

Kiran Musunuru

List of Publications by Year in Descending Order

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

118
papers

14,109
citations

42
h-index

118
g-index

133
ext. papers

17,093
ext. citations

13.6
avg, IF

6.27
L-index

#	Paper	IF	Citations
118	Adipocyte-Specific Modulation of KLF14 Expression in Mice Leads to Sex-Dependent Impacts on Adiposity and Lipid Metabolism.. <i>Diabetes</i> , 2022 ,	0.9	1
117	Engineered virus-like particles for efficient in vivo delivery of therapeutic proteins.. <i>Cell</i> , 2022 , 185, 250-265.e166	36.5	166
116	Moving toward genome-editing therapies for cardiovascular diseases.. <i>Journal of Clinical Investigation</i> , 2022 , 132,	15.9	2
115	Mutations and Cardiovascular Function in Breast Cancer Survivors.. <i>Frontiers in Cardiovascular Medicine</i> , 2022 , 9, 833171	5.4	0
114	Detoxifying chemotherapy with genetics-guided stem cell modeling: A personalized affair. <i>Cell Stem Cell</i> , 2021 , 28, 2039-2040	18	
113	Active learning-based STEM education for in-person and online learning. <i>Cell</i> , 2021 , 184, 1409-1414	56.2	4
112	Chronobiology of Natriuretic Peptides and Blood Pressure in Lean and Obese Individuals. <i>Journal of the American College of Cardiology</i> , 2021 , 77, 2291-2303	15.1	4
111	In vivo CRISPR base editing of PCSK9 durably lowers cholesterol in primates. <i>Nature</i> , 2021 , 593, 429-434	50.4	96
110	Pathogenic LMNA variants disrupt cardiac lamina-chromatin interactions and de-repress alternative fate genes. <i>Cell Stem Cell</i> , 2021 , 28, 938-954.e9	18	23
109	Treating Coronary Artery Disease: Beyond Statins, Ezetimibe, and PCSK9 Inhibition. <i>Annual Review of Medicine</i> , 2021 , 72, 447-458	17.4	3
108	Genome editing for cellular disease modeling 2021 , 145-167		
107	Assessing for off-target mutagenesis 2021 , 81-100		0
106	Base editing 2021 , 101-121		
105	Therapeutic genome editing 2021 , 193-211		
104	In utero adenine base editing corrects multi-organ pathology in a lethal lysosomal storage disease. <i>Nature Communications</i> , 2021 , 12, 4291	17.4	5
103	CRISPR Hits Home in a First-in-Human Study. <i>CRISPR Journal</i> , 2021 , 4, 460-461	2.5	0
102	Genome editing for functional experiments and screens 2021 , 169-191		

101	Therapeutic application of genome editing in dyslipidemia.. <i>Current Opinion in Lipidology</i> , 2021 , 33,	4.4	1
100	EDEM3 Modulates Plasma Triglyceride Level through Its Regulation of LRP1 Expression. <i>IScience</i> , 2020 , 23, 100973	6.1	4
99	Noncoding RNAs in Cardiovascular Disease: Current Knowledge, Tools and Technologies for Investigation, and Future Directions: A Scientific Statement From the American Heart Association. <i>Circulation Genomic and Precision Medicine</i> , 2020 , 13, e000062	5.2	18
98	Correcting tyrosinaemia via a point mutation. <i>Nature Biomedical Engineering</i> , 2020 , 4, 14-15	19	
97	Genetic Testing for Inherited Cardiovascular Diseases: A Scientific Statement From the American Heart Association. <i>Circulation Genomic and Precision Medicine</i> , 2020 , 13, e000067	5.2	59
96	Self-Organizing Human Induced Pluripotent Stem Cell Hepatocyte 3D Organoids Inform the Biology of the Pleiotropic Gene. <i>Hepatology Communications</i> , 2020 , 4, 1316-1331	6	5
95	Base editing: a brief review and a practical example. <i>Journal of Biomedical Research</i> , 2020 , 35, 107-114	1.5	
94	Challenges and advances of CRISPR-Cas9 genome editing in therapeutics. <i>Cardiovascular Research</i> , 2019 , 115, e12-e14	9.9	3
93	From Hypertrophy to Heart Failure: What Is New in Genetic Cardiomyopathies. <i>Current Heart Failure Reports</i> , 2019 , 16, 157-167	2.8	7
92	Cardioprotective Effects of MTSS1 Enhancer Variants. <i>Circulation</i> , 2019 , 139, 2073-2076	16.7	2
91	In utero gene editing for monogenic lung disease. <i>Science Translational Medicine</i> , 2019 , 11,	17.5	51
90	Genetics of Common, Complex Coronary Artery Disease. <i>Cell</i> , 2019 , 177, 132-145	56.2	84
89	The clinical application of gene editing: ethical and social issues. <i>Personalized Medicine</i> , 2019 , 16, 337-350.	2	15
88	Race, Natriuretic Peptides, and High-Carbohydrate Challenge: A Clinical Trial. <i>Circulation Research</i> , 2019 , 125, 957-968	15.7	21
87	Angiotensin-Like 3: From Discovery to Therapeutic Gene Editing. <i>JACC Basic To Translational Science</i> , 2019 , 4, 755-762	8.7	17
86	Investigation of a dilated cardiomyopathy-associated variant in BAG3 using genome-edited iPSC-derived cardiomyocytes. <i>JCI Insight</i> , 2019 , 4,	9.9	20
85	Why Human Embryo Editing Should Be Banned. <i>CRISPR Journal</i> , 2019 , 2, 356-358	2.5	2
84	Reduced Blood Lipid Levels With In Vivo CRISPR-Cas9 Base Editing of ANGPTL3. <i>Circulation</i> , 2018 , 137, 975-977	16.7	68

83	Functional Assays to Screen and Dissect Genomic Hits: Doubling Down on the National Investment in Genomic Research. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e002178	5.2	16
82	Regulatory variants at KLF14 influence type 2 diabetes risk via a female-specific effect on adipocyte size and body composition. <i>Nature Genetics</i> , 2018 , 50, 572-580	36.3	82
81	From Genotype to Phenotype. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11,	5.2	11
80	Induced Pluripotent Stem Cells for Cardiovascular Disease Modeling and Precision Medicine: A Scientific Statement From the American Heart Association. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e000043	5.2	95
79	Stem cell modeling of lipid genetics. <i>Current Opinion in Lipidology</i> , 2018 , 29, 151-155	4.4	1
78	Turning up the Heat with Therapeutic Epigenome Editing. <i>Cell Stem Cell</i> , 2018 , 22, 10-11	18	8
77	Role of angiopoietin-like 3 (ANGPTL3) in regulating plasma level of low-density lipoprotein cholesterol. <i>Atherosclerosis</i> , 2018 , 268, 196-206	3.1	52
76	CRISPR-Cas9 Genome Editing for Treatment of Atherogenic Dyslipidemia. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2018 , 38, 12-18	9.4	13
75	Genetic and Chemical Screenings Identify HDAC3 as a Key Regulator in Hepatic Differentiation of Human Pluripotent Stem Cells. <i>Stem Cell Reports</i> , 2018 , 11, 22-31	8	20
74	From Genotype to Phenotype: A Primer on the Functional Follow-up of Genome-Wide Association Studies in Cardiovascular Disease. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11,	5.2	4
73	Interrogation of the Atherosclerosis-Associated (Sortilin 1) Locus With Primary Human Hepatocytes, Induced Pluripotent Stem Cell-Hepatocytes, and Locus-Humanized Mice. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2018 , 38, 76-82	9.4	23
72	Functional Annotation of TNNT2 Variants of Uncertain Significance With Genome-Edited Cardiomyocytes. <i>Circulation</i> , 2018 , 138, 2852-2854	16.7	23
71	In utero CRISPR-mediated therapeutic editing of metabolic genes. <i>Nature Medicine</i> , 2018 , 24, 1513-1518	50.5	112
70	Interdisciplinary Models for Research and Clinical Endeavors in Genomic Medicine: A Scientific Statement From the American Heart Association. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e000046	5.2	7
69	Confirmation of Causal rs9349379- PHACTR1 Expression Quantitative Trait Locus in Human-Induced Pluripotent Stem Cell Endothelial Cells. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e002327	5.2	19
68	Novel tricyclic glycol-based inducers that reprogram LDL metabolism in hepatic cells. <i>MedChemComm</i> , 2018 , 9, 1831-1842	5	3
67	Genome Editing for the Study of Cardiovascular Diseases. <i>Current Cardiology Reports</i> , 2017 , 19, 22	4.2	17
66	Treatment of Dyslipidemia Using CRISPR/Cas9 Genome Editing. <i>Current Atherosclerosis Reports</i> , 2017 , 19, 32	6	6

65	The Hope and Hype of CRISPR-Cas9 Genome Editing: A Review. <i>JAMA Cardiology</i> , 2017 , 2, 914-919	16.2	30
64	NLRP2 is a suppressor of NF- κ B signaling and HLA-C expression in human trophoblasts. <i>Biology of Reproduction</i> , 2017 , 96, 831-842	3.9	31
63	ANGPTL3 Deficiency and Protection Against Coronary Artery Disease. <i>Journal of the American College of Cardiology</i> , 2017 , 69, 2054-2063	15.1	226
62	Large, Diverse Population Cohorts of hiPSCs and Derived Hepatocyte-like Cells Reveal Functional Genetic Variation at Blood Lipid-Associated Loci. <i>Cell Stem Cell</i> , 2017 , 20, 558-570.e10	18	102
61	Induced Pluripotent Stem Cell Differentiation Enables Functional Validation of GWAS Variants in Metabolic Disease. <i>Cell Stem Cell</i> , 2017 , 20, 547-557.e7	18	86
60	ESRP1 Mutations Cause Hearing Loss due to Defects in Alternative Splicing that Disrupt Cochlear Development. <i>Developmental Cell</i> , 2017 , 43, 318-331.e5	10.2	26
59	Exome-wide association study of plasma lipids in >300,000 individuals. <i>Nature Genetics</i> , 2017 , 49, 1758-1766	13.6	310
58	In Vivo Base Editing of PCSK9 (Proprotein Convertase Subtilisin/Kexin Type 9) as a Therapeutic Alternative to Genome Editing. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2017 , 37, 1741-1747	9.4	128
57	Human Germline Genome Editing. <i>American Journal of Human Genetics</i> , 2017 , 101, 167-176	11	105
56	Genome Editing: The Recent History and Perspective in Cardiovascular Diseases. <i>Journal of the American College of Cardiology</i> , 2017 , 70, 2808-2821	15.1	14
55	Cardiovascular endocrinology: Is ANGPTL3 the next PCSK9?. <i>Nature Reviews Endocrinology</i> , 2017 , 13, 503-504	15.2	14
54	Genome editing in cardiovascular diseases. <i>Nature Reviews Cardiology</i> , 2017 , 14, 11-20	14.8	57
53	Targeted Application of Human Genetic Variation Can Improve Red Blood Cell Production from Stem Cells. <i>Cell Stem Cell</i> , 2016 , 18, 73-78	18	57
52	Circular non-coding RNA ANRIL modulates ribosomal RNA maturation and atherosclerosis in humans. <i>Nature Communications</i> , 2016 , 7, 12429	17.4	619
51	Genome engineering tools for building cellular models of disease. <i>FEBS Journal</i> , 2016 , 283, 3222-31	5.7	21
50	Transcriptome-Wide Analysis Reveals Modulation of Human Macrophage Inflammatory Phenotype Through Alternative Splicing. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2016 , 36, 1434-47	9.4	30
49	Genome-Edited Human Pluripotent Stem Cell-Derived Macrophages as a Model of Reverse Cholesterol Transport--Brief Report. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2016 , 36, 15-8	9.4	27
48	Surprises From Genetic Analyses of Lipid Risk Factors for Atherosclerosis. <i>Circulation Research</i> , 2016 , 118, 579-85	15.7	98

47	A dual AAV system enables the Cas9-mediated correction of a metabolic liver disease in newborn mice. <i>Nature Biotechnology</i> , 2016 , 34, 334-8	44.5	360
46	CRISPR-Cas9 Targeting of PCSK9 in Human Hepatocytes In Vivo-Brief Report. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2016 , 36, 783-6	9.4	85
45	Rapid generation of novel models of RAG1 deficiency by CRISPR/Cas9-induced mutagenesis in murine zygotes. <i>Oncotarget</i> , 2016 , 7, 12962-74	3.3	8
44	Asialoglycoprotein receptor 1 is a specific cell-surface marker for isolating hepatocytes derived from human pluripotent stem cells. <i>Development (Cambridge)</i> , 2016 , 143, 1475-81	6.6	36
43	Enhancing Literacy in Cardiovascular Genetics: A Scientific Statement From the American Heart Association. <i>Circulation: Cardiovascular Genetics</i> , 2016 , 9, 448-467		37
42	Improving Genomic Literacy Among Cardiovascular Practitioners via a Flipped-Classroom Workshop at a National Meeting. <i>Circulation: Cardiovascular Genetics</i> , 2016 , 9, 287-90		7
41	Myocardial Infarction-Associated SNP at 6p24 Interferes With MEF2 Binding and Associates With PHACTR1 Expression Levels in Human Coronary Arteries. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2015 , 35, 1472-1479	9.4	62
40	Modelling kidney disease with CRISPR-mutant kidney organoids derived from human pluripotent epiblast spheroids. <i>Nature Communications</i> , 2015 , 6, 8715	17.4	410
39	Novel Genome-Editing Tools to Model and Correct Primary Immunodeficiencies. <i>Frontiers in Immunology</i> , 2015 , 6, 250	8.4	27
38	Modulators of hepatic lipoprotein metabolism identified in a search for small-molecule inducers of tribbles pseudokinase 1 expression. <i>PLoS ONE</i> , 2015 , 10, e0120295	3.7	21
37	Basic concepts and potential applications of genetics and genomics for cardiovascular and stroke clinicians: a scientific statement from the American Heart Association. <i>Circulation: Cardiovascular Genetics</i> , 2015 , 8, 216-42		33
36	Personalized genomes and cardiovascular disease. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2014 , 5, a014068	5.4	5
35	Efficient ablation of genes in human hematopoietic stem and effector cells using CRISPR/Cas9. <i>Cell Stem Cell</i> , 2014 , 15, 643-52	18	324
34	Low incidence of off-target mutations in individual CRISPR-Cas9 and TALEN targeted human stem cell clones detected by whole-genome sequencing. <i>Cell Stem Cell</i> , 2014 , 15, 27-30	18	394
33	Permanent alteration of PCSK9 with in vivo CRISPR-Cas9 genome editing. <i>Circulation Research</i> , 2014 , 115, 488-92	15.7	345
32	Expanding the genetic editing tool kit: ZFNs, TALENs, and CRISPR-Cas9. <i>Journal of Clinical Investigation</i> , 2014 , 124, 4154-61	15.9	252
31	A TALEN genome-editing system for generating human stem cell-based disease models. <i>Cell Stem Cell</i> , 2013 , 12, 238-51	18	407
30	Enhanced efficiency of human pluripotent stem cell genome editing through replacing TALENs with CRISPRs. <i>Cell Stem Cell</i> , 2013 , 12, 393-4	18	391

29	Genome editing of human pluripotent stem cells to generate human cellular disease models. <i>DMM Disease Models and Mechanisms</i> , 2013 , 6, 896-904	4.1	94
28	Identification of a growth factor that rejuvenates the heart. <i>Circulation: Cardiovascular Genetics</i> , 2013 , 6, 435-6		
27	Enduring mystery of the chromosome 9p21.3 locus. <i>Circulation: Cardiovascular Genetics</i> , 2013 , 6, 224-5		6
26	Plasma HDL cholesterol and risk of myocardial infarction: a mendelian randomisation study. <i>Lancet, The</i> , 2012 , 380, 572-80	40	1523
25	Transforming growth factor β mutations and familial thoracic aortic aneurysms. <i>Circulation: Cardiovascular Genetics</i> , 2012 , 5, 593-4		1
24	Induction of cardiomyocytes from cardiac fibroblasts. <i>Circulation: Cardiovascular Genetics</i> , 2012 , 5, 481-2		
23	Exome sequencing to identify novel genes in hypertension. <i>Circulation: Cardiovascular Genetics</i> , 2012 , 5, 267-8		
22	Functional evaluation of genetic variation in complex human traits. <i>Human Molecular Genetics</i> , 2012 , 21, R18-23	5.6	9
21	Cardiovascular pharmacogenomics: current status and future directions-report of a national heart, lung, and blood institute working group. <i>Journal of the American Heart Association</i> , 2012 , 1, e000554	6	9
20	Hepatic sortilin regulates both apolipoprotein B secretion and LDL catabolism. <i>Journal of Clinical Investigation</i> , 2012 , 122, 2807-16	15.9	151
19	Large-scale association analysis identifies 13 new susceptibility loci for coronary artery disease. <i>Nature Genetics</i> , 2011 , 43, 333-8	36.3	1394
18	Current role of pharmacogenomics in cardiovascular medicine. <i>Current Treatment Options in Cardiovascular Medicine</i> , 2011 , 13, 302-12	2.1	1
17	Regulatory elements in noncoding DNA in the chromosome 9p21 locus. <i>Circulation: Cardiovascular Genetics</i> , 2011 , 4, 330-1		3
16	Lack of association of KIF6 genotype with vascular disease and statin response. <i>Circulation: Cardiovascular Genetics</i> , 2011 , 4, 467-8		2
15	From noncoding variant to phenotype via SORT1 at the 1p13 cholesterol locus. <i>Nature</i> , 2010 , 466, 714-9	50.4	820
14	Biological, clinical and population relevance of 95 loci for blood lipids. <i>Nature</i> , 2010 , 466, 707-13	50.4	2742
13	Association of single nucleotide polymorphisms on chromosome 9p21.3 with platelet reactivity: a potential mechanism for increased vascular disease. <i>Circulation: Cardiovascular Genetics</i> , 2010 , 3, 445-53		48
12	Stem cell models of cardiac development and disease. <i>Annual Review of Cell and Developmental Biology</i> , 2010 , 26, 667-87	12.6	53

11	Exome sequencing, ANGPTL3 mutations, and familial combined hypolipidemia. <i>New England Journal of Medicine</i> , 2010 , 363, 2220-7	59.2	485
10	Genetics of coronary artery disease. <i>Annual Review of Genomics and Human Genetics</i> , 2010 , 11, 91-108	9.7	67
9	Atherogenic dyslipidemia: cardiovascular risk and dietary intervention. <i>Lipids</i> , 2010 , 45, 907-14	1.6	182
8	Ion mobility analysis of lipoprotein subfractions identifies three independent axes of cardiovascular risk. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2009 , 29, 1975-80	9.4	113
7	Paraneoplastic opsoclonus-myoclonus ataxia associated with non-small-cell lung carcinoma. <i>Journal of Neuro-Oncology</i> , 2008 , 90, 213-6	4.8	22
6	A synergistic relationship of elevated low-density lipoprotein cholesterol levels and systolic blood pressure with coronary artery calcification. <i>Atherosclerosis</i> , 2008 , 200, 368-73	3.1	0
5	The use of high-sensitivity assays for C-reactive protein in clinical practice. <i>Nature Clinical Practice Cardiovascular Medicine</i> , 2008 , 5, 621-35		95
4	HapMap and mapping genes for cardiovascular disease. <i>Circulation: Cardiovascular Genetics</i> , 2008 , 1, 66-71		12
3	Lipid nanoparticles incorporating a GalNAc ligand enable in vivo liver ANGPTL3 editing in wild-type and somatic LDLR knockout non-human primates		1
2	High-Throughput Screening and CRISPR-Cas9 Modeling of Causal Lipid-Associated Expression Quantitative Trait Locus Variants		3
1	Global-scale CRISPR gene editor specificity profiling by ONE-seq identifies population-specific, variant off-target effects		2