Archie Campbell

List of Publications by Citations

Source: https://exaly.com/author-pdf/7275617/archie-campbell-publications-by-citations.pdf

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

 170
 6,680
 41
 79

 papers
 citations
 h-index
 g-index

 203
 10,401
 11
 4.85

 ext. papers
 ext. citations
 avg, IF
 L-index

#	Paper	IF	Citations
170	Genomic insights into the origin of farming in the ancient Near East. <i>Nature</i> , 2016 , 536, 419-24	50.4	485
169	Genetic analysis of over 1 million people identifies 535 new loci associated with blood pressure traits. <i>Nature Genetics</i> , 2018 , 50, 1412-1425	36.3	386
168	Multi-ethnic genome-wide association study for atrial fibrillation. <i>Nature Genetics</i> , 2018 , 50, 1225-1233	36.3	277
167	Genomic analyses identify hundreds of variants associated with age at menarche and support a role for puberty timing in cancer risk. <i>Nature Genetics</i> , 2017 , 49, 834-841	36.3	257
166	Cohort Profile: Generation Scotland: Scottish Family Health Study (GS:SFHS). The study, its participants and their potential for genetic research on health and illness. <i>International Journal of Epidemiology</i> , 2013 , 42, 689-700	7.8	237
165	Large-scale genomic analyses link reproductive aging to hypothalamic signaling, breast cancer susceptibility and BRCA1-mediated DNA repair. <i>Nature Genetics</i> , 2015 , 47, 1294-1303	36.3	226
164	PCSK9 genetic variants and risk of type 2 diabetes: a mendelian randomisation study. <i>Lancet Diabetes and Endocrinology,the</i> , 2017 , 5, 97-105	18.1	225
163	Genome-wide association study of alcohol consumption and genetic overlap with other health-related traits in UK Biobank (N=112 117). <i>Molecular Psychiatry</i> , 2017 , 22, 1376-1384	15.1	225
162	Ultra-High-Throughput Clinical Proteomics Reveals Classifiers of COVID-19 Infection. <i>Cell Systems</i> , 2020 , 11, 11-24.e4	10.6	219
161	A catalog of genetic loci associated with kidney function from analyses of a million individuals. <i>Nature Genetics</i> , 2019 , 51, 957-972	36.3	217
160	Large-scale analyses of common and rare variants identify 12 new loci associated with atrial fibrillation. <i>Nature Genetics</i> , 2017 , 49, 946-952	36.3	176
159	Genome-wide association analyses for lung function and chronic obstructive pulmonary disease identify new loci and potential druggable targets. <i>Nature Genetics</i> , 2017 , 49, 416-425	36.3	170
158	Molecular genetic contributions to socioeconomic status and intelligence. <i>Intelligence</i> , 2014 , 44, 26-32	3	131
157	Mental health before and during the COVID-19 pandemic in two longitudinal UK population cohorts. <i>British Journal of Psychiatry</i> , 2020 , 1-10	5.4	123
156	Target genes, variants, tissues and transcriptional pathways influencing human serum urate levels. Nature Genetics, 2019 , 51, 1459-1474	36.3	122
155	Directional dominance on stature and cognition indiverse human populations. <i>Nature</i> , 2015 , 523, 459-4	63 0.4	119
154	Investigating the possible causal association of smoking with depression and anxiety using Mendelian randomisation meta-analysis: the CARTA consortium. <i>BMJ Open</i> , 2014 , 4, e006141	3	115

153	Cardiac Troponin T and Troponin I in the General Population. Circulation, 2019, 139, 2754-2764	16.7	90
152	Genetic variants linked to education predict longevity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016 , 113, 13366-13371	11.5	90
151	Genetic Evidence for a Link Between Favorable Adiposity and Lower Risk of Type 2 Diabetes, Hypertension, and Heart Disease. <i>Diabetes</i> , 2016 , 65, 2448-60	0.9	86
150	Exploration of haplotype research consortium imputation for genome-wide association studies in 20,032 Generation Scotland participants. <i>Genome Medicine</i> , 2017 , 9, 23	14.4	85
149	Novel Blood Pressure Locus and Gene Discovery Using Genome-Wide Association Study and Expression Data Sets From Blood and the Kidney. <i>Hypertension</i> , 2017 ,	8.5	85
148	Association analyses identify 31 new risk loci for colorectal cancer susceptibility. <i>Nature Communications</i> , 2019 , 10, 2154	17.4	81
147	Genome-wide meta-analysis associates HLA-DQA1/DRB1 and LPA and lifestyle factors with human longevity. <i>Nature Communications</i> , 2017 , 8, 910	17.4	78
146	Epigenetic prediction of complex traits and death. <i>Genome Biology</i> , 2018 , 19, 136	18.3	77
145	Epidemiology and Heritability of Major Depressive Disorder, Stratified by Age of Onset, Sex, and Illness Course in Generation Scotland: Scottish Family Health Study (GS:SFHS). <i>PLoS ONE</i> , 2015 , 10, e014	1 27 97	75
144	Effect of Smoking on Blood Pressure and Resting Heart Rate: A Mendelian Randomization Meta-Analysis in the CARTA Consortium. <i>Circulation: Cardiovascular Genetics</i> , 2015 , 8, 832-41		70
143	Comparison between High-Sensitivity Cardiac Troponin T and Cardiac Troponin I in a Large General Population Cohort. <i>Clinical Chemistry</i> , 2018 , 64, 1607-1616	5.5	61
142	Genome-wide meta-analysis of 158,000 individuals of European ancestry identifies three loci associated with chronic back pain. <i>PLoS Genetics</i> , 2018 , 14, e1007601	6	60
141	Multi-ancestry genome-wide gene-smoking interaction study of 387,272 individuals identifies new loci associated with serum lipids. <i>Nature Genetics</i> , 2019 , 51, 636-648	36.3	59
140	A Large-Scale Multi-ancestry Genome-wide Study Accounting for Smoking Behavior Identifies Multiple Significant Loci for Blood Pressure. <i>American Journal of Human Genetics</i> , 2018 , 102, 375-400	11	59
139	Investigating the relationship between DNA methylation age acceleration and risk factors for Alzheimer's disease. <i>Alzheimer's and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2018 , 10, 429-437	5.2	57
138	Pedigree- and SNP-Associated Genetics and Recent Environment are the Major Contributors to Anthropometric and Cardiometabolic Trait Variation. <i>PLoS Genetics</i> , 2016 , 12, e1005804	6	50
137	Differential effects of the APOE e4 allele on different domains of cognitive ability across the life-course. <i>European Journal of Human Genetics</i> , 2016 , 24, 919-23	5.3	47
136	Genome-Wide and Abdominal MRI Data Provide Evidence That a Genetically Determined Favorable Adiposity Phenotype Is Characterized by Lower Ectopic Liver Fat and Lower Risk of Type 2 Diabetes, Heart Disease, and Hypertension. <i>Diabetes</i> , 2019 , 68, 207-219	0.9	46

135	The trans-ancestral genomic architecture of glycemic traits. <i>Nature Genetics</i> , 2021 , 53, 840-860	36.3	44
134	An epigenome-wide association study of sex-specific chronological ageing. <i>Genome Medicine</i> , 2019 , 12, 1	14.4	43
133	Mental health during the COVID-19 pandemic in two longitudinal UK population cohorts		42
132	Multi-ancestry study of blood lipid levels identifies four loci interacting with physical activity. <i>Nature Communications</i> , 2019 , 10, 376	17.4	41
131	Pedigree and genotyping quality analyses of over 10,000 DNA samples from the Generation Scotland: Scottish Family Health Study. <i>BMC Medical Genetics</i> , 2013 , 14, 38	2.1	41
130	Genetic and Environmental Risk for Chronic Pain and the Contribution of Risk Variants for Major Depressive Disorder: A Family-Based Mixed-Model Analysis. <i>PLoS Medicine</i> , 2016 , 13, e1002090	11.6	41
129	Exploring causality in the association between circulating 25-hydroxyvitamin D and colorectal cancer risk: a large Mendelian randomisation study. <i>BMC Medicine</i> , 2018 , 16, 142	11.4	40
128	Associations of autozygosity with a broad range of human phenotypes. <i>Nature Communications</i> , 2019 , 10, 4957	17.4	40
127	Stratification by smoking status reveals an association of CHRNA5-A3-B4 genotype with body mass index in never smokers. <i>PLoS Genetics</i> , 2014 , 10, e1004799	6	40
126	Chronic pain, depression and cardiovascular disease linked through a shared genetic predisposition: Analysis of a family-based cohort and twin study. <i>PLoS ONE</i> , 2017 , 12, e0170653	3.7	40
125	Epigenetic measures of ageing predict the prevalence and incidence of leading causes of death and disease burden. <i>Clinical Epigenetics</i> , 2020 , 12, 115	7.7	40
124	Multiancestry Genome-Wide Association Study of Lipid Levels Incorporating Gene-Alcohol Interactions. <i>American Journal of Epidemiology</i> , 2019 , 188, 1033-1054	3.8	39
123	Heavier smoking may lead to a relative increase in waist circumference: evidence for a causal relationship from a Mendelian randomisation meta-analysis. The CARTA consortium. <i>BMJ Open</i> , 2015 , 5, e008808	3	39
122	Self-reported medication use validated through record linkage to national prescribing data. <i>Journal of Clinical Epidemiology</i> , 2018 , 94, 132-142	5.7	38
121	A meta-analysis of genome-wide association studies of epigenetic age acceleration. <i>PLoS Genetics</i> , 2019 , 15, e1008104	6	38
120	Epigenetic signatures of starting and stopping smoking. <i>EBioMedicine</i> , 2018 , 37, 214-220	8.8	36
119	Recent genomic heritage in Scotland. <i>BMC Genomics</i> , 2015 , 16, 437	4.5	33
118	Factors associated with sharing e-mail information and mental health survey participation in large population cohorts. <i>International Journal of Epidemiology</i> , 2020 , 49, 410-421	7.8	32

117	A time-resolved proteomic and prognostic map of COVID-19. Cell Systems, 2021, 12, 780-794.e7	10.6	32
116	Novel genetic associations for blood pressure identified via gene-alcohol interaction in up to 570K individuals across multiple ancestries. <i>PLoS ONE</i> , 2018 , 13, e0198166	3.7	31
115	and Loci Identified through Large-Scale Exome Chip Analysis Regulate Kidney Development and Function. <i>Journal of the American Society of Nephrology: JASN</i> , 2017 , 28, 981-994	12.7	30
114	Common genetic variants explain the majority of the correlation between height and intelligence: the generation Scotland study. <i>Behavior Genetics</i> , 2014 , 44, 91-6	3.2	30
113	Quantifying the extent to which index event biases influence large genetic association studies. <i>Human Molecular Genetics</i> , 2017 , 26, 1018-1030	5.6	30
112	Assessing the genetic overlap between BMI and cognitive function. <i>Molecular Psychiatry</i> , 2016 , 21, 1477	7 -83 1	29
111	Genetic insights into biological mechanisms governing human ovarian ageing. <i>Nature</i> , 2021 , 596, 393-39	930.4	28
110	Meta-analysis of 49 549 individuals imputed with the 1000 Genomes Project reveals an exonic damaging variant in ANGPTL4 determining fasting TG levels. <i>Journal of Medical Genetics</i> , 2016 , 53, 441-	9 ^{5.8}	27
109	Shared Genetics and Couple-Associated Environment Are Major Contributors to the Risk of Both Clinical and Self-Declared Depression. <i>EBioMedicine</i> , 2016 , 14, 161-167	8.8	26
108	Evidence for large-scale gene-by-smoking interaction effects on pulmonary function. <i>International Journal of Epidemiology</i> , 2017 , 46, 894-904	7.8	25
107	The power of genetic diversity in genome-wide association studies of lipids. <i>Nature</i> , 2021 ,	50.4	24
106	Genome-wide association study of antidepressant treatment resistance in a population-based cohort using health service prescription data and meta-analysis with GENDEP. <i>Pharmacogenomics Journal</i> , 2020 , 20, 329-341	3.5	24
105	The Dementias Platform UK (DPUK) Data Portal. European Journal of Epidemiology, 2020, 35, 601-611	12.1	23
104	Flexible and scalable diagnostic filtering of genomic variants using G2P with Ensembl VEP. <i>Nature Communications</i> , 2019 , 10, 2373	17.4	22
103	Parent of origin genetic effects on methylation in humans are common and influence complex trait variation. <i>Nature Communications</i> , 2019 , 10, 1383	17.4	21
102	Genetic Stratification to Identify Risk Groups for Alzheimer& Disease. <i>Journal of Alzheimerr</i> Disease, 2017 , 57, 275-283	4.3	21
101	Recurrent Coding Sequence Variation Explains Only A Small Fraction of the Genetic Architecture of Colorectal Cancer. <i>Scientific Reports</i> , 2015 , 5, 16286	4.9	21
100	Blunted medial prefrontal cortico-limbic reward-related effective connectivity and depression. <i>Brain</i> , 2020 , 143, 1946-1956	11.2	19

99	Dissection of major depressive disorder using polygenic risk scores for schizophrenia in two independent cohorts. <i>Translational Psychiatry</i> , 2016 , 6, e938	8.6	19
98	Bayesian reassessment of the epigenetic architecture of complex traits. <i>Nature Communications</i> , 2020 , 11, 2865	17.4	18
97	Homozygous loss-of-function variants in European cosmopolitan and isolate populations. <i>Human Molecular Genetics</i> , 2015 , 24, 5464-74	5.6	18
96	Novel Urinary Peptidomic Classifier Predicts Incident Heart Failure. <i>Journal of the American Heart Association</i> , 2017 , 6,	6	17
95	Meta-analysis of exome array data identifies six novel genetic loci for lung function. <i>Wellcome Open Research</i> , 2018 , 3, 4	4.8	16
94	Characterisation of an inflammation-related epigenetic score and its association with cognitive ability. <i>Clinical Epigenetics</i> , 2020 , 12, 113	7.7	15
93	A multi-ancestry genome-wide study incorporating gene-smoking interactions identifies multiple new loci for pulse pressure and mean arterial pressure. <i>Human Molecular Genetics</i> , 2019 , 28, 2615-2633	5.6	14
92	Within-sibship GWAS improve estimates of direct genetic effects		14
91	Urinary peptides in heart failure: a link to molecular pathophysiology. <i>European Journal of Heart Failure</i> , 2021 , 23, 1875-1887	12.3	14
90	Structural brain correlates of serum and epigenetic markers of inflammation in major depressive disorder. <i>Brain, Behavior, and Immunity</i> , 2021 , 92, 39-48	16.6	14
89	Identification of novel differentially methylated sites with potential as clinical predictors of impaired respiratory function and COPD. <i>EBioMedicine</i> , 2019 , 43, 576-586	8.8	13
88	Polygenic risk of ischemic stroke is associated with cognitive ability. <i>Neurology</i> , 2016 , 86, 611-8	6.5	13
87	Genetic and environmental determinants of stressful life events and their overlap with depression and neuroticism. <i>Wellcome Open Research</i> , 2018 , 3, 11	4.8	13
86	An epigenetic score for BMI based on DNA methylation correlates with poor physical health and major disease in the Lothian Birth Cohort. <i>International Journal of Obesity</i> , 2019 , 43, 1795-1802	5.5	12
85	Regional variation in health is predominantly driven by lifestyle rather than genetics. <i>Nature Communications</i> , 2017 , 8, 801	17.4	11
84	Genetic and environmental determinants of stressful life events and their overlap with depression and neuroticism. <i>Wellcome Open Research</i> , 2018 , 3, 11	4.8	11
83	Identification of epigenome-wide DNA methylation differences between carriers of APOE A and APOE alleles. <i>Genome Medicine</i> , 2021 , 13, 1	14.4	11
82	The genetic landscape of Scotland and the Isles. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019 , 116, 19064-19070	11.5	10

(2022-2015)

81	Current versus lifetime depression, APOE variation, and their interaction on cognitive performance in younger and older adults. <i>Psychosomatic Medicine</i> , 2015 , 77, 480-92	3.7	10	
80	Sex-Differences in the Metabolic Health of Offspring of Parents with Diabetes: A Record-Linkage Study. <i>PLoS ONE</i> , 2015 , 10, e0134883	3.7	10	
79	Cohort profile for the STratifying Resilience and Depression Longitudinally (STRADL) study: A depression-focused investigation of Generation Scotland, using detailed clinical, cognitive, and neuroimaging assessments <i>Wellcome Open Research</i> , 2019 , 4, 185	4.8	10	
78	A genome-wide association study finds genetic variants associated with neck or shoulder pain in UK Biobank. <i>Human Molecular Genetics</i> , 2020 , 29, 1396-1404	5.6	10	
77	Electronic health record and genome-wide genetic data in Generation Scotland participants. <i>Wellcome Open Research</i> , 2017 , 2, 85	4.8	9	
76	Insights into the genetic basis of retinal detachment. <i>Human Molecular Genetics</i> , 2020 , 29, 689-702	5.6	9	
75	Genetic stratification of depression in UK Biobank. <i>Translational Psychiatry</i> , 2020 , 10, 163	8.6	8	
74	DNA methylation outlier burden, health, and ageing in Generation Scotland and the Lothian Birth Cohorts of 1921 and 1936. <i>Clinical Epigenetics</i> , 2020 , 12, 49	7.7	8	
73	Phenome-wide association analysis of LDL-cholesterol lowering genetic variants in PCSK9. <i>BMC Cardiovascular Disorders</i> , 2019 , 19, 240	2.3	8	
72	Investigating genetic links between grapheme-colour synaesthesia and neuropsychiatric traits. <i>Philosophical Transactions of the Royal Society B: Biological Sciences</i> , 2019 , 374, 20190026	5.8	8	
71	Assessment of dried blood spots for DNA methylation profiling. Wellcome Open Research, 2019, 4, 44	4.8	8	
70	Epigenetic clocks predict prevalence and incidence of leading causes of death and disease burden		8	
69	Genetic comorbidity between major depression and cardio-metabolic traits, stratified by age at onset of major depression. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2020 , 183, 309-330	3.5	8	
68	Phenotypic and genetic analysis of cognitive performance in Major Depressive Disorder in the Generation Scotland: Scottish Family Health Study. <i>Translational Psychiatry</i> , 2018 , 8, 63	8.6	7	
67	Birth weight associations with DNA methylation differences in an adult population. <i>Epigenetics</i> , 2021 , 16, 783-796	5.7	7	
66	DNA methylation signatures of aggression and closely related constructs: A meta-analysis of epigenome-wide studies across the lifespan. <i>Molecular Psychiatry</i> , 2021 , 26, 2148-2162	15.1	7	
65	CovidLife: a resource to understand mental health, well-being and behaviour during the COVID-19 pandemic in the UK. <i>Wellcome Open Research</i> ,6, 176	4.8	7	
64	Polygenic prediction of educational attainment within and between families from genome-wide association analyses in 3 million individuals <i>Nature Genetics</i> , 2022 ,	36.3	7	

63	Exome sequencing to detect rare variants associated with general cognitive ability: a pilot study. <i>Twin Research and Human Genetics</i> , 2015 , 18, 117-25	2.2	6
62	A proteomic survival predictor for COVID-19 patients in intensive care 2022 , 1, e0000007		6
61	Meta-analysis of exome array data identifies six novel genetic loci for lung function. <i>Wellcome Open Research</i> , 3, 4	4.8	6
60	Global Biobank Meta-analysis Initiative: powering genetic discovery across human diseases		6
59	Genetic stratification of depression in UK Biobank suggests a subgroup linked to age of natural menop	ause	6
58	Prescreening for European Prevention of Alzheimer Dementia (EPAD) trial-ready cohort: impact of AD risk factors and recruitment settings. <i>Alzheimerns Research and Therapy</i> , 2020 , 12, 8	9	6
57	Within-sibship genome-wide association analyses decrease bias in estimates of direct genetic effects <i>Nature Genetics</i> , 2022 , 54, 581-592	36.3	6
56	Birth weight associations with psychiatric and physical health, cognitive function, and DNA methylation differences in an adult population		5
55	Genome-wide association study of cardiac troponin I in the general population. <i>Human Molecular Genetics</i> , 2021 , 30, 2027-2039	5.6	5
54	Early life predictors of late life cerebral small vessel disease in four prospective cohort studies. Brain, 2021 ,	11.2	5
53	Acute COVID-19 severity and mental health morbidity trajectories in patient populations of six nations: an observational study <i>Lancet Public Health, The</i> , 2022 ,	22.4	5
52	Molecular Genetic Risk for Psychosis Is Associated With Psychosis Risk Symptoms in a Population-Based UK Cohort: Findings From Generation Scotland. <i>Schizophrenia Bulletin</i> , 2020 ,	1.3	4
51	Pathology of Tumors Associated With Pathogenic Germline Variants in 9 Breast Cancer Susceptibility Genes <i>JAMA Oncology</i> , 2022 ,	13.4	4
50	Genetic and shared couple environmental contributions to smoking and alcohol use in the UK population. <i>Molecular Psychiatry</i> , 2021 , 26, 4344-4354	15.1	4
49	Pharmaco-epidemiology of antidepressant exposure in a UK cohort record-linkage study. <i>Journal of Psychopharmacology</i> , 2019 , 33, 482-493	4.6	3
48	Gene-educational attainment interactions in a multi-ancestry genome-wide meta-analysis identify novel blood pressure loci. <i>Molecular Psychiatry</i> , 2021 , 26, 2111-2125	15.1	3
47	Using tree-based methods for detection of gene-gene interactions in the presence of a polygenic signal: simulation study with application to educational attainment in the Generation Scotland Cohort Study. <i>Bioinformatics</i> , 2019 , 35, 181-188	7.2	3
46	Using symptom-based case predictions to identify host genetic factors that contribute to COVID-19 susceptibility. <i>PLoS ONE</i> , 2021 , 16, e0255402	3.7	3

45	Genome-Wide Association Study of NAFLD Using Electronic Health Records. <i>Hepatology Communications</i> , 2021 ,	6	3
44	Epigenetic scores for the circulating proteome as tools for disease prediction <i>ELife</i> , 2022 , 11,	8.9	2
43	Cohort Profile: COVIDMENT: COVID-19 cohorts on mental health across six nations <i>International Journal of Epidemiology</i> , 2021 ,	7.8	2
42	TeenCovidLife: The resource to understand the impact of the COVID-19 pandemic on adolescents in Scotland. Wellcome Open Research, 6, 277	4.8	2
41	Creating and validating a DNA methylation-based proxy for Interleukin-6		2
40	Generation Scotland participant survey on data collection. Wellcome Open Research, 2019, 4, 111	4.8	2
39	Electronic Health Record and Genome-wide Genetic Data in Generation Scotland Participants		2
38	Epigenetic scores for the circulating proteome as tools for disease prediction		2
37	A meta-analysis of genome-wide association studies of epigenetic age acceleration		2
36	Blood-based epigenome-wide analyses of cognitive abilities		2
35	Sex Differences in Cardiac Troponin I and T and the Prediction of Cardiovascular Events in the General Population. <i>Clinical Chemistry</i> , 2021 , 67, 1351-1360	5.5	2
34	Meta-GWAS Reveals Novel Genetic Variants Associated with Urinary Excretion of Uromodulin Journal of the American Society of Nephrology: JASN, 2022 , 33, 511-529	12.7	2
33	Blood-based epigenome-wide analyses of cognitive abilities <i>Genome Biology</i> , 2022 , 23, 26	18.3	1
32	Lifestyle and Genetic Factors Modify Parent-of-Origin Effects on the Human Methylome. <i>EBioMedicine</i> , 2021 , 74, 103730	8.8	1
31	Meta-analysis of exome array data identifies six novel genetic loci for lung function. <i>Wellcome Open Research</i> , 3, 4	4.8	1
30	Genome-wide interaction study with major depression identifies novel variants associated with cognitive function. <i>Molecular Psychiatry</i> , 2021 ,	15.1	1
29	Characterisation of an inflammation-related epigenetic score and its association with cognitive ability		1
28	Genetic and shared couple environmental contributions to smoking and alcohol use in the UK population	on	1

27	Epigenome-wide analyses identify DNA methylation signatures of dementia risk. <i>Alzheimerns and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2020 , 12, e12078	5.2	1
26	Socioeconomic position and mental health during the COVID-19 pandemic: a cross-sectional analysis of the CovidLife study. <i>Wellcome Open Research</i> ,6, 139	4.8	1
25	Creating and Validating a DNA Methylation-Based Proxy for Interleukin-6. <i>Journals of Gerontology - Series A Biological Sciences and Medical Sciences</i> , 2021 , 76, 2284-2292	6.4	1
24	Genome-wide methylation data improves dissection of the effect of smoking on body mass index. <i>PLoS Genetics</i> , 2021 , 17, e1009750	6	1
23	Methylome-wide association study of antidepressant use in Generation Scotland and the Netherlands Twin Register implicates the innate immune system. <i>Molecular Psychiatry</i> , 2021 ,	15.1	1
22	Differential and shared genetic effects on kidney function between diabetic and non-diabetic individuals. <i>Communications Biology</i> , 2022 , 5,	6.7	1
21	Association of low-frequency and rare coding variants with information processing speed. <i>Translational Psychiatry</i> , 2021 , 11, 613	8.6	0
20	Epigenome-wide association study of global cortical volumes in generation Scotland: Scottish family health study. <i>Epigenetics</i> , 2021 , 1-17	5.7	O
19	Association of Genetic Variant at Chromosome 12q23.1 With Neuropathic Pain Susceptibility. <i>JAMA Network Open</i> , 2021 , 4, e2136560	10.4	0
18	Hair glucocorticoids are associated with childhood adversity, depressive symptoms and reduced global and lobar grey matter in Generation Scotland. <i>Translational Psychiatry</i> , 2021 , 11, 523	8.6	O
17	Variants associated with HHIP expression have sex-differential effects on lung function. <i>Wellcome Open Research</i> , 2020 , 5, 111	4.8	0
16	Genome-wide association study of susceptibility to hospitalised respiratory infections. <i>Wellcome Open Research</i> ,6, 290	4.8	O
15	Variants associated with expression have sex-differential effects on lung function. <i>Wellcome Open Research</i> , 2020 , 5, 111	4.8	O
14	Cohort profile for the STratifying Resilience and Depression Longitudinally (STRADL) study: A depression-focused investigation of Generation Scotland, using detailed clinical, cognitive, and neuroimaging assessments. <i>Wellcome Open Research</i> ,4, 185	4.8	O
13	Spectral clustering based on structural magnetic resonance imaging and its relationship with major depressive disorder and cognitive ability. <i>European Journal of Neuroscience</i> , 2021 , 54, 6281-6303	3.5	O
12	Identification of plasma proteins relating to brain neurodegeneration and vascular pathology in cognitively normal individuals. <i>Alzheimerns and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2021 , 13, e12240	5.2	O
11	Complex trait methylation scores in the prediction of major depressive disorder <i>EBioMedicine</i> , 2022 , 79, 104000	8.8	O
10	Breast cancer risks associated with missense variants in breast cancer susceptibility genes <i>Genome Medicine</i> , 2022 , 14, 51	14.4	O

LIST OF PUBLICATIONS

9	TeenCovidLife: a resource to understand the impact of the COVID-19 pandemic on adolescents in Scotland. Wellcome Open Research, 6, 277	4.8	О
8	A Computer Based Follow-up of Hypertension in the Community: Baseline Characteristics of 502 Patients. <i>Clinical Science</i> , 1988 , 74, 17P-17P		
7	SNP and Haplotype Regional Heritability Mapping (SNHap-RHM): Joint Mapping of Common and Rare Variation Affecting Complex Traits <i>Frontiers in Genetics</i> , 2021 , 12, 791712	4.5	
6	RuralCovidLife: Study protocol and description of the data. Wellcome Open Research,6, 317	4.8	
5	Generation Scotland participant survey on data collection. Wellcome Open Research, 4, 111	4.8	
4	Face covering adherence is positively associated with better mental health and wellbeing: a longitudinal analysis of the CovidLife surveys. <i>Wellcome Open Research</i> , 6, 62	4.8	
3	General Framework for Meta-Analysis of Haplotype Association Tests. <i>Genetic Epidemiology</i> , 2016 , 40, 244-52	2.6	
2	Genome- and epigenome-wide studies of plasma protein biomarkers for Alzheimers disease implicate TBCA and TREM2 in disease risk <i>Alzheimeris and Dementia: Diagnosis, Assessment and Disease Monitoring</i> , 2022 , 14, e12280	5.2	
1	RuralCovidLife: A new resource for the impact of the pandemic on rural Scotland <i>Wellcome Open Research</i> ,6, 317	4.8	