

Keith L Keene

List of Publications by Year in descending order

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Version: 2024-02-01

31
papers

2,693
citations

471371

17
h-index

395590

33
g-index

33
all docs

33
docs citations

33
times ranked

6725
citing authors

#	ARTICLE	IF	CITATIONS
1	Estimating Clinical Research Project Duration from Idea to Publication. <i>Journal of Investigative Medicine</i> , 2022, 70, 108-109.	0.7	3
2	Multi-phenotype analyses of hemostatic traits with cardiovascular events reveal novel genetic associations. <i>Journal of Thrombosis and Haemostasis</i> , 2022, 20, 1331-1349.	1.9	12
3	The Impact of COVID-19 on Racial-Ethnic Health Disparities in the US: Now Is the Time To Address the Problem. <i>Journal of the National Medical Association</i> , 2021, 113, 195-198.	0.6	3
4	Multi-omic analysis of stroke recurrence in African Americans from the Vitamin Intervention for Stroke Prevention (VISP) clinical trial. <i>PLoS ONE</i> , 2021, 16, e0247257.	1.1	4
5	Genetic landscape of Gullah African Americans. <i>American Journal of Physical Anthropology</i> , 2021, 175, 905-919.	2.1	9
6	DNA methylation analyses identify an intronic ZDHHC6 locus associated with time to recurrent stroke in the Vitamin Intervention for Stroke Prevention (VISP) clinical trial. <i>PLoS ONE</i> , 2021, 16, e0254562.	1.1	5
7	Metabolic Traits and Stroke Risk in Individuals of African Ancestry: Mendelian Randomization Analysis. <i>Stroke</i> , 2021, 52, 2680-2684.	1.0	22
8	Genome-Wide Association Study Meta-Analysis of Stroke in 22 000 Individuals of African Descent Identifies Novel Associations With Stroke. <i>Stroke</i> , 2020, 51, 2454-2463.	1.0	26
9	Differential expression of PHACTR1 in atheromatous versus normal carotid artery tissue. <i>Journal of Clinical Neuroscience</i> , 2020, 74, 265-267.	0.8	3
10	Cervical Artery Dissection in Patients of African Ancestry. <i>Cerebrovascular Diseases</i> , 2018, 46, 218-222.	0.8	3
11	Epigenome-Wide Analyses Identify Two Novel Associations With Recurrent Stroke in the Vitamin Intervention for Stroke Prevention Clinical Trial. <i>Frontiers in Genetics</i> , 2018, 9, 358.	1.1	12
12	Multiancestry genome-wide association study of 520,000 subjects identifies 32 loci associated with stroke and stroke subtypes. <i>Nature Genetics</i> , 2018, 50, 524-537.	9.4	1,124
13	Genetic Drivers of von Willebrand Factor Levels in an Ischemic Stroke Population and Association With Risk for Recurrent Stroke. <i>Stroke</i> , 2017, 48, 1444-1450.	1.0	21
14	Type 2 Diabetes Variants Disrupt Function of SLC16A11 through Two Distinct Mechanisms. <i>Cell</i> , 2017, 170, 199-212.e20.	13.5	121
15	Loci associated with ischaemic stroke and its subtypes (SiGN): a genome-wide association study. <i>Lancet Neurology</i> , The, 2016, 15, 174-184.	4.9	217
16	Genetic associations at 53 loci highlight cell types and biological pathways relevant for kidney function. <i>Nature Communications</i> , 2016, 7, 10023.	5.8	412
17	Shared genetic susceptibility of vascular-related biomarkers with ischemic and recurrent stroke. <i>Neurology</i> , 2016, 86, 351-359.	1.5	33
18	Meta-Analysis of Genome-Wide Association Studies Identifies Genetic Risk Factors for Stroke in African Americans. <i>Stroke</i> , 2015, 46, 2063-2068.	1.0	63

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19	Genetic Associations with Plasma B12, B6, and Folate Levels in an Ischemic Stroke Population from the Vitamin Intervention for Stroke Prevention (VISP) Trial. <i>Frontiers in Public Health</i> , 2014, 2, 112.	1.3	23
20	Genome-Wide Association Study for Circulating Tissue Plasminogen Activator Levels and Functional Follow-Up Implicates Endothelial <i>STXBP5</i> and <i>STX2</i> . <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2014, 34, 1093-1101.	1.1	43
21	Genome-Wide Meta-Analysis of Homocysteine and Methionine Metabolism Identifies Five One Carbon Metabolism Loci and a Novel Association of <i>ALDH1L1</i> with Ischemic Stroke. <i>PLoS Genetics</i> , 2014, 10, e1004214.	1.5	69
22	Meta-Analysis of Genome-Wide Association Studies in African Americans Provides Insights into the Genetic Architecture of Type 2 Diabetes. <i>PLoS Genetics</i> , 2014, 10, e1004517.	1.5	191
23	Fine Mapping and Functional Studies of Risk Variants for Type 1 Diabetes at Chromosome 16p13.13. <i>Diabetes</i> , 2014, 63, 4360-4368.	0.3	17
24	Chromosome 7p linkage and association study for diabetes related traits and type 2 diabetes in an African-American population enriched for nephropathy. <i>BMC Medical Genetics</i> , 2010, 11, 22.	2.1	13
25	Comprehensive evaluation of the estrogen receptor β gene reveals further evidence for association with type 2 diabetes enriched for nephropathy in an African American population. <i>Human Genetics</i> , 2008, 123, 333-341.	1.8	28
26	Evaluation of a SNP map of 6q24 confirms diabetic nephropathy loci and identifies novel associations in type 2 diabetes patients with nephropathy from an African-American population. <i>Human Genetics</i> , 2008, 124, 63-71.	1.8	14
27	Exploration of the utility of ancestry informative markers for genetic association studies of African Americans with type 2 diabetes and end stage renal disease. <i>Human Genetics</i> , 2008, 124, 147-154.	1.8	29
28	Association of the Distal Region of the Ectonucleotide Pyrophosphatase/Phosphodiesterase 1 Gene With Type 2 Diabetes in an African-American Population Enriched for Nephropathy. <i>Diabetes</i> , 2008, 57, 1057-1062.	0.3	28
29	Association Analysis of the Ephrin-B2 Gene in African-Americans with End-Stage Renal Disease. <i>American Journal of Nephrology</i> , 2008, 28, 914-920.	1.4	7
30	Variants of the Transcription Factor 7-Like 2 (<i>TCF7L2</i>) Gene Are Associated With Type 2 Diabetes in an African-American Population Enriched for Nephropathy. <i>Diabetes</i> , 2007, 56, 2638-2642.	0.3	89
31	Investigation of the Estrogen Receptor- β Gene With Type 2 Diabetes and/or Nephropathy in African-American and European-American Populations. <i>Diabetes</i> , 2007, 56, 675-684.	0.3	30