Hannah R Elliott

List of Publications by Year in Descending Order

 $\textbf{Source:} \ https://exaly.com/author-pdf/3113783/hannah-r-elliott-publications-by-year.pdf$

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.

50
papers

1,854
citations

18
h-index

9-index

53
ext. papers

2,370
ext. citations

7.9
avg, IF

L-index

#	Paper	IF	Citations
50	The EWAS Catalog: a database of epigenome-wide association studies <i>Wellcome Open Research</i> , 2022 , 7, 41	4.8	3
49	Investigating DNA methylation as a potential mediator between pigmentation genes, pigmentary traits and skin cancer. <i>Pigment Cell and Melanoma Research</i> , 2021 , 34, 892-904	4.5	2
48	Comparison of DNA methylation clocks in Black South African men. <i>Epigenomics</i> , 2021 , 13, 437-449	4.4	2
47	Association of medically assisted reproduction with offspring cord blood DNA methylation across cohorts. <i>Human Reproduction</i> , 2021 , 36, 2403-2413	5.7	2
46	Epigenetic age acceleration in the emerging burden of cardiometabolic diseases among migrant and non-migrant African populations: the population based cross-sectional RODAM study <i>The Lancet Healthy Longevity</i> , 2021 , 2, E327-E339	9.5	4
45	DNA methylation of blood cells is associated with prevalent type 2 diabetes in a meta-analysis of four European cohorts. <i>Clinical Epigenetics</i> , 2021 , 13, 40	7.7	8
44	Genomic and phenotypic insights from an atlas of genetic effects on DNA methylation. <i>Nature Genetics</i> , 2021 , 53, 1311-1321	36.3	27
43	Identifying epigenetic biomarkers of established prognostic factors and survival in a clinical cohort of individuals with oropharyngeal cancer. <i>Clinical Epigenetics</i> , 2020 , 12, 95	7.7	4
42	DNA hypomethylation during MSC chondrogenesis occurs predominantly at enhancer regions. <i>Scientific Reports</i> , 2020 , 10, 1169	4.9	9
41	Associations between high blood pressure and DNA methylation. <i>PLoS ONE</i> , 2020 , 15, e0227728	3.7	18
40	Replication and expansion of epigenome-wide association literature in a black South African population. <i>Clinical Epigenetics</i> , 2020 , 12, 6	7.7	9
39	Methylation vs. Protein Inflammatory Biomarkers and Their Associations With Cardiovascular Function. <i>Frontiers in Immunology</i> , 2020 , 11, 1577	8.4	0
38	Leveraging the urban-rural divide for epigenetic research. <i>Epigenomics</i> , 2020 , 12, 1071-1081	4.4	2
37	Associations between high blood pressure and DNA methylation 2020 , 15, e0227728		
36	Associations between high blood pressure and DNA methylation 2020 , 15, e0227728		
35	Associations between high blood pressure and DNA methylation 2020 , 15, e0227728		
34	Associations between high blood pressure and DNA methylation 2020 , 15, e0227728		

33 Associations between high blood pressure and DNA methylation **2020**, 15, e0227728

32	Associations between high blood pressure and DNA methylation 2020 , 15, e0227728		
31	Epigenetics and gestational diabetes: a review of epigenetic epidemiology studies and their use to explore epigenetic mediation and improve prediction. <i>Diabetologia</i> , 2019 , 62, 2171-2178	10.3	26
30	DNA Methylation and Type 2 Diabetes: the Use of Mendelian Randomization to Assess Causality. <i>Current Genetic Medicine Reports</i> , 2019 , 7, 191-207	2.2	5
29	miR-324-5p is up regulated in end-stage osteoarthritis and regulates Indian Hedgehog signalling by differing mechanisms in human and mouse. <i>Matrix Biology</i> , 2019 , 77, 87-100	11.4	27
28	Differential methylation of the type 2 diabetes susceptibility locus KCNQ1 is associated with insulin sensitivity and is predicted by CpG site specific genetic variation. <i>Diabetes Research and Clinical Practice</i> , 2019 , 148, 189-199	7.4	8
27	The Value of Biosamples in Smoking Cessation Trials: A Review of Genetic, Metabolomic, and Epigenetic Findings. <i>Nicotine and Tobacco Research</i> , 2018 , 20, 403-413	4.9	10
26	Leveraging Genomic Data in Smoking Cessation Trials in the Era of Precision Medicine: Why and How. <i>Nicotine and Tobacco Research</i> , 2018 , 20, 414-424	4.9	9
25	Personal Genome Project UK (PGP-UK): a research and citizen science hybrid project in support of personalized medicine. <i>BMC Medical Genomics</i> , 2018 , 11, 108	3.7	20
24	Role of DNA Methylation in Type 2 Diabetes Etiology: Using Genotype as a Causal Anchor. <i>Diabetes</i> , 2017 , 66, 1713-1722	0.9	25
23	Commentary: Migrant study designs for epigenetic studies of disease risk. <i>International Journal of Epidemiology</i> , 2017 , 46, 772	7.8	
22	Epigenome-wide association of DNA methylation markers in peripheral blood from Indian Asians and Europeans with incident type 2 diabetes: a nested case-control study. <i>Lancet Diabetes and Endocrinology,the</i> , 2015 , 3, 526-534	18.1	277
21	Commentary: Migrant study designs for epigenetic studies of disease risk. <i>International Journal of Epidemiology</i> , 2015 ,	7.8	
20	The Effects of Being in a New Relationshiplbn Levels of Testosterone in Men. <i>Evolutionary Psychology</i> , 2015 , 13, 147470491501300	1.5	10
19	Commentary: Migrant study designs for epigenetic studies of disease risk. <i>International Journal of Epidemiology</i> , 2015 , 44, 1449-1451	7.8	2
18	The effects of being in a "new relationship" on levels of testosterone in men. <i>Evolutionary Psychology</i> , 2015 , 13, 250-61	1.5	1
17	Differences in smoking associated DNA methylation patterns in South Asians and Europeans. <i>Clinical Epigenetics</i> , 2014 , 6, 4	7.7	176
16	Valproic acid triggers increased mitochondrial biogenesis in POLG-deficient fibroblasts. <i>Molecular Genetics and Metabolism</i> , 2014 , 112, 57-63	3.7	33

15	Titin founder mutation is a common cause of myofibrillar myopathy with early respiratory failure. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 331-8	:.5	58
14	Competitors who choose to be red have higher testosterone levels. <i>Psychological Science</i> , 2013 , 24, 2122 ₇	· .	19
13	Migration and DNA methylation: a comparison of methylation patterns in type 2 diabetes susceptibility genes between indians and europeans. <i>Journal of Diabetes Research & Clinical Metabolism</i> , 2013 , 2, 6)	4
12	Epigenetics, epidemiology and mitochondrial DNA diseases. <i>International Journal of Epidemiology</i> , 2012 , 41, 177-87	2.8	124
11	The expression and function of microRNAs in chondrogenesis and osteoarthritis. <i>Arthritis and Rheumatism</i> , 2012 , 64, 1909-19		167
10	Titin mutation segregates with hereditary myopathy with early respiratory failure. <i>Brain</i> , 2012 , 135, 1695	-7.1 3	98
9	Epigenetics and child health: basic principles. <i>Archives of Disease in Childhood</i> , 2011 , 96, 863-9	2	38
8	Mitochondrial DNA haplogroups and risk of transient ischaemic attack and ischaemic stroke: a genetic association study. <i>Lancet Neurology, The</i> , 2010 , 9, 498-503	.4.1	74
7	An investigation of mitochondrial haplogroups in autism. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2008 , 147B, 987-9	.5	13
6	Pathogenic mitochondrial DNA mutations are common in the general population. <i>American Journal of Human Genetics</i> , 2008 , 83, 254-60	1	431
5	Episodic ataxia and hemiplegia caused by the 8993T->C mitochondrial DNA mutation. <i>Journal of Medical Genetics</i> , 2007 , 44, 797-9	:.8	24
4	The mitochondrial DNA A3243A>G mutation must be an infrequent cause of Asperger syndrome. <i>Journal of Pediatrics</i> , 2006 , 149, 280-1	.6	6
3	Role of the mitochondrial DNA 16184-16193 poly-C tract in type 2 diabetes. Lancet, The, 2005 , 366, 1650 $_{\uparrow}$. ð	59
2	PGP-UK: a research and citizen science hybrid project in support of personalized medicine		1
1	The EWAS Catalog: a database of epigenome-wide association studies		13