psychiatric Research, 2014, 50, 92-99.	3.1	34
21	Partitioning the Heritability of Tourette Syndrome and Obsessive Compulsive Disorder Reveals Differences in Genetic Architecture. PLoS Genetics, 2013, 9, e1003864.	3.5	241
22	CNV Analysis in Tourette Syndrome Implicates Large Genomic Rearrangements in COL8A1 and NRXN1. PLoS ONE, 2013, 8, e59061.	2.5	70
23	<i>CYP2C19</i> genotype predicts steady state escitalopram concentration in GENDEP. Journal of Psychopharmacology, 2012, 26, 398-407.	4.0	69
24	Reconstructing Native American population history. Nature, 2012, 488, 370-374.	27.8	699
25	Amerind Ancestry, Socioeconomic Status and the Genetics of Type 2 Diabetes in a Colombian Population. PLoS ONE, 2012, 7, e33570.	2.5	47
26	Software for generating liability distributions for pedigrees conditional on their observed disease states and covariates. Genetic Epidemiology, 2010, 34, 159-170.	1.3	11
27	Suicidal ideation during treatment of depression with escitalopram and nortriptyline in Genome-Based Therapeutic Drugs for Depression (GENDEP): a clinical trial. BMC Medicine, 2009, 7, 60.	5 . 5	43
28	DSMâ€N combined type ADHD shows familial association with sibling trait scores: A sampling strategy for QTL linkage. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 1450-1460.	1.7	129
29	Linkage to Chromosome 1p36 for Attention-Deficit/Hyperactivity Disorder Traits in School and Home Settings. Biological Psychiatry, 2008, 64, 571-576.	1.3	41