

Sandosh Padmanabhan

List of Publications by Citations

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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.

219
papers

14,906
citations

68
h-index

118
g-index

235
ext. papers

19,126
ext. citations

10.5
avg, IF

5.72
L-index

#	Paper	IF	Citations
219	Spironolactone versus placebo, bisoprolol, and doxazosin to determine the optimal treatment for drug-resistant hypertension (PATHWAY-2): a randomised, double-blind, crossover trial. <i>Lancet, The</i> , 2015 , 386, 2059-2068	40	632
218	Genetically distinct subsets within ANCA-associated vasculitis. <i>New England Journal of Medicine</i> , 2012 , 367, 214-23	59.2	627
217	Mendelian randomization of blood lipids for coronary heart disease. <i>European Heart Journal</i> , 2015 , 36, 539-50	9.5	417
216	Rare and low-frequency coding variants alter human adult height. <i>Nature</i> , 2017 , 542, 186-190	50.4	412
215	Association between alcohol and cardiovascular disease: Mendelian randomisation analysis based on individual participant data. <i>BMJ, The</i> , 2014 , 349, g4164	5.9	406
214	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
213	Risk HLA-DQA1 and PLA(2)R1 alleles in idiopathic membranous nephropathy. <i>New England Journal of Medicine</i> , 2011 , 364, 616-26	59.2	350
212	Genome-wide association analysis identifies novel blood pressure loci and offers biological insights into cardiovascular risk. <i>Nature Genetics</i> , 2017 , 49, 403-415	36.3	313
211	Exome-wide association study of plasma lipids in >300,000 individuals. <i>Nature Genetics</i> , 2017 , 49, 1758-1766	36.3	310
210	Multi-ethnic genome-wide association study for atrial fibrillation. <i>Nature Genetics</i> , 2018 , 50, 1225-1233	36.3	277
209	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
208	Seventy-five genetic loci influencing the human red blood cell. <i>Nature</i> , 2012 , 492, 369-75	50.4	257
207	Genome-wide association study of blood pressure extremes identifies variant near UMOD associated with hypertension. <i>PLoS Genetics</i> , 2010 , 6, e1001177	6	255
206	Common variants in 22 loci are associated with QRS duration and cardiac ventricular conduction. <i>Nature Genetics</i> , 2010 , 42, 1068-76	36.3	249
205	Polygenic Risk Score Identifies Subgroup With Higher Burden of Atherosclerosis and Greater Relative Benefit From Statin Therapy in the Primary Prevention Setting. <i>Circulation</i> , 2017 , 135, 2091-2101	16.7	244
204	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
203	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

202	Identification of heart rate-associated loci and their effects on cardiac conduction and rhythm disorders. <i>Nature Genetics</i> , 2013 , 45, 621-31	36.3	219
201	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
200	Large-scale gene-centric meta-analysis across 39 studies identifies type 2 diabetes loci. <i>American Journal of Human Genetics</i> , 2012 , 90, 410-25	11	214
199	Genetic association study of QT interval highlights role for calcium signaling pathways in myocardial repolarization. <i>Nature Genetics</i> , 2014 , 46, 826-36	36.3	199
198	Large-scale gene-centric meta-analysis across 32 studies identifies multiple lipid loci. <i>American Journal of Human Genetics</i> , 2012 , 91, 823-38	11	189
197	Protein-altering variants associated with body mass index implicate pathways that control energy intake and expenditure in obesity. <i>Nature Genetics</i> , 2018 , 50, 26-41	36.3	186
196	Trans-ancestry meta-analyses identify rare and common variants associated with blood pressure and hypertension. <i>Nature Genetics</i> , 2016 , 48, 1151-1161	36.3	181
195	Association between genetic variants on chromosome 15q25 locus and objective measures of tobacco exposure. <i>Journal of the National Cancer Institute</i> , 2012 , 104, 740-8	9.7	178
194	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
193	HLA has strongest association with IgA nephropathy in genome-wide analysis. <i>Journal of the American Society of Nephrology: JASN</i> , 2010 , 21, 1791-7	12.7	173
192	Obesity paradox in a cohort of 4880 consecutive patients undergoing percutaneous coronary intervention. <i>European Heart Journal</i> , 2010 , 31, 222-6	9.5	173
191	Whole-exome sequencing identifies rare and low-frequency coding variants associated with LDL cholesterol. <i>American Journal of Human Genetics</i> , 2014 , 94, 233-45	11	170
190	Genetic and molecular aspects of hypertension. <i>Circulation Research</i> , 2015 , 116, 937-59	15.7	165
189	Gene-centric association signals for lipids and apolipoproteins identified via the HumanCVD BeadChip. <i>American Journal of Human Genetics</i> , 2009 , 85, 628-42	11	163
188	Copy-number disorders are a common cause of congenital kidney malformations. <i>American Journal of Human Genetics</i> , 2012 , 91, 987-97	11	161
187	Low-frequency and rare exome chip variants associate with fasting glucose and type 2 diabetes susceptibility. <i>Nature Communications</i> , 2015 , 6, 5897	17.4	147
186	Should diabetes be considered a coronary heart disease risk equivalent?: results from 25 years of follow-up in the Renfrew and Paisley survey. <i>Diabetes Care</i> , 2005 , 28, 1588-93	14.6	147
185	Blood pressure loci identified with a gene-centric array. <i>American Journal of Human Genetics</i> , 2011 , 89, 688-700	11	137

184	Molecular genetic contributions to socioeconomic status and intelligence. <i>Intelligence</i> , 2014 , 44, 26-32	3	131
183	Gene-centric meta-analysis in 87,736 individuals of European ancestry identifies multiple blood-pressure-related loci. <i>American Journal of Human Genetics</i> , 2014 , 94, 349-60	11	131
182	Genome-wide Association for Major Depression Through Age at Onset Stratification: Major Depressive Disorder Working Group of the Psychiatric Genomics Consortium. <i>Biological Psychiatry</i> , 2017 , 81, 325-335	7.9	129
181	Endocrine and haemodynamic changes in resistant hypertension, and blood pressure responses to spironolactone or amiloride: the PATHWAY-2 mechanisms substudies. <i>Lancet Diabetes and Endocrinology</i> , 2018 , 6, 464-475	18.1	126
180	Genomewide association study using a high-density single nucleotide polymorphism array and case-control design identifies a novel essential hypertension susceptibility locus in the promoter region of endothelial NO synthase. <i>Hypertension</i> , 2012 , 59, 248-55	8.5	124
179	Target genes, variants, tissues and transcriptional pathways influencing human serum urate levels. <i>Nature Genetics</i> , 2019 , 51, 1459-1474	36.3	122
178	Directional dominance on stature and cognition in diverse human populations. <i>Nature</i> , 2015 , 523, 459-462	50.4	119
177	Loci influencing blood pressure identified using a cardiovascular gene-centric array. <i>Human Molecular Genetics</i> , 2013 , 22, 1663-78	5.6	119
176	Variants in the fetal genome near FLT1 are associated with risk of preeclampsia. <i>Nature Genetics</i> , 2017 , 49, 1255-1260	36.3	118
175	Genome-wide meta-analysis of 241,258 adults accounting for smoking behaviour identifies novel loci for obesity traits. <i>Nature Communications</i> , 2017 , 8, 14977	17.4	105
174	Meta-analysis of Dense Genecentric Association Studies Reveals Common and Uncommon Variants Associated with Height. <i>American Journal of Human Genetics</i> , 2011 , 88, 6-18	11	103
173	Genome-wide physical activity interactions in adiposity - A meta-analysis of 200,452 adults. <i>PLoS Genetics</i> , 2017 , 13, e1006528	6	103
172	Resting heart rate pattern during follow-up and mortality in hypertensive patients. <i>Hypertension</i> , 2010 , 55, 567-74	8.5	101
171	Meta-analysis of genome-wide association studies identifies six new Loci for serum calcium concentrations. <i>PLoS Genetics</i> , 2013 , 9, e1003796	6	100
170	Genetic basis of blood pressure and hypertension. <i>Trends in Genetics</i> , 2012 , 28, 397-408	8.5	99
169	Genetic variation at CHRNA5-CHRNA3-CHRNA4 interacts with smoking status to influence body mass index. <i>International Journal of Epidemiology</i> , 2011 , 40, 1617-28	7.8	92
168	Cardiac Troponin T and Troponin I in the General Population. <i>Circulation</i> , 2019 , 139, 2754-2764	16.7	90
167	Systems genetics identifies a convergent gene network for cognition and neurodevelopmental disease. <i>Nature Neuroscience</i> , 2016 , 19, 223-32	25.5	88

166	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
165	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
164	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
163	Genetic variation at the SLC23A1 locus is associated with circulating concentrations of L-ascorbic acid (vitamin C): evidence from 5 independent studies with >15,000 participants. <i>American Journal of Clinical Nutrition</i> , 2010 , 92, 375-82	7	84
162	Validation of uromodulin as a candidate gene for human essential hypertension. <i>Hypertension</i> , 2014 , 63, 551-8	8.5	83
161	Large-scale GWAS identifies multiple loci for hand grip strength providing biological insights into muscular fitness. <i>Nature Communications</i> , 2017 , 8, 16015	17.4	80
160	Sixteen new lung function signals identified through 1000 Genomes Project reference panel imputation. <i>Nature Communications</i> , 2015 , 6, 8658	17.4	79
159	Genomic association analysis of common variants influencing antihypertensive response to hydrochlorothiazide. <i>Hypertension</i> , 2013 , 62, 391-7	8.5	79
158	Meta-analysis of Dense Genecentric Association Studies Reveals Common and Uncommon Variants Associated with Height. <i>American Journal of Human Genetics</i> , 2012 , 90, 1116-1117	11	78
157	52 Genetic Loci Influencing Myocardial Mass. <i>Journal of the American College of Cardiology</i> , 2016 , 68, 1435-1448	15.1	76
156	No Evidence of a Common DNA Variant Profile Specific to World Class Endurance Athletes. <i>PLoS ONE</i> , 2016 , 11, e0147330	3.7	74
155	Allopurinol and Cardiovascular Outcomes in Adults With Hypertension. <i>Hypertension</i> , 2016 , 67, 535-40	8.5	72
154	Effect of amiloride, or amiloride plus hydrochlorothiazide, versus hydrochlorothiazide on glucose tolerance and blood pressure (PATHWAY-3): a parallel-group, double-blind randomised phase 4 trial. <i>Lancet Diabetes and Endocrinology</i> , 2016 , 4, 136-47	18.1	72
153	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
152	The Y chromosome effect on blood pressure in two European populations. <i>Hypertension</i> , 2002 , 39, 353-68.5		70
151	Long-term and ultra long-term blood pressure variability during follow-up and mortality in 14,522 patients with hypertension. <i>Hypertension</i> , 2013 , 62, 698-705	8.5	68
150	Monotherapy With Major Antihypertensive Drug Classes and Risk of Hospital Admissions for Mood Disorders. <i>Hypertension</i> , 2016 , 68, 1132-1138	8.5	68
149	Adult height, coronary heart disease and stroke: a multi-locus Mendelian randomization meta-analysis. <i>International Journal of Epidemiology</i> , 2016 , 45, 1927-1937	7.8	65

148	Metabolomic identification of a novel pathway of blood pressure regulation involving hexadecanedioate. <i>Hypertension</i> , 2015 , 66, 422-9	8.5	63
147	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
146	Towards Precision Medicine for Hypertension: A Review of Genomic, Epigenomic, and Microbiomic Effects on Blood Pressure in Experimental Rat Models and Humans. <i>Physiological Reviews</i> , 2017 , 97, 1469-1528	17.9	60
145	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
144	Combination Therapy Is Superior to Sequential Monotherapy for the Initial Treatment of Hypertension: A Double-Blind Randomized Controlled Trial. <i>Journal of the American Heart Association</i> , 2017 , 6,	6	57
143	Serum chloride is an independent predictor of mortality in hypertensive patients. <i>Hypertension</i> , 2013 , 62, 836-43	8.5	51
142	Implications of discoveries from genome-wide association studies in current cardiovascular practice. <i>World Journal of Cardiology</i> , 2011 , 3, 230-47	2.1	50
141	The hidden hand of chloride in hypertension. <i>Pflugers Archiv European Journal of Physiology</i> , 2015 , 467, 595-603	4.6	48
140	Genetics of hypertension: from experimental animals to humans. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2010 , 1802, 1299-308	6.9	47
139	The trans-ancestral genomic architecture of glycemic traits. <i>Nature Genetics</i> , 2021 , 53, 840-860	36.3	44
138	Elevated heart rate and cardiovascular outcomes in patients with coronary artery disease: clinical evidence and pathophysiological mechanisms. <i>Atherosclerosis</i> , 2010 , 212, 1-8	3.1	42
137	Associations of autozygosity with a broad range of human phenotypes. <i>Nature Communications</i> , 2019 , 10, 4957	17.4	40
136	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
135	Blood pressure response to patterns of weather fluctuations and effect on mortality. <i>Hypertension</i> , 2013 , 62, 190-6	8.5	40
134	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
133	PR interval genome-wide association meta-analysis identifies 50 loci associated with atrial and atrioventricular electrical activity. <i>Nature Communications</i> , 2018 , 9, 2904	17.4	39
132	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
131	Allopurinol initiation and change in blood pressure in older adults with hypertension. <i>Hypertension</i> , 2014 , 64, 1102-7	8.5	38

130	Familial and phenotypic associations of the aldosterone Renin ratio. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009 , 94, 4324-33	5.6	38
129	Metabolomic study of carotid-femoral pulse-wave velocity in women. <i>Journal of Hypertension</i> , 2015 , 33, 791-6; discussion 796	1.9	36
128	Genomics of elite sporting performance: what little we know and necessary advances. <i>Advances in Genetics</i> , 2013 , 84, 123-49	3.3	36
127	Genomics of hypertension: the road to precision medicine. <i>Nature Reviews Cardiology</i> , 2021 , 18, 235-250	14.8	34
126	Unsupervised Discovery and Comparison of Structural Families Across Multiple Samples in Untargeted Metabolomics. <i>Analytical Chemistry</i> , 2017 , 89, 7569-7577	7.8	33
125	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
124	Common polymorphisms in the CYP11B1 and CYP11B2 genes: evidence for a digenic influence on hypertension. <i>Hypertension</i> , 2013 , 61, 232-9	8.5	31
123	Serum uric acid level, longitudinal blood pressure, renal function, and long-term mortality in treated hypertensive patients. <i>Hypertension</i> , 2013 , 62, 105-11	8.5	31
122	Chromosome 2p shows significant linkage to antihypertensive response in the British Genetics of Hypertension Study. <i>Hypertension</i> , 2006 , 47, 603-8	8.5	31
121	The effects of sex and method of blood pressure measurement on genetic associations with blood pressure in the PAMELA study. <i>Journal of Hypertension</i> , 2010 , 28, 465-77	1.9	30
120	Glutathione S-transferase variants and hypertension. <i>Journal of Hypertension</i> , 2008 , 26, 1343-52	1.9	30
119	Genetic dysregulation of endothelin-1 is implicated in coronary microvascular dysfunction. <i>European Heart Journal</i> , 2020 , 41, 3239-3252	9.5	29
118	PTPRD gene associated with blood pressure response to atenolol and resistant hypertension. <i>Journal of Hypertension</i> , 2015 , 33, 2278-85	1.9	29
117	Hypertension and genome-wide association studies: combining high fidelity phenotyping and hypercontrols. <i>Journal of Hypertension</i> , 2008 , 26, 1275-81	1.9	29
116	Uromodulin, an emerging novel pathway for blood pressure regulation and hypertension. <i>Hypertension</i> , 2014 , 64, 918-23	8.5	28
115	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
114	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
113	Pharmacogenomic association of nonsynonymous SNPs in SIGLEC12, A1BG, and the selectin region and cardiovascular outcomes. <i>Hypertension</i> , 2013 , 62, 48-54	8.5	26

112	Discovery of rare variants associated with blood pressure regulation through meta-analysis of 1.3 million individuals. <i>Nature Genetics</i> , 2020 , 52, 1314-1332	36.3	26
111	A Combined Pathway and Regional Heritability Analysis Indicates NETRIN1 Pathway Is Associated With Major Depressive Disorder. <i>Biological Psychiatry</i> , 2017 , 81, 336-346	7.9	25
110	Exome-chip meta-analysis identifies novel loci associated with cardiac conduction, including ADAMTS6. <i>Genome Biology</i> , 2018 , 19, 87	18.3	25
109	Genome-Wide and Gene-Based Meta-Analyses Identify Novel Loci Influencing Blood Pressure Response to Hydrochlorothiazide. <i>Hypertension</i> , 2017 , 69, 51-59	8.5	25
108	Hematocrit predicts long-term mortality in a nonlinear and sex-specific manner in hypertensive adults. <i>Hypertension</i> , 2012 , 60, 631-8	8.5	25
107	Fibroblast growth factor 1 gene and hypertension: from the quantitative trait locus to positional analysis. <i>Circulation</i> , 2007 , 116, 1915-24	16.7	25
106	Resting heart rate and outcomes in patients with cardiovascular disease: where do we currently stand?. <i>Cardiovascular Therapeutics</i> , 2013 , 31, 215-23	3.3	24
105	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
104	Rare coding variants and X-linked loci associated with age at menarche. <i>Nature Communications</i> , 2015 , 6, 7756	17.4	23
103	Family history of premature cardiovascular disease: blood pressure control and long-term mortality outcomes in hypertensive patients. <i>European Heart Journal</i> , 2014 , 35, 563-70	9.5	23
102	Gene-centric meta-analyses for central adiposity traits in up to 57 412 individuals of European descent confirm known loci and reveal several novel associations. <i>Human Molecular Genetics</i> , 2014 , 23, 2498-510	5.6	22
101	Four genetic loci influencing electrocardiographic indices of left ventricular hypertrophy. <i>Circulation: Cardiovascular Genetics</i> , 2011 , 4, 626-35		22
100	Genetics and hypertension: is it time to change my practice?. <i>Canadian Journal of Cardiology</i> , 2012 , 28, 296-304	3.8	21
99	Polygenic risk for alcohol dependence associates with alcohol consumption, cognitive function and social deprivation in a population-based cohort. <i>Addiction Biology</i> , 2016 , 21, 469-80	4.6	21
98	Discontinuation of beta-blockers in cardiovascular disease: UK primary care cohort study. <i>International Journal of Cardiology</i> , 2013 , 167, 2695-9	3.2	20
97	Investigating shared aetiology between type 2 diabetes and major depressive disorder in a population based cohort. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2017 , 174, 227-234	3.5	20
96	TET2 and CSMD1 genes affect SBP response to hydrochlorothiazide in never-treated essential hypertensives. <i>Journal of Hypertension</i> , 2015 , 33, 1301-9	1.9	20
95	Genomics and Precision Medicine for Clinicians and Scientists in Hypertension. <i>Hypertension</i> , 2017 , 69, e10-e13	8.5	19

94	A PROgramme of Lifestyle Intervention in Families for Cardiovascular risk reduction (PROLIFIC Study): design and rationale of a family based randomized controlled trial in individuals with family history of premature coronary heart disease. <i>BMC Public Health</i> , 2017 , 17, 10	4.1	19
93	Acetaminophen use and risk of myocardial infarction and stroke in a hypertensive cohort. <i>Hypertension</i> , 2015 , 65, 1008-14	8.5	18
92	Exome-wide analysis of rare coding variation identifies novel associations with COPD and airflow limitation in MOCS3, IFIT3 and SERPINA12. <i>Thorax</i> , 2016 , 71, 501-9	7.3	18
91	Diastolic Blood Pressure J-Curve Phenomenon in a Tertiary-Care Hypertension Clinic. <i>Hypertension</i> , 2019 , 74, 767-775	8.5	18
90	Longitudinal Blood Pressure Control, Long-Term Mortality, and Predictive Utility of Serum Liver Enzymes and Bilirubin in Hypertensive Patients. <i>Hypertension</i> , 2015 , 66, 37-43	8.5	18
89	Discovery of novel heart rate-associated loci using the Exome Chip. <i>Human Molecular Genetics</i> , 2017 , 26, 2346-2363	5.6	17
88	Genome-wide Regional Heritability Mapping Identifies a Locus Within the TOX2 Gene Associated With Major Depressive Disorder. <i>Biological Psychiatry</i> , 2017 , 82, 312-321	7.9	17
87	Novel Urinary Peptidomic Classifier Predicts Incident Heart Failure. <i>Journal of the American Heart Association</i> , 2017 , 6,	6	17
86	Urinary antihypertensive drug metabolite screening using molecular networking coupled to high-resolution mass spectrometry fragmentation. <i>Metabolomics</i> , 2016 , 12, 125	4.7	17
85	Multi-ancestry GWAS of the electrocardiographic PR interval identifies 202 loci underlying cardiac conduction. <i>Nature Communications</i> , 2020 , 11, 2542	17.4	16
84	Genomic approaches to coronary artery disease. <i>Indian Journal of Medical Research</i> , 2010 , 132, 567-78	2.9	16
83	An Empirical Comparison of Joint and Stratified Frameworks for Studying G x E Interactions: Systolic Blood Pressure and Smoking in the CHARGE Gene-Lifestyle Interactions Working Group. <i>Genetic Epidemiology</i> , 2016 , 40, 404-15	2.6	15
82	Rationale and design of the British Heart Foundation (BHF) Coronary Microvascular Angina (CorMicA) stratified medicine clinical trial. <i>American Heart Journal</i> , 2018 , 201, 86-94	4.9	14
81	Heritability analyses show visit-to-visit blood pressure variability reflects different pathological phenotypes in younger and older adults: evidence from UK twins. <i>Journal of Hypertension</i> , 2013 , 31, 2356-81	1.9	14
80	Acetaminophen use and change in blood pressure in a hypertensive population. <i>Journal of Hypertension</i> , 2013 , 31, 1485-90; discussion 1490	1.9	14
79	Association between ADRA1A gene and the metabolic syndrome: candidate genes and functional counterpart in the PAMELA population. <i>Journal of Hypertension</i> , 2011 , 29, 1121-7	1.9	14
78	ExomeChip-Wide Analysis of 95 626 Individuals Identifies 10 Novel Loci Associated With QT and JT Intervals. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e001758	5.2	14
77	Genomics of hypertension. <i>Pharmacological Research</i> , 2017 , 121, 219-229	10.2	13

76	Risk of Neuropsychiatric Adverse Effects of Lipid-Lowering Drugs: A Mendelian Randomization Study. <i>International Journal of Neuropsychopharmacology</i> , 2018 , 21, 1067-1075	5.8	13
75	Gene and environmental interactions according to the components of lifestyle modifications in hypertension guidelines. <i>Environmental Health and Preventive Medicine</i> , 2019 , 24, 19	4.2	12
74	Variation in the SLC23A1 gene does not influence cardiometabolic outcomes to the extent expected given its association with L-ascorbic acid. <i>American Journal of Clinical Nutrition</i> , 2015 , 101, 202-9	7	12
73	Effects of Calcium, Magnesium, and Potassium Concentrations on Ventricular Repolarization in Unselected Individuals. <i>Journal of the American College of Cardiology</i> , 2019 , 73, 3118-3131	15.1	12
72	Rationale and design of the Medical Research Council's Precision Medicine with Zibotentan in Microvascular Angina (PRIZE) trial. <i>American Heart Journal</i> , 2020 , 229, 70-80	4.9	12
71	Genomics of Blood Pressure and Hypertension: Extending the Mosaic Theory Toward Stratification. <i>Canadian Journal of Cardiology</i> , 2020 , 36, 694-705	3.8	11
70	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
69	Common and Rare Coding Genetic Variation Underlying the Electrocardiographic PR Interval. <i>Circulation Genomic and Precision Medicine</i> , 2018 , 11, e002037	5.2	11
68	Contrasting mortality risks among subgroups of treated hypertensive patients developing new-onset diabetes. <i>European Heart Journal</i> , 2016 , 37, 968-74	9.5	10
67	Recent Findings in the Genetics of Blood Pressure: How to Apply in Practice or Is a Moonshot Required?. <i>Current Hypertension Reports</i> , 2018 , 20, 54	4.7	9
66	Pharmacogenomics and Stratified Medicine 2014 , 3-25		9
65	The Pharmacogenomics of Anti-Hypertensive Therapy. <i>Pharmaceuticals</i> , 2010 , 3, 1779-1791	5.2	9
64	The genetics of cardiovascular disease. <i>Trends in Endocrinology and Metabolism</i> , 2008 , 19, 309-16	8.8	9
63	Development, Evaluation, and Comparison of Land Use Regression Modeling Methods to Estimate Residential Exposure to Nitrogen Dioxide in a Cohort Study. <i>Environmental Science & Technology</i> , 2016 , 50, 11085-11093	10.3	9
62	The relationship between antihypertensive medications and mood disorders: analysis of linked healthcare data for 1.8 million patients. <i>Psychological Medicine</i> , 2021 , 51, 1183-1191	6.9	9
61	Gene Variants at Loci Related to Blood Pressure Account for Variation in Response to Antihypertensive Drugs Between Black and White Individuals. <i>Hypertension</i> , 2019 , 74, 614-622	8.5	8
60	Phenome-wide association analysis of LDL-cholesterol lowering genetic variants in PCSK9. <i>BMC Cardiovascular Disorders</i> , 2019 , 19, 240	2.3	8
59	Pharmacokinetic Pharmacogenomics 2014 , 341-364		8

58	Genome-wide association studies of hypertension: light at the end of the tunnel. <i>International Journal of Hypertension</i> , 2010 , 2010, 509581	2.4	8
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