Meta-analysis of genome-wide association studies identity with blood pressure variation in east Asians

Nature Genetics

43, 531-538

DOI: 10.1038/ng.834

Citation Report

#	Article	IF	CITATIONS
1	KAREBrowser: SNP database of Korea Association REsource project., 2010,,.		0
2	Genetic variants in novel pathways influence blood pressure and cardiovascular disease risk. Nature, 2011, 478, 103-109.	13.7	1,855
3	Large-scale genome-wide association studies in east Asians identify new genetic loci influencing metabolic traits. Nature Genetics, 2011, 43, 990-995.	9.4	270
4	Top Three Pharmacogenomics and Personalized Medicine Applications at the Nexus of Renal Pathophysiology and Cardiovascular Medicine. Current Pharmacogenomics and Personalized Medicine, 2011, 9, 299-322.	0.2	9
5	Under pressure: the search for the essential mechanisms of hypertension. Nature Medicine, 2011, 17, 1402-1409.	15.2	247
6	Blood Pressure Loci Identified with a Gene-Centric Array. American Journal of Human Genetics, 2011, 89, 688-700.	2.6	159
8	Beyond Genome-Wide Association Studies: New Strategies for Identifying Genetic Determinants of Hypertension. Current Hypertension Reports, 2011, 13, 442-451.	1.5	35
9	Human genetics, natriuretic peptides and hypertension. BMC Pharmacology, 2011, 11, .	0.4	O
10	A Rare Variant at the <i>KYNU </i> Gene Is Associated With Kynureninase Activity and Essential Hypertension in the Han Chinese Population. Circulation: Cardiovascular Genetics, 2011, 4, 687-694.	5.1	14
11	ALDH2 Activator Inhibits Increased Myocardial Infarction Injury by Nitroglycerin Tolerance. Science Translational Medicine, 2011, 3, 107ra111.	5.8	73
12	Genome-wide association study identifies six new loci influencing pulse pressure and mean arterial pressure. Nature Genetics, 2011, 43, 1005-1011.	9.4	403
13	A Common Genetic Variant of <i>FCN3/CD164L2</i> Is Associated With Essential Hypertension in a Chinese Population. Clinical and Experimental Hypertension, 2012, 34, 377-382.	0.5	6
14	Novel findings and future directions on the genetics of hypertension. Current Opinion in Nephrology and Hypertension, 2012, 21, 500-507.	1.0	49
15	Task3 Potassium Channel Gene Invalidation Causes Low Renin and Salt-Sensitive Arterial Hypertension. Endocrinology, 2012, 153, 4740-4748.	1.4	63
16	Efficiency of trans-ethnic genome-wide meta-analysis and fine-mapping. European Journal of Human Genetics, 2012, 20, 1300-1307.	1.4	20
17	Common genetic factors for hematological traits in Humans. Journal of Human Genetics, 2012, 57, 161-169.	1.1	31
18	Association of Natriuretic Peptide Receptor-C Gene with Ischemic Stroke and Hypertension in Chinese Han Population. Clinical and Experimental Hypertension, 2012, 34, 504-509.	0.5	7
19	Ethnic differences in genetic predisposition to hypertension. Hypertension Research, 2012, 35, 574-581.	1.5	51

#	ARTICLE	IF	CITATIONS
20	Effect of mitochondrial aldehyde dehydrogenase-2 genotype on cardioprotection in patients with congenital heart disease. European Heart Journal, 2012, 33, 1606-1614.	1.0	35
21	Recapitulation of genome-wide association studies on pulse pressure and mean arterial pressure in the Korean population. Journal of Human Genetics, 2012, 57, 391-393.	1.1	6
22	Genomic epidemiology of blood pressure salt sensitivity. Journal of Hypertension, 2012, 30, 861-873.	0.3	53
23	Novel genes as primary triggers for polygenic hypertension. Journal of Hypertension, 2012, 30, 81-86.	0.3	17
24	Essential hypertension. Journal of Hypertension, 2012, 30, 42-45.	0.3	18
26	Genetic basis of blood pressure and hypertension. Trends in Genetics, 2012, 28, 397-408.	2.9	117
27	Genome-wide analysis of copy number variations reveals that aging processes influence body fat distribution in Korea Associated Resource (KARE) cohorts. Human Genetics, 2012, 131, 1795-1804.	1.8	6
28	Characterization of the ATP2B gene family in blood pressure. Genes and Genomics, 2012, 34, 539-547.	0.5	1
29	Genome-wide association analysis identifies three new breast cancer susceptibility loci. Nature Genetics, 2012, 44, 312-318.	9.4	256
30	Advancing management of hypertension through pharmacogenomics. Annals of Medicine, 2012, 44, S17-S22.	1.5	41
31	The stroke-prone spontaneously hypertensive rat: still a useful model for post-GWAS genetic studies?. Hypertension Research, 2012, 35, 477-484.	1.5	36
32	Gene–environment interactions of selected pharmacogenes in arterial hypertension. Expert Review of Clinical Pharmacology, 2012, 5, 677-686.	1.3	8
33	Reevaluation of the association of seven candidate genes with blood pressure and hypertension: a replication study and meta-analysis with a larger sample size. Hypertension Research, 2012, 35, 825-831.	1.5	44
34	Common ALDH2 genetic variants predict development of hypertension in the SAPPHIRe prospective cohort: Gene-environmental interaction with alcohol consumption. BMC Cardiovascular Disorders, 2012, 12, 58.	0.7	39
35	Evaluation of the imputation performance of the program IMPUTE in an admixed sample from Mexico City using several model designs. BMC Medical Genomics, 2012, 5, 12.	0.7	9
36	The pharmacogenetics of \hat{l}^2 -adrenergic receptor antagonists in the treatment of hypertension and heart failure. Expert Opinion on Drug Metabolism and Toxicology, 2012, 8, 767-790.	1.5	14
37	Parameters in Dynamic Models of Complex Traits are Containers of Missing Heritability. PLoS Computational Biology, 2012, 8, e1002459.	1.5	24
38	Hunting for genes for hypertension: the Millennium Genome Project for Hypertension. Hypertension Research, 2012, 35, 567-573.	1.5	14

#	ARTICLE	IF	CITATIONS
39	Genetic variation in metabolic phenotypes: study designs and applications. Nature Reviews Genetics, 2012, 13, 759-769.	7.7	165
40	Genome-wide association study of coronary artery disease in the Japanese. European Journal of Human Genetics, 2012, 20, 333-340.	1.4	156
41	Physical activity modifies the associations between genetic variants and hypertension in the Chinese children. Atherosclerosis, 2012, 225, 376-380.	0.4	25
42	Regulation of insulin and typeÂ1 insulinâ€like growth factor signaling and action by the <scp>G</scp> rb10/14 and <scp>SH</scp> 2 <scp>B</scp> 1/ <scp>B</scp> 2 adaptor proteins. FEBS Journal, 2013, 280, 794-816.	2.2	49
43	Meta-analysis identifies multiple loci associated with kidney function–related traits in east Asian populations. Nature Genetics, 2012, 44, 904-909.	9.4	254
45	Gene–Sodium Interaction and Blood Pressure: Findings from Genomics Research of Blood Pressure Salt Sensitivity. Progress in Molecular Biology and Translational Science, 2012, 108, 237-260.	0.9	3
46	Blood Pressure Variability and Vascular Dysfunction in Essential Hypertension. Journal of the Korean Society of Hypertension, 2012, 18, 75.	0.2	2
47	Genomewide Association Studies in Cardiovascular Diseaseâ€"An Update 2011. Clinical Chemistry, 2012, 58, 92-103.	1.5	64
48	Genome-wide association study in Han Chinese identifies four new susceptibility loci for coronary artery disease. Nature Genetics, 2012, 44, 890-894.	9.4	295
49	Strong protective effect of the aldehyde dehydrogenase gene (ALDH2) 504lys (*2) allele against alcoholism and alcohol-induced medical diseases in Asians. Human Genetics, 2012, 131, 725-737.	1.8	132
50	Current understanding of human genetics and genetic analysis of psoriasis. Journal of Dermatology, 2012, 39, 231-241.	0.6	54
51	Between Candidate Genes and Whole Genomes: Time for Alternative Approaches in Blood Pressure Genetics. Current Hypertension Reports, 2012, 14, 46-61.	1.5	37
52	Mendelian Randomization: Application to Cardiovascular Disease. Current Hypertension Reports, 2012, 14, 29-37.	1.5	38
53	cGMP becomes a drug target. Naunyn-Schmiedeberg's Archives of Pharmacology, 2012, 385, 243-252.	1.4	33
54	Association of a functional single-nucleotide polymorphism in the ALDH2 gene with essential hypertension depends on drinking behavior in a Chinese Han population. Journal of Human Hypertension, 2013, 27, 181-186.	1.0	46
55	Characterization of functional variants in 33 blood pressure loci using 1000 genomes project data. Genes and Genomics, 2013, 35, 387-393.	0.5	3
56	Common variants in the ATP2B1 gene are associated with hypertension and arterial stiffness in Chinese population. Molecular Biology Reports, 2013, 40, 1867-1873.	1.0	20
57	Genetics of Atherosclerotic Cardiovascular Disease. , 2013, , 1-37.		2

#	Article	IF	CITATIONS
58	Pharmacogenetics and Pharmacogenomics. , 2013, , 1-27.		0
59	P.R4810K, a polymorphism of RNF213, the susceptibility gene for moyamoya disease, is associated with blood pressure. Environmental Health and Preventive Medicine, 2013, 18, 121-129.	1.4	59
60	Genome-wide Association Analysis of Blood-Pressure Traits in African-Ancestry Individuals Reveals Common Associated Genes in African and Non-African Populations. American Journal of Human Genetics, 2013, 93, 545-554.	2.6	189
61	Mapping eQTLs in the Norfolk Island Genetic Isolate Identifies Candidate Genes for CVD Risk Traits. American Journal of Human Genetics, 2013, 93, 1087-1099.	2.6	28
62	Recapitulation of four hypertension susceptibility genes (CSK, CYP17A1, MTHFR, and FGF5) in East Asians. Metabolism: Clinical and Experimental, 2013, 62, 196-203.	1.5	38
63	Association between aldehyde dehydrogenase 2 polymorphisms and the incidence of diabetic retinopathy among Japanese subjects with type 2 diabetes mellitus. Cardiovascular Diabetology, 2013, 12, 132.	2.7	26
64	Progress and Future Aspects in Genetics of Human Hypertension. Current Hypertension Reports, 2013, 15, 676-686.	1.5	20
65	Genetics and Genomics for the Prevention and Treatment of Cardiovascular Disease: Update. Circulation, 2013, 128, 2813-2851.	1.6	100
66	Deep Whole-Genome Sequencing of 100 Southeast Asian Malays. American Journal of Human Genetics, 2013, 92, 52-66.	2.6	153
67	Genes for blood pressure: an opportunity to understand hypertension. European Heart Journal, 2013, 34, 951-961.	1.0	163
68	The molecular basis of blood pressure variation. Pediatric Nephrology, 2013, 28, 387-399.	0.9	28
69	Influence of Adiponectin and Resistin Gene Polymorphisms on Quantitative Traits Related to Metabolic Syndrome Among Malay, Chinese, and Indian Men in Malaysia. Biochemical Genetics, 2013, 51, 166-174.	0.8	4
70	Single nucleotide polymorphism in genome-wide association of human population: A tool for broad spectrum service. Egyptian Journal of Medical Human Genetics, 2013, 14, 123-134.	0.5	55
71	Genetic modification of hypertension by $sGC\hat{l}\pm 1$. Trends in Cardiovascular Medicine, 2013, 23, 312-318.	2.3	4
72	Gene-based copy number variation study reveals a microdeletion at 12q24 that influences height in the Korean population. Genomics, 2013, 101, 134-138.	1.3	17
74	Pathophysiology of Hypertension in the Absence of Nitric Oxide/Cyclic GMP Signaling. Current Hypertension Reports, 2013, 15, 47-58.	1.5	41
75	The Power of Meta-Analysis in Genome-Wide Association Studies. Annual Review of Genomics and Human Genetics, 2013, 14, 441-465.	2.5	107
76	Genetic Polymorphisms of Alcohol Dehydrogenaseâ€1 <scp>B</scp> and Aldehyde Dehydrogenaseâ€2 and Liver Cirrhosis, Chronic Calcific Pancreatitis, Diabetes Mellitus, and Hypertension Among <scp>J</scp> apanese Alcoholic Men. Alcoholism: Clinical and Experimental Research, 2013, 37, 1391-1401.	1.4	63

#	Article	IF	Citations
77	Association of a single nucleotide polymorphism of the NPR3 gene promoter with early onset ischemic stroke in an Italian cohort. European Journal of Internal Medicine, 2013, 24, 80-82.	1.0	13
78	Common Polymorphism rs11191548 Near the CYP17A1 Gene Is Associated With Hypertension and Systolic Blood Pressure in the Han Chinese Population. American Journal of Hypertension, 2013, 26, 465-472.	1.0	15
79	Genetic discoveries in hypertension: steps on the road to therapeutic translation. Heart, 2013, 99, 1645-1651.	1.2	15
80	Exploring causal associations between alcohol and coronary heart disease risk factors: findings from a Mendelian randomization study in the Copenhagen General Population Study. European Heart Journal, 2013, 34, 2519-2528.	1.0	81
81	Genome-Wide Analysis of Blood Pressure Variability and Ischemic Stroke. Stroke, 2013, 44, 2703-2709.	1.0	17
82	ALDH2, a novel protector against stroke?. Cell Research, 2013, 23, 874-875.	5.7	18
83	Common Genetic Variants in the Endothelial System Predict Blood Pressure Response to Sodium Intake: The GenSalt Study. American Journal of Hypertension, 2013, 26, 643-656.	1.0	24
84	Silencing of Atp2b1 increases blood pressure through vasoconstriction. Journal of Hypertension, 2013, 31, 1575-1583.	0.3	23
85	Advances in Blood Pressure Genomics. Circulation Research, 2013, 112, 1365-1379.	2.0	106
86	ATP2B1 and blood pressure. Current Opinion in Nephrology and Hypertension, 2013, 22, 177-184.	1.0	18
87	Natriuretic Peptide Receptor-3 Gene (NPR3). Circulation: Cardiovascular Genetics, 2013, 6, 201-210.	5.1	12
88	Genome-Wide Association Study Identifies 8 Novel Loci Associated With Blood Pressure Responses to Interventions in Han Chinese. Circulation: Cardiovascular Genetics, 2013, 6, 598-607.	5.1	64
89	Comparing methods for performing trans-ethnic meta-analysis of genome-wide association studies. Human Molecular Genetics, 2013, 22, 2303-2311.	1.4	63
90	Analysis of Sex Hormone Genes Reveals Gender Differences in the Genetic Etiology of Blood Pressure Salt Sensitivity: The GenSalt Study. American Journal of Hypertension, 2013, 26, 191-200.	1.0	24
91	Modularization and epistatic hierarchy determine homeostatic actions of multiple blood pressure quantitative trait loci. Human Molecular Genetics, 2013, 22, 4451-4459.	1.4	29
92	Candidate genes revisited in the genetics of hypertension and blood pressure. Hypertension Research, 2013, 36, 1032-1034.	1.5	1
93	A genome-wide association study of a coronary artery disease risk variant. Journal of Human Genetics, 2013, 58, 120-126.	1.1	135
94	Genome-Wide Association Study Meta-Analysis Reveals Transethnic Replication of Mean Arterial and Pulse Pressure Loci. Hypertension, 2013, 62, 853-859.	1.3	63

#	Article	IF	Citations
95	Influence of Obesity on Association Between Genetic Variants Identified by Genome-Wide Association Studies and Hypertension Risk in Chinese Children. American Journal of Hypertension, 2013, 26, 990-996.	1.0	36
96	Lack of association between STK39 and hypertension in the Chinese population. Journal of Human Hypertension, 2013, 27, 294-297.	1.0	14
97	Associations Between Polymorphisms in the Glucocorticoid-Receptor Gene and Cardiovascular Risk Factors in a Chinese Population. Journal of Epidemiology, 2013, 23, 389-395.	1.1	18
98	The combination of mitochondrial low enzyme-activity aldehyde dehydrogenase 2 allele and superoxide dismutase 2 genotypes increases the risk of hypertension in relation to alcohol consumption. Pharmacogenetics and Genomics, 2013, 23, 34-37.	0.7	29
99	MicroRNA-34a Promotes Cardiomyocyte Apoptosis Post Myocardial Infarction Through Down-regulating Aldehyde Dehydrogenase 2. Current Pharmaceutical Design, 2013, 19, 4865-4873.	0.9	67
100	Hypertension and Genetic Variation in Endothelial-Specific Genes. PLoS ONE, 2013, 8, e62035.	1.1	9
101	Gene-alcohol interactions identify several novel blood pressure loci including a promising locus near SLC16A9. Frontiers in Genetics, 2013, 4, 277.	1.1	33
102	The role of SNP-loop diuretic interactions in hypertension across ethnic groups in HyperGEN. Frontiers in Genetics, 2013, 4, 304.	1.1	11
103	Atrial Fibrillation Associated Chromosome 4q25 Variants Are Not Associated with PITX2c Expression in Human Adult Left Atrial Appendages. PLoS ONE, 2014, 9, e86245.	1.1	56
104	Cross-Sectional and Longitudinal Replication Analyses of Genome-Wide Association Loci of Type 2 Diabetes in Han Chinese. PLoS ONE, 2014, 9, e91790.	1.1	17
105	On the Analysis of a Repeated Measure Design in Genome-Wide Association Analysis. International Journal of Environmental Research and Public Health, 2014, 11, 12283-12303.	1.2	8
106	Genomic architecture of sickle cell disease in West African children. Frontiers in Genetics, 2014, 5, 26.	1.1	11
107	Genome-Wide Association Study Identifies Two Novel Loci with Sex-Specific Effects for Type 2 Diabetes Mellitus and Glycemic Traits in a Korean Population. Diabetes and Metabolism Journal, 2014, 38, 375.	1.8	30
108	Hypertension Pharmacogenomics. , 2014, , 747-778.		0
109	Mapping of a blood pressure QTL on chromosome 17 in American Indians of the strong heart family study. BMC Cardiovascular Disorders, 2014, 14, 158.	0.7	3
110	Prevention of Hypertension and Cardiovascular Diseases. Hypertension, 2014, 63, 655-660.	1.3	54
111	Disease Risk Factors Identified Through Shared Genetic Architecture and Electronic Medical Records. Science Translational Medicine, 2014, 6, 234ra57.	5 . 8	58
112	Chapter 13. Secondary hypertension. Hypertension Research, 2014, 37, 349-361.	1.5	5

#	Article	IF	CITATIONS
113	ANTXR2 is a potential causative gene in the genome-wide association study of the blood pressure locus 4q21. Hypertension Research, 2014, 37, 811-817.	1.5	13
115	Meta-analysis of genome-wide association studies in East Asian-ancestry populations identifies four new loci for body mass index. Human Molecular Genetics, 2014, 23, 5492-5504.	1.4	192
116	Probing genetic overlap in the regulation of systolic and diastolic blood pressure in Danish and Chinese twins. Hypertension Research, 2014, 37, 954-959.	1.5	4
117	Methodologies in the Era of Cardiovascular "Omics― , 2014, , 15-55.		O
118	Aldehyde dehydrogenase 2 as a potential protective factor for renal insufficiency in Japanese subjects with heart failure: a pilot study. Journal of Human Hypertension, 2014, 28, 279-281.	1.0	2
119	Epidemiology of Hypertension in Children. , 2014, , 1-57.		0
120	The Association of the Vanin-1 N131S Variant with Blood Pressure Is Mediated by Endoplasmic Reticulum-Associated Degradation and Loss of Function. PLoS Genetics, 2014, 10, e1004641.	1.5	16
121	Gene expression responses of threespine stickleback to salinity: implications for salt-sensitive hypertension. Frontiers in Genetics, 2014, 5, 312.	1.1	39
122	A meta-analysis of genome-wide association studies for adiponectin levels in East Asians identifies a novel locus near WDR11-FGFR2. Human Molecular Genetics, 2014, 23, 1108-1119.	1.4	68
123	Atrial natriuretic peptide gene variants and circulating levels: implications in cardiovascular diseases. Clinical Science, 2014, 127, 1-13.	1.8	29
124	Association of common variants in/near six genes (ATP2B1, CSK, MTHFR, CYP17A1, STK39 and FGF5) with blood pressure/hypertension risk in Chinese children. Journal of Human Hypertension, 2014, 28, 32-36.	1.0	43
125	Blood Pressure Responses to Dietary Sodium and Potassium Interventions and the Cold Pressor Test: The GenSalt Replication Study in Rural North China. American Journal of Hypertension, 2014, 27, 72-80.	1.0	8
126	Cardiovascular consequences of a polygenetic component of blood pressure in an urban-based longitudinal study. Journal of Hypertension, 2014, 32, 1424-1428.	0.3	7
127	NEDD4L in essential hypertension. Journal of Hypertension, 2014, 32, 230-232.	0.3	3
128	Risk of progression to hypertension in nonhypertensive Japanese workers aged 20–64 years. Journal of Hypertension, 2014, 32, 236-244.	0.3	18
129	A Three-Stage Genome-Wide Association Study Combining Multilocus Test and Gene Expression Analysis for Young-Onset Hypertension in Taiwan Han Chinese. American Journal of Hypertension, 2014, 27, 819-827.	1.0	12
130	The â°344C/T polymorphism in the <i>CYP11B2</i> gene is associated with essential hypertension in the Chinese. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2014, 15, 150-155.	1.0	13
131	Body Mass Index Modulates Blood Pressure Heritability: The Family Blood Pressure Program. American Journal of Hypertension, 2014, 27, 610-619.	1.0	7

#	ARTICLE	IF	CITATIONS
132	Genetic Variants in Renalase and Blood Pressure Responses to Dietary Salt and Potassium Interventions: A Family-Based Association Study. Kidney and Blood Pressure Research, 2014, 39, 497-506.	0.9	17
133	Mitochondrial aldehyde dehydrogenase 2 accentuates aging-induced cardiac remodeling and contractile dysfunction: role of AMPK, Sirt1, and mitochondrial function. Free Radical Biology and Medicine, 2014, 71, 208-220.	1.3	112
134	Targeting Aldehyde Dehydrogenase 2: New Therapeutic Opportunities. Physiological Reviews, 2014, 94, 1-34.	13.1	465
135	Blood Pressure Genetics and Hypertension: Genome-Wide Analysis and Role of Ancestry. Current Genetic Medicine Reports, 2014, 2, 13-22.	1.9	3
136	On individual genome-wide association studies and their meta-analysis. Human Genetics, 2014, 133, 265-279.	1.8	30
137	Male-specific genetic effect on hypertension and metabolic disorders. Human Genetics, 2014, 133, 311-319.	1.8	29
138	Rare variants and cardiovascular disease. Briefings in Functional Genomics, 2014, 13, 384-391.	1.3	12
139	Natriuretic peptides in cardiovascular diseases: current use and perspectives. European Heart Journal, 2014, 35, 419-425.	1.0	221
140	Ethanol Reduces Lifespan, Body Weight, and Serum Alanine Aminotransferase Level of <i> Aldehyde Dehydrogenase 2 < /i > Knockout Mouse. Alcoholism: Clinical and Experimental Research, 2014, 38, 1883-1893.</i>	1.4	13
141	Genetics of hypertension: discoveries from the bench to human populations. American Journal of Physiology - Renal Physiology, 2014, 306, F1-F11.	1.3	28
142	The Path to Personalized Cardiovascular Medicine. , 2014, , 837-871.		0
143	Gene-Age Interactions in Blood Pressure Regulation: A Large-Scale Investigation with the CHARGE, Global BPgen, and ICBP Consortia. American Journal of Human Genetics, 2014, 95, 24-38.	2.6	109
144	Cardioprotective effects of moderate red wine consumption: Polyphenols vs. ethanol. Journal of Applied Biomedicine, 2014, 12, 193-202.	0.6	27
145	Association between alcohol and cardiovascular disease: Mendelian randomisation analysis based on individual participant data. BMJ, The, 2014, 349, g4164-g4164.	3.0	528
146	Genetics, Ancestry, and Hypertension: Implications for Targeted Antihypertensive Therapies. Current Hypertension Reports, 2014, 16, 461.	1.5	34
147	Genome-Wide Association Studies of Genetic Impact on Cardiovascular and Metabolic Diseases in Asians: Opportunity for Discovery. Current Cardiovascular Risk Reports, 2014, 8, 1.	0.8	2
148	Identification of a genetic variant at $2q12.1$ associated with blood pressure in East-Asians by genome-wide scan including gene-environment interactions. BMC Medical Genetics, 2014, 15, 65.	2.1	14
149	Replication of the top 10 most significant polymorphisms from a large blood pressure genome-wide association study of northeastern Han Chinese East Asians. Hypertension Research, 2014, 37, 134-138.	1.5	25

#	Article	IF	Citations
150	Elevated blood pressure: Our family's fault? The genetics of essential hypertension. World Journal of Cardiology, 2014, 6, 327.	0.5	27
151	Etiology of Individual Differences in Human Health and Longevity. Annual Review of Gerontology and Geriatrics, 2014, 34, 189-227.	0.5	6
152	Role of the Na ⁺ H ⁺ exchanger 3 in angiotensin II-induced hypertension in NHE3-deficient mice with transgenic rescue of NHE3 in small intestines. Physiological Reports, 2015, 3, e12605.	0.7	28
153	Heterogeneous Effects of Association Between Blood Pressure Loci and Coronary Artery Disease in East Asian Individuals. Circulation Journal, 2015, 79, 830-838.	0.7	6
154	Natriuretic Peptides and Cardiometabolic Health. Circulation Journal, 2015, 79, 1647-1655.	0.7	73
155	Alcohol intake and cardiovascular risk factors: A Mendelian randomisation study. Scientific Reports, 2015, 5, 18422.	1.6	78
156	Exploring causal associations of alcohol with cardiovascular and metabolic risk factors in a Chinese population using Mendelian randomization analysis. Scientific Reports, 2015, 5, 14005.	1.6	38
157	Epigenomic mapping and effect sizes of noncoding variants associated with psychotropic drug response. Pharmacogenomics, 2015, 16, 1565-1583.	0.6	21
158	Expression of STK39 in peripheral blood of hypertension patients and the relationship between its genetic polymorphism and blood pressure. Genetics and Molecular Research, 2015, 14, 16461-16468.	0.3	3
159	Association of NPRA and NPRC gene variants and hypertension in Mongolian population. Genetics and Molecular Research, 2015, 14, 18494-18502.	0.3	1
160	Aldehyde Dehydrogenase Polymorphisms and Blood Pressure Elevation in the Japanese: A Cross-Sectional and a Longitudinal Study over 20 Years in the Shimane CoHRE Study. Disease Markers, 2015, 2015, 1-4.	0.6	5
161	A Single Nucleotide Polymorphism near the CYP17A1 Gene Is Associated with Left Ventricular Mass in Hypertensive Patients under Pharmacotherapy. International Journal of Molecular Sciences, 2015, 16, 17456-17468.	1.8	9
162	The Role of Genetic Risk Score in Predicting the Risk of Hypertension in the Korean population: Korean Genome and Epidemiology Study. PLoS ONE, 2015, 10, e0131603.	1.1	28
163	CYP17A1and Blood Pressure Reactivity to Stress in Adolescence. International Journal of Hypertension, 2015, 2015, 1-9.	0.5	6
164	High Blood Pressure and Its Association With Incident Diabetes Over 10 Years in the Korean Genome and Epidemiology Study (KoGES). Diabetes Care, 2015, 38, 1333-1338.	4.3	39
165	Genome-Wide Association Studies (Gwas) of Blood Pressure in Different Populations. , 2015, , 157-184.		О
166	Pharmacogenomics of Hypertension: A Genomeâ€Wide, Placeboâ€Controlled Crossâ€Over Study, Using Four Classes of Antihypertensive Drugs. Journal of the American Heart Association, 2015, 4, e001521.	1.6	74
167	Arterial Stiffness and Decline in Kidney Function. Clinical Journal of the American Society of Nephrology: CJASN, 2015, 10, 2190-2197.	2.2	117

#	Article	IF	CITATIONS
168	Role of the Na ⁺ /H ⁺ exchanger 3 in angiotensin II-induced hypertension. Physiological Genomics, 2015, 47, 479-487.	1.0	28
169	Associations Between Genetic Variants of the Natriuretic Peptide System and Blood Pressure Response to Dietary Sodium Intervention: The GenSalt Study. American Journal of Hypertension, 2016, 29, 397-404.	1.0	2
170	The Role of Rare Variants in Systolic Blood Pressure: Analysis of ExomeChip Data in HyperGEN African Americans. Human Heredity, 2015, 79, 20-27.	0.4	13
171	Endothelial Gata5 transcription factor regulates blood pressure. Nature Communications, 2015, 6, 8835.	5.8	35
172	Genetic Markers in Prediction of Cardiovascular Disease., 2015,, 239-260.		0
173	Epithelial Sodium Transport and Its Control by Aldosterone: The Story of Our Internal Environment Revisited. Physiological Reviews, 2015, 95, 297-340.	13.1	217
174	A genome-wide association study identifies PLCL2 and AP3D1-DOT1L-SF3A2 as new susceptibility loci for myocardial infarction in Japanese. European Journal of Human Genetics, 2015, 23, 374-380.	1.4	48
175	Exploring hypertension genomeâ€wide association studies findings and impact on pathophysiology, pathways, and pharmacogenetics. Wiley Interdisciplinary Reviews: Systems Biology and Medicine, 2015, 7, 73-90.	6.6	30
176	Dysregulation of T cell Subsets in the Pathogenesis of Hypertension. Current Hypertension Reports, 2015, 17, 8.	1.5	19
177	Integromic Analysis of Genetic Variation and Gene Expression Identifies Networks for Cardiovascular Disease Phenotypes. Circulation, 2015, 131, 536-549.	1.6	65
178	Genetic and Functional Evidence Supports <i>LPAR1</i> as a Susceptibility Gene for Hypertension. Hypertension, 2015, 66, 641-646.	1.3	16
179	Long-range regulatory interactions at the 4q25 atrial fibrillation risk locus involve PITX2c and ENPEP. BMC Biology, 2015, 13, 26.	1.7	53
180	Variant Near <i>FGF5</i> Has Stronger Effects on Blood Pressure in Chinese With a Higher Body Mass Index. American Journal of Hypertension, 2015, 28, 1031-1037.	1.0	17
181	Genetic Contribution to the Variance of Blood Pressure and Heart Rate: A Systematic Review and Meta-Regression of Twin Studies. Twin Research and Human Genetics, 2015, 18, 158-170.	0.3	39
182	Genetic and Molecular Aspects of Hypertension. Circulation Research, 2015, 116, 937-959.	2.0	218
183	Joseph A. Vita, MD, 1956–2014. Journal of the American Heart Association, 2015, 4, e001778.	1.6	7
184	Generation of Hypertension-Associated STK39 Polymorphism Knockin Cell Lines With the Clustered Regularly Interspaced Short Palindromic Repeats/Cas9 System. Hypertension, 2015, 66, 1199-1206.	1.3	10
185	Trans-ancestry genome-wide association study identifies 12 genetic loci influencing blood pressure and implicates a role for DNA methylation. Nature Genetics, 2015, 47, 1282-1293.	9.4	294

#	Article	IF	CITATIONS
186	Genetic Predisposition to Higher Blood Pressure Increases Risk of Incident Hypertension and Cardiovascular Diseases in Chinese. Hypertension, 2015, 66, 786-792.	1.3	22
187	An update on genome-wide association studies of hypertension. Applied Informatics, 2015, 2, .	0.5	12
188	Genetics of Cardiovascular Disease. , 2015, , 117-127.		0
189	Genome-wide association studies and contribution to cardiovascular physiology. Physiological Genomics, 2015, 47, 365-375.	1.0	11
190	Prevention of Cardiovascular Diseases., 2015,,.		1
191	Effects of established blood pressure loci on blood pressure values and hypertension risk in an Algerian population sample. Journal of Human Hypertension, 2015, 29, 296-302.	1.0	6
192	Combined effects of current-smoking and the aldehyde dehydrogenase 2*2 allele on the risk of myocardial infarction in Japanese patients. Toxicology Letters, 2015, 232, 221-225.	0.4	10
193	A Personalized Medicine Approach for Asian Americans with the Aldehyde Dehydrogenase 2*2 Variant. Annual Review of Pharmacology and Toxicology, 2015, 55, 107-127.	4.2	112
194	Genome-wide association study in Chinese identifies novel loci for blood pressure and hypertension. Human Molecular Genetics, 2015, 24, 865-874.	1.4	157
195	Chronic Kidney Disease and Hypertension. , 2015, , .		0
196	Epigenetics and arterial hypertension: the challenge of emerging evidence. Translational Research, 2015, 165, 154-165.	2.2	83
197	Interactions of Methylenetetrahydrofolate Reductase C677T Polymorphism with Environmental Factors on Hypertension Susceptibility. International Journal of Environmental Research and Public Health, 2016, 13, 601.	1.2	17
198	Cognitive and Physical Aging., 2016,, 125-146.		2
199	Gender-Specific Association of <i>ATP2B1 < /i>Variants with Susceptibility to Essential Hypertension in the Han Chinese Population. BioMed Research International, 2016, 2016, 1-7.</i>	0.9	8
200	Using the Generalized Index of Dissimilarity to Detect Gene-Gene Interactions in Multi-Class Phenotypes. PLoS ONE, 2016, 11, e0158668.	1.1	3
201	G = E: What GWAS Can Tell Us about the Environment. PLoS Genetics, 2016, 12, e1005765.	1.5	120
202	Gene Silencing and Haploinsufficiency of Csk Increase Blood Pressure. PLoS ONE, 2016, 11, e0146841.	1.1	16
203	Variant Discovery and Fine Mapping of Genetic Loci Associated with Blood Pressure Traits in Hispanics and African Americans. PLoS ONE, 2016, 11, e0164132.	1.1	24

#	ARTICLE	IF	Citations
204	Analysis and validation of traits associated with a single nucleotide polymorphism Gly364Ser in catestatin using humanized chromogranin A mouse models. Journal of Hypertension, 2016, 34, 68-78.	0.3	7
205	Recent Updates on the Proximal Tubule Renin-Angiotensin System in Angiotensin II-Dependent Hypertension. Current Hypertension Reports, 2016, 18, 63.	1.5	27
207	Recent Advances in the Genetics of Hypertension. Advances in Experimental Medicine and Biology, 2016, 956, 561-581.	0.8	17
208	Xanthine oxidase gene variants and their association with blood pressure and incident hypertension. Journal of Hypertension, 2016, 34, 2147-2154.	0.3	30
209	CNV analysis in Chinese children of mental retardation highlights a sex differentiation in parental contribution to de novo and inherited mutational burdens. Scientific Reports, 2016, 6, 25954.	1.6	19
210	Analysis of multiple related phenotypes in genome-wide association studies. Journal of Bioinformatics and Computational Biology, 2016, 14, 1644005.	0.3	5
211	Behavioural Genetics for Education. , 2016, , .		9
212	Mechanisms of Vascular Smooth Muscle Contraction and the Basis for Pharmacologic Treatment of Smooth Muscle Disorders. Pharmacological Reviews, 2016, 68, 476-532.	7.1	365
213	Molecular Genetic Investigations of Personality: From Candidate Genes to Genome-wide Associations. , 2016, , 130-154.		1
214	Variant Aldehyde Dehydrogenase 2 (<i>ALDH2*2</i>) Is a Risk Factor for Coronary Spasm and STâ€Segment Elevation Myocardial Infarction. Journal of the American Heart Association, 2016, 5, .	1.6	34
215	Heme oxygenase 1 protects ethanol-administered liver tissue in Aldh2 knockout mice. Alcohol, 2016, 52, 49-54.	0.8	12
216	Genetic Variation in the Human SORBS1 Gene is Associated With Blood Pressure Regulation and Age at Onset of Hypertension. Medicine (United States), 2016, 95, e2970.	0.4	12
217	Salt Sensitivity of Blood Pressure. Hypertension, 2016, 68, e7-e46.	1.3	347
218	Mendelian randomization analysis in three Japanese populations supports a causal role of alcohol consumption in lowering low-density lipid cholesterol levels and particle numbers. Atherosclerosis, 2016, 254, 242-248.	0.4	27
219	Roles of defective ALDH2 polymorphism on liver protection and cancer development. Environmental Health and Preventive Medicine, 2016, 21, 395-402.	1.4	20
220	Homozygous <i>ALDH2*2</i> Is an Independent Risk Factor for Ischemic Stroke in Taiwanese Men. Stroke, 2016, 47, 2174-2179.	1.0	38
221	The mitochondrial calcium uniporter is involved in mitochondrial calcium cycle dysfunction: Underlying mechanism of hypertension associated with mitochondrial tRNA lle A4263G mutation. International Journal of Biochemistry and Cell Biology, 2016, 78, 307-314.	1.2	16
222	Retinoblastoma-associated protein 140 as a candidate for a novel etiological gene to hypertension. Clinical and Experimental Hypertension, 2016, 38, 533-540.	0.5	5

#	Article	IF	CITATIONS
223	Association of Parental Hypertension With Arterial Stiffness in Nonhypertensive Offspring. Hypertension, 2016, 68, 584-589.	1.3	29
224	The Role of DNA Methylation in Hypertension. Advances in Experimental Medicine and Biology, 2016, 956, 583-598.	0.8	10
226	The genetics of blood pressure regulation and its target organs from association studies in 342,415 individuals. Nature Genetics, 2016, 48, 1171-1184.	9.4	362
227	Genome-wide association studies in East Asians identify new loci for waist-hip ratio and waist circumference. Scientific Reports, 2016, 6, 17958.	1.6	58
228	Genome Wide Association Study Identifies L3MBTL4 as a Novel Susceptibility Gene for Hypertension. Scientific Reports, 2016, 6, 30811.	1.6	15
229	Androgen-sensitive hypertension associated with soluble guanylate cyclase-α ₁ deficiency is mediated by 20-HETE. American Journal of Physiology - Heart and Circulatory Physiology, 2016, 310, H1790-H1800.	1.5	27
230	Assessing the Causality between Blood Pressure and Retinal Vascular Caliber through Mendelian Randomisation. Scientific Reports, 2016, 6, 22031.	1.6	5
231	Polymorphisms in the <i>SLC12A3</i> Gene Encoding Sodium-Chloride Cotransporter are Associated with Hypertension: A Family-Based Study in the Mongolian Population. Kidney and Blood Pressure Research, 2016, 41, 18-28.	0.9	7
232	Genome-wide association of trajectories of systolic blood pressure change. BMC Proceedings, 2016, 10, 321-327.	1.8	8
233	Comparison of 2 models for gene–environment interactions: an example of simulated gene–medication interactions on systolic blood pressure in family-based data. BMC Proceedings, 2016, 10, 371-377.	1.8	3
234	Zinc finger transcription factor Casz1 expression is regulated by homeodomain transcription factor Prrxl1 in embryonic spinal dorsal horn lateâ€born excitatory interneurons. European Journal of Neuroscience, 2016, 43, 1449-1459.	1.2	7
235	Genome-Wide Gene–Sodium Interaction Analyses on Blood Pressure. Hypertension, 2016, 68, 348-355.	1.3	44
236	The longitudinal effect of the aldehyde dehydrogenase 2*2 allele on the risk for nonalcoholic fatty liver disease. Nutrition and Diabetes, 2016, 6, e210-e210.	1.5	20
237	Evaluation of pleiotropic effects among common genetic loci identified for cardio-metabolic traits in a Korean population. Cardiovascular Diabetology, 2016, 15, 20.	2.7	20
238	Rare Exome Sequence Variants in <i>CLCN6</i> Reduce Blood Pressure Levels and Hypertension Risk. Circulation: Cardiovascular Genetics, 2016, 9, 64-70.	5.1	44
239	Associations between aldehyde dehydrogenase 2 (ALDH2) genetic polymorphisms, drinking status, and hypertension risk in Japanese adult male workers: a caseâ€"control study. Environmental Health and Preventive Medicine, 2016, 21, 1-8.	1.4	18
240	International Genome-Wide Association Study Consortium Identifies Novel Loci Associated With Blood Pressure in Children and Adolescents. Circulation: Cardiovascular Genetics, 2016, 9, 266-278.	5.1	48
241	Do You Want to Ditch Sodium? Meet Nitric Oxide Synthase $1\hat{l}^2$ at the Macula Densa. Journal of the American Society of Nephrology: JASN, 2016, 27, 2217-2218.	3.0	0

#	Article	IF	CITATIONS
242	Polymorphisms of three genes (<i>ACE</i> , <i>AGT</i> and <i>CYP11B2</i>) in the renin–angiotensin–aldosterone system are not associated with blood pressure salt sensitivity: A systematic meta-analysis. Blood Pressure, 2016, 25, 117-122.	0.7	8
243	CYP17A1 Enzyme Activity Is Linked to Ambulatory Blood Pressure in a Family-Based Population Study. American Journal of Hypertension, 2016, 29, 484-493.	1.0	13
244	Genome-wide association study of myelosuppression in non-small-cell lung cancer patients with platinum-based chemotherapy. Pharmacogenomics Journal, 2016, 16, 41-46.	0.9	27
245	Genome-wide association analysis identifies novel blood pressure loci and offers biological insights into cardiovascular risk. Nature Genetics, 2017, 49, 403-415.	9.4	492
246	Gut microbiota dysbiosis contributes to the development of hypertension. Microbiome, 2017, 5, 14.	4.9	1,086
247	Alcohol and coronary artery calcification: an investigation using alcohol flushing as an instrumental variable. International Journal of Epidemiology, 2017, 46, dyw237.	0.9	18
248	Increased NBCn1 expression, Na ⁺ /HCO ₃ ⁻ co-transport and intracellular pH in human vascular smooth muscle cells with a risk allele for hypertension. Human Molecular Genetics, 2017, 26, ddx015.	1.4	21
249	Vascular transcriptome profiling identifies Sphingosine kinase 1 as a modulator of angiotensin ll-induced vascular dysfunction. Scientific Reports, 2017, 7, 44131.	1.6	36
250	Are genetic polymorphisms in the renin–angiotensin–aldosterone system associated with essential hypertension? Evidence from genome-wide association studies. Journal of Human Hypertension, 2017, 31, 695-698.	1.0	39
251	Epigenetics and Arterial Hypertension. , 2017, , 159-184.		3
252	Complex inhibition of autophagy by mitochondrial aldehyde dehydrogenase shortens lifespan and exacerbates cardiac aging. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2017, 1863, 1919-1932.	1.8	81
253	Genomic Approaches to Hypertension. Cardiology Clinics, 2017, 35, 185-196.	0.9	12
254	Genomics of hypertension. Pharmacological Research, 2017, 121, 219-229.	3.1	17
255	Association of <i>NPR3 </i> polymorphism with risk of essential hypertension in a Chinese population. Journal of Clinical Pharmacy and Therapeutics, 2017, 42, 554-560.	0.7	3
256	Renal function of cyclin M2 Mg2+ transporter maintains blood pressure. Journal of Hypertension, 2017, 35, 585-592.	0.3	46
258	The Plasma Membrane Calcium ATP <scp>ases</scp> and Their Role as Major New Players in Human Disease. Physiological Reviews, 2017, 97, 1089-1125.	13.1	94
259	Associations Between Genetic Variants of NADPH Oxidase-Related Genes and Blood Pressure Responses to Dietary Sodium Intervention: The GenSalt Study. American Journal of Hypertension, 2017, 30, 427-434.	1.0	14
260	The ALDH2 gene rs671 polymorphism is not associated with essential hypertension. Clinical and Experimental Hypertension, 2017, 39, 691-695.	0.5	9

#	Article	IF	Citations
261	Rare coding variants associated with blood pressure variation in 15 914 individuals of African ancestry. Journal of Hypertension, 2017, 35, 1381-1389.	0.3	15
262	Genome-Wide Association Study Meta-Analysis of Long-Term Average Blood Pressure in East Asians. Circulation: Cardiovascular Genetics, 2017, 10, e001527.	5.1	26
263	A Review of the Genetics of Hypertension with a Focus on Gene-Environment Interactions. Current Hypertension Reports, 2017, 19, 23.	1.5	39
264	<i>ANTXR2</i> Knock-Out Does Not Result in the Development of Hypertension in Rats. American Journal of Hypertension, 2017, 30, 182-187.	1.0	8
265	Pharmacogenomics of the Natriuretic Peptide System in Heart Failure. Current Heart Failure Reports, 2017, 14, 536-542.	1.3	17
266	Blood Pressure Genetic Risk Score Predicts Blood Pressure Responses to Dietary Sodium and Potassium. Hypertension, 2017, 70, 1106-1112.	1.3	24
267	Genetics of NO Deficiency. American Journal of Cardiology, 2017, 120, S80-S88.	0.7	13
268	Genetic mechanisms of human hypertension and their implications for blood pressure physiology. Physiological Genomics, 2017, 49, 630-652.	1.0	32
269	Towards Precision Medicine for Hypertension: A Review of Genomic, Epigenomic, and Microbiomic Effects on Blood Pressure in Experimental Rat Models and Humans. Physiological Reviews, 2017, 97, 1469-1528.	13.1	85
270	Novel Blood Pressure Locus and Gene Discovery Using Genome-Wide Association Study and Expression Data Sets From Blood and the Kidney. Hypertension, 2017, 70, .	1.3	123
271	Dissecting the genetic components of a quantitative trait locus for blood pressure and renal pathology on rat chromosome 3. Journal of Hypertension, 2017, 35, 319-329.	0.3	6
272	DNA Methylation Analysis Identifies Loci for Blood Pressure Regulation. American Journal of Human Genetics, 2017, 101, 888-902.	2.6	154
273	Genome-Wide Gene–Potassium Interaction Analyses on Blood Pressure. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	9
274	Longitudinal exome-wide association study to identify genetic susceptibility loci for hypertension in a Japanese population. Experimental and Molecular Medicine, 2017, 49, e409-e409.	3.2	8
275	Differences in Natriuretic Peptide Levels by Race/Ethnicity (From the Multi-Ethnic Study of) Tj ETQq0 0 0 rgBT /Ov	verlock 10 0.7	Tf ₄₉ 0 182 Td
276	Genome-wide association analyses using electronic health records identify new loci influencing blood pressure variation. Nature Genetics, 2017, 49, 54-64.	9.4	281
277	A genome-wide association study of essential hypertension in an Australian population using a DNA pooling approach. Molecular Genetics and Genomics, 2017, 292, 307-324.	1.0	13
278	Targeting acetaldehyde dehydrogenase 2 (ALDH2) in heart failureâ€"Recent insights and perspectives. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2017, 1863, 1933-1941.	1.8	32

#	Article	IF	CITATIONS
279	Cohort Profile: The Korean Genome and Epidemiology Study (KoGES) Consortium. International Journal of Epidemiology, 2017, 46, e20-e20.	0.9	576
280	Gender-specific Association of ATP2B1 (rs2681472) Gene Polymorphism with Essential Hypertension in South Indian Population. International Journal of Human Genetics, 2017, 17, 169-176.	0.1	0
281	Meta-Analysis on the Association of ALDH2 Polymorphisms and Type 2 Diabetic Mellitus, Diabetic Retinopathy. International Journal of Environmental Research and Public Health, 2017, 14, 165.	1.2	22
282	ACTH and Polymorphisms at Steroidogenic Loci as Determinants of Aldosterone Secretion and Blood Pressure. International Journal of Molecular Sciences, 2017, 18, 579.	1.8	5
283	Urinary Exosomes and Their Cargo: Potential Biomarkers for Mineralocorticoid Arterial Hypertension?. Frontiers in Endocrinology, 2017, 8, 230.	1.5	36
284	Amyloid Î ² Modification: A Key to the Sporadic Alzheimer's Disease?. Frontiers in Genetics, 2017, 8, 58.	1.1	52
285	Positive association between ALDH2 rs671 polymorphism and essential hypertension: A case-control study and meta-analysis. PLoS ONE, 2017, 12, e0177023.	1.1	21
286	Association between kidney function and genetic polymorphisms in atherosclerotic and chronic kidney diseases: A cross-sectional study in Japanese male workers. PLoS ONE, 2017, 12, e0185476.	1.1	5
287	The fine-scale genetic structure and evolution of the Japanese population. PLoS ONE, 2017, 12, e0185487.	1.1	27
288	Development of Personal Data Handling Policy in Human Genome Research: a Historical Perspective in Japan. Asian Bioethics Review, 2017, 9, 183-197.	0.9	2
289	Mammalian Glutamyl Aminopeptidase Genes (ENPEP) and Proteins: Comparative Studies of a Major Contributor to Arterial Hypertension. Journal of Data Mining in Genomics & Proteomics, 2017, 08, .	0.5	25
290	Genetic polymorphisms in the <i>ALDH2</i> gene and the risk of ischemic stroke in a Chinese han population. Oncotarget, 2017, 8, 101936-101943.	0.8	11
291	Associations of NADPH oxidase-related genes with blood pressure changes and incident hypertension: The GenSalt Study. Journal of Human Hypertension, 2018, 32, 287-293.	1.0	11
292	Polymorphisms of genes involved in inflammation and blood vessel development influence the risk of varicose veins. Clinical Genetics, 2018, 94, 191-199.	1.0	15
293	Cohort Profile: The Singapore Multi-Ethnic Cohort (MEC) study. International Journal of Epidemiology, 2018, 47, 699-699j.	0.9	67
294	The potential of aldehyde dehydrogenase 2 as a therapeutic target in cardiovascular disease. Expert Opinion on Therapeutic Targets, 2018, 22, 217-231.	1.5	20
295	Genetics of Blood Pressure and Hypertension. Updates in Hypertension and Cardiovascular Protection, 2018, , 135-154.	0.1	0
297	Genetics and Genomics of Systemic Hypertension. , 2018, , 723-740.		O

#	Article	IF	CITATIONS
298	CrossTalk proposal: CNNM proteins are Na ⁺ /Mg ²⁺ exchangers playing a central role in transepithelial Mg ²⁺ (re)absorption. Journal of Physiology, 2018, 596, 743-746.	1.3	36
299	The PMCA pumps in genetically determined neuronal pathologies. Neuroscience Letters, 2018, 663, 2-11.	1.0	21
300	Genetic factors contributing to hypertension in Africanâ€based populations: A systematic review and metaâ€analysis. Journal of Clinical Hypertension, 2018, 20, 485-495.	1.0	21
301	Effect of dietary energy and polymorphisms in BRAP and GHRL on obesity and metabolic traits. Obesity Research and Clinical Practice, 2018, 12, 39-48.	0.8	22
302	Replication of a genome-wide association study on essential hypertension in Mongolians. Clinical and Experimental Hypertension, 2018, 40, 79-89.	0.5	1
303	The biological impact of blood pressure-associated genetic variants in the natriuretic peptide receptor C gene on human vascular smooth muscle. Human Molecular Genetics, 2018, 27, 199-210.	1.4	21
304	Regular aerobic exercise and blood pressure in East Asians: A meta-analysis of randomized controlled trials. Clinical and Experimental Hypertension, 2018, 40, 378-389.	0.5	30
305	Bevacizumab-induced hypertension: Clinical presentation and molecular understanding. , 2018, 182, 152-160.		99
307	Genome-Wide Association Studies of Hypertension and Several Other Cardiovascular Diseases. Pulse, 2018, 6, 169-186.	0.9	34
308	Csk Regulates Blood Pressure by Controlling the Synthetic Pathways of Aldosterone. Circulation Journal, 2018, 82, 168-175.	0.7	11
309	Interethnic analyses of blood pressure loci in populations of East Asian and European descent. Nature Communications, 2018, 9, 5052.	5.8	75
310	Characteristics and treatment of African-American and European-American patients with resistant hypertension identified using the electronic health record in an academic health centre: a caseâr control study. BMJ Open, 2018, 8, e021640.	0.8	15
311	Uncovering association networks through an eQTL analysis involving human miRNAs and lincRNAs. Scientific Reports, 2018, 8, 15050.	1.6	6
312	Vascular Smooth Muscle Remodeling in Conductive and Resistance Arteries in Hypertension. Arteriosclerosis, Thrombosis, and Vascular Biology, 2018, 38, 1969-1985.	1.1	157
313	Expanding the clinical relevance of the 5′-nucleotidase cN-II/NT5C2. Purinergic Signalling, 2018, 14, 321-329.	1.1	12
314	ATP2B1 rs2681472 and STK39 rs35929607 polymorphisms and risk of Hypertension in Iranian Population. Medical Journal of the Islamic Republic of Iran, 2018, 32, 78-82.	0.9	6
315	Genome-Wide Association Studies of Hypertension and Several Other Cardiovascular Diseases. Translational Bioinformatics, 2018, , 1-29.	0.0	3
316	Progress of Genomics in Hypertension–Cardiac Hypertrophy. Translational Bioinformatics, 2018, , 179-217.	0.0	0

#	Article	IF	CITATIONS
318	Gene-gene interactions and associations of six hypertension related single nucleotide polymorphisms with obesity risk in a Chinese children population. Gene, 2018, 679, 320-327.	1.0	18
319	Association between <i>ATP2B1</i> and <i>CACNB2</i> polymorphisms and high blood pressure in a population of Lithuanian children and adolescents: a cross-sectional study. BMJ Open, 2018, 8, e019902.	0.8	6
320	Identification of 13 novel susceptibility loci for early-onset myocardial infarction, hypertension, or chronic kidney disease. International Journal of Molecular Medicine, 2018, 42, 2415-2436.	1.8	33
321	Resistant Hypertension: Detection, Evaluation, and Management: A Scientific Statement From the American Heart Association. Hypertension, 2018, 72, e53-e90.	1.3	629
322	Contributions of rare coding variants in hypotension syndrome genes to population blood pressure variation. Medicine (United States), 2018, 97, e11865.	0.4	6
324	ALDH2 Protects Against Ischemic Stroke in Rats by Facilitating 4-HNE Clearance and AQP4 Down-Regulation. Neurochemical Research, 2018, 43, 1339-1347.	1.6	30
325	Brief Overview of a Decade of Genome-Wide Association Studies on Primary Hypertension. International Journal of Endocrinology, 2018, 2018, 1-14.	0.6	11
326	Integrative Bioinformatics Approaches for Identification of Drug Targets in Hypertension. Frontiers in Cardiovascular Medicine, 2018, 5, 25.	1.1	3
327	Modulation of the orthostatic blood pressure response by acute nitrate consumption is dependent upon ethnic origin. Clinical and Experimental Pharmacology and Physiology, 2018, 45, 1106-1117.	0.9	0
328	Reduced secretion of parathyroid hormone and hypocalcemia in systemic heterozygous ATP2B1-null hypertensive mice. Hypertension Research, 2018, 41, 699-707.	1.5	6
329	Generalization and fine mapping of red blood cell trait genetic associations to multiâ€ethnic populations: The PAGE study. American Journal of Hematology, 2018, 93, 1061-1073.	2.0	5
330	Common variant rs11191548 near the CYP17A1 gene is associated with hypertension and the serum 25(OH) D levels in Han Chinese. Journal of Human Genetics, 2018, 63, 731-737.	1.1	6
331	Synthesis, secretion, function, metabolism and application of natriuretic peptides in heart failure. Journal of Biological Engineering, 2018, 12, 2.	2.0	108
332	Omics of Blood Pressure and Hypertension. Circulation Research, 2018, 122, 1409-1419.	2.0	74
333	Genetics of Hypertension., 2018,, 52-59.		1
334	Applicability of Precision Medicine Approaches to Managing Hypertension in Rural Populations. Journal of Personalized Medicine, 2018, 8, 16.	1.1	5
335	Identification of nine genes as novel susceptibility loci for earlyâ€onset ischemic stroke, intracerebral hemorrhage, or subarachnoid hemorrhage. Biomedical Reports, 2018, 9, 8-20.	0.9	10
336	A decade in psychiatric GWAS research. Molecular Psychiatry, 2019, 24, 378-389.	4.1	78

#	Article	IF	Citations
337	Multivariate Cluster-Based Multifactor Dimensionality Reduction to Identify Genetic Interactions for Multiple Quantitative Phenotypes. BioMed Research International, 2019, 2019, 1-10.	0.9	6
338	Evolutionary history of diseaseâ€susceptibility loci identified in longitudinal exomeâ€wide association studies. Molecular Genetics & Enomic Medicine, 2019, 7, e925.	0.6	1
339	Over 1000 genetic loci influencing blood pressure with multiple systems and tissues implicated. Human Molecular Genetics, 2019, 28, R151-R161.	1.4	39
340	ALDH2 and Stroke: A Systematic Review of the Evidence. Advances in Experimental Medicine and Biology, 2019, 1193, 195-210.	0.8	17
341	The Role of ALDH2 in Sepsis and the To-Be-Discovered Mechanisms. Advances in Experimental Medicine and Biology, 2019, 1193, 175-194.	0.8	11
342	ALDH2 and Cardiovascular Disease. Advances in Experimental Medicine and Biology, 2019, 1193, 53-67.	0.8	40
343	The Bidirectional Effect of Defective ALDH2 Polymorphism and Disease Prevention. Advances in Experimental Medicine and Biology, 2019, 1193, 69-87.	0.8	17
344	Differential Analysis of Hypertension-Associated Intestinal Microbiota. International Journal of Medical Sciences, 2019, 16, 872-881.	1.1	91
345	A Prediction Model of Essential Hypertension Based on Genetic and Environmental Risk Factors in Northern Han Chinese. International Journal of Medical Sciences, 2019, 16, 793-799.	1.1	27
347	Characteristics of Gut Microbiota in Patients with Hypertension and/or Hyperlipidemia: A Cross-Sectional Study on Rural Residents in Xinxiang County, Henan Province. Microorganisms, 2019, 7, 399.	1.6	36
348	A genomeâ€wide association and replication study of blood pressure in Ugandan early adolescents. Molecular Genetics & Denomic Medicine, 2019, 7, e00950.	0.6	15
349	Familial aggregation and heritability of hypertension in Han population in Shanghai China: a case-control study. Clinical Hypertension, 2019, 25, 17.	0.7	7
350	Aldehyde dehydrogenase-2 as a therapeutic target. Expert Opinion on Therapeutic Targets, 2019, 23, 955-966.	1.5	37
351	Blood Pressure-Associated Genetic Variants in the Natriuretic Peptide Receptor 1 Gene Modulate Guanylate Cyclase Activity. Circulation Genomic and Precision Medicine, 2019, 12, e002472.	1.6	10
352	A Social Determinant of Health May Modify Genetic Associations for Blood Pressure: Evidence From a SNP by Education Interaction in an African American Population. Frontiers in Genetics, 2019, 10, 428.	1.1	7
353	Genetic linkage analysis of a large family identifies <i>FIGN</i> as a candidate modulator of reduced penetrance in heritable pulmonary arterial hypertension. Journal of Medical Genetics, 2019, 56, 481-490.	1.5	3
354	Generality of genomic findings on blood pressure traits and its usefulness in precision medicine in diverse populations: A systematic review. Clinical Genetics, 2019, 96, 17-27.	1.0	8
355	Identifying Multi-Omics Causers and Causal Pathways for Complex Traits. Frontiers in Genetics, 2019, 10, 110.	1.1	13

#	Article	IF	CITATIONS
356	The effect of body mass index and its interaction with family history on hypertension: a case–control study. Clinical Hypertension, 2019, 25, 6.	0.7	12
357	Shared genes between Alzheimer's disease and ischemic stroke. CNS Neuroscience and Therapeutics, 2019, 25, 855-864.	1.9	36
358	Epigenetic modification: a regulatory mechanism in essential hypertension. Hypertension Research, 2019, 42, 1099-1113.	1.5	57
359	Genetics of Hypertension and Heart Failure. Updates in Hypertension and Cardiovascular Protection, 2019, , 15-29.	0.1	0
360	Mitochondrial ALDH2 protects against lipopolysaccharide-induced myocardial contractile dysfunction by suppression of ER stress and autophagy. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2019, 1865, 1627-1641.	1.8	60
361	Effects of angiotensin converting enzyme gene polymorphism on hypertension in Africa: A meta-analysis and systematic review. PLoS ONE, 2019, 14, e0211054.	1.1	26
362	Gene–environment interactions related to blood pressure traits in two communityâ€based Korean cohorts. Genetic Epidemiology, 2019, 43, 402-413.	0.6	4
363	Genetic Variation in Steroid and Xenobiotic Metabolizing Pathways and Enterolactone Excretion Before and After Flaxseed Intervention in African American and European American Women. Cancer Epidemiology Biomarkers and Prevention, 2019, 28, 265-274.	1.1	9
364	Disorders of renal NaCl transport and implications for blood pressure regulation. Medizinische Genetik, 2019, 31, 1-7.	0.1	3
365	The Korea Biobank Array: Design and Identification of Coding Variants Associated with Blood Biochemical Traits. Scientific Reports, 2019, 9, 1382.	1.6	179
366	A new approach to identifying hypertension-associated genes in the mesenteric artery of spontaneously hypertensive rats and stroke-prone spontaneously hypertensive rats. Journal of Hypertension, 2019, 37, 1644-1656.	0.3	13
367	Trans-ethnic association study of blood pressure determinants in over 750,000 individuals. Nature Genetics, 2019, 51, 51-62.	9.4	328
368	Genetic Polymorphisms in Hypertension: Are We Missing the Immune Connection?. American Journal of Hypertension, 2019, 32, 113-122.	1.0	9
369	Pharmacogenomics of amlodipine and hydrochlorothiazide therapy and the quest for improved control of hypertension: a mini review. Heart Failure Reviews, 2019, 24, 343-357.	1.7	13
370	Genome-wide association and gene-environment interaction study identifies variants in ALDH2 associated with serum ferritin in a Chinese population. Gene, 2019, 685, 196-201.	1.0	6
371	Combined linkage and association analysis identifies rare and low frequency variants for blood pressure at 1q31. European Journal of Human Genetics, 2019, 27, 269-277.	1.4	5
372	Genome-wide meta-analysis of variant-by-diuretic interactions as modulators of lipid traits in persons of European and African ancestry. Pharmacogenomics Journal, 2020, 20, 482-493.	0.9	4
373	Systemic Adeno-Associated Virus-Mediated Gene Therapy Prevents the Multiorgan Disorders Associated with Aldehyde Dehydrogenase 2 Deficiency and Chronic Ethanol Ingestion. Human Gene Therapy, 2020, 31, 163-182.	1.4	6

#	Article	IF	CITATIONS
374	An association study of C9orf3, a novel component of the renin-angiotensin system, and hypertension in diabetes. Scientific Reports, 2020, 10, 16111.	1.6	5
375	Discovery of rare variants associated with blood pressure regulation through meta-analysis of 1.3 million individuals. Nature Genetics, 2020, 52, 1314-1332.	9.4	91
376	Genome-Wide and Candidate Gene Association Analyses Identify a 14-SNP Combination for Hypertension in Patients With Type 2 Diabetes. American Journal of Hypertension, 2020, 34, 651-661.	1.0	6
377	Association between the incidence of hypertension and alcohol consumption pattern and the alcohol flushing response: A 12-year follow-up study. Alcohol, 2020, 89, 43-48.	0.8	6
378	Birth weight associations with DNA methylation differences in an adult population. Epigenetics, 2021, 16, 783-796.	1.3	18
379	Impact of Gut Microbiome on Hypertensive Patients With Low-Salt Intake: Shika Study Results. Frontiers in Medicine, 2020, 7, 475.	1.2	8
380	Exploration of CYP21A2 and CYP17A1 polymorphisms and preeclampsia risk among Chinese Han population: a large-scale case-control study based on 5021 subjects. Human Genomics, 2020, 14, 33.	1.4	3
381	Integrative analysis identifies the association between CASZ1 methylation and ischemic stroke. Neurology: Genetics, 2020, 6, e509.	0.9	11
382	Prevention of aortic dissection and aneurysm via an ALDH2-mediated switch in vascular smooth muscle cell phenotype. European Heart Journal, 2020, 41, 2442-2453.	1.0	92
383	Identification of type 2 diabetes loci in 433,540 East Asian individuals. Nature, 2020, 582, 240-245.	13.7	282
384	Pharmacogenomics of Hypertension Treatment. International Journal of Molecular Sciences, 2020, 21, 4709.	1.8	40
385	Hypertension in the Dog and Cat. , 2020, , .		2
386	ALDH2 rs671 polymorphisms and the risk of cerebral microbleeds in Chinese elderly: the Taizhou Imaging Study. Annals of Translational Medicine, 2020, 8, 229-229.	0.7	4
387	CASZ1b is a novel transcriptional corepressor of mineralocorticoid receptor. Hypertension Research, 2021, 44, 407-416.	1.5	10
388	Lessons Learned from the Jackson Heart Study. Contemporary Cardiology, 2021, , 105-122.	0.0	0
389	Chromosome 2 Fragment Substitutions in Dahl Salt-Sensitive Rats and RNA Sequencing Identified Enpep and Hs2st1 as Vascular Inflammatory Modulators. Hypertension, 2021, 77, 178-189.	1.3	3
390	Membrane Transport Plasma Membrane Calcium Pump: Structure and Function., 2021, , 1063-1069.		0
391	Genome-Wide Association Study of Korean Asthmatics: A Comparison With UK Asthmatics. Allergy, Asthma and Immunology Research, 2021, 13, 609.	1.1	4

#	Article	IF	CITATIONS
392	The interaction on hypertension between family history and diabetes and other risk factors. Scientific Reports, 2021, 11, 4716.	1.6	25
393	Renal Natriuretic Peptide Receptor-C Deficiency Attenuates NaCl Cotransporter Activity in Angiotensin Il–Induced Hypertension. Hypertension, 2021, 77, 868-881.	1.3	8
394	Integrated metagenome and metabolome analyses of blood pressure studies in early postmenopausal Chinese women. Journal of Hypertension, 2021, 39, 1800-1809.	0.3	2
395	Aminopeptidase A (ENPEP) gene polymorphisms and preeclampsia: Descriptive analysis. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2021, 258, 70-74.	0.5	2
396	Hypertension Gene Risk Score in Diagnosis and Prediction of Complications. RUHS Journal of Health Sciences, 2021, 6, .	0.1	1
397	Concentrations of vanadium in urine with hypertension prevalence and blood pressure levels. Ecotoxicology and Environmental Safety, 2021, 213, 112028.	2.9	11
398	Characteristics and variation of fecal bacterial communities and functions in isolated systolic and diastolic hypertensive patients. BMC Microbiology, 2021, 21, 128.	1.3	9
399	Prospective study: Aldehyde dehydrogenaseÂ2 gene is associated with cardioâ€erebrovascular complications in typeÂ2 diabetes patients. Journal of Diabetes Investigation, 2021, 12, 1845-1854.	1.1	5
400	Hypertension in African Populations: Review and Computational Insights. Genes, 2021, 12, 532.	1.0	12
401	Progress in Defining the Genetic Contribution to Type 2 Diabetes in Individuals of East Asian Ancestry. Current Diabetes Reports, 2021, 21, 17.	1.7	5
402	ATP2B1 gene polymorphisms rs2681472 and rs17249754 are associated with susceptibility to hypertension and blood pressure levels. Medicine (United States), 2021, 100, e25530.	0.4	5
403	UPLC-MS-Based Serum Metabolomics Reveals Potential Biomarkers of Ang II-Induced Hypertension in Mice. Frontiers in Cardiovascular Medicine, 2021, 8, 683859.	1.1	1
404	Acetaldehyde Dehydrogenase 2 regulates HMG-CoA reductase stability and cholesterol synthesis in the liver. Redox Biology, 2021, 41, 101919.	3.9	27
405	Integrative genomic analysis of blood pressure and related phenotypes in rats. DMM Disease Models and Mechanisms, 2021, 14, .	1.2	6
406	<i>ALDH2</i> rs671 Is Associated With Elevated FPG, Reduced Glucose Clearance and Hepatic Insulin Resistance in Japanese Men. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e3573-e3581.	1.8	8
407	Hypertension genetics past, present and future applications. Journal of Internal Medicine, 2021, 290, 1130-1152.	2.7	20
408	Interaction between ALDH2 rs671 and life habits affects the risk of hypertension in Koreans. Medicine (United States), 2021, 100, e26664.	0.4	2
409	Effects of Isometric Handgrip Training on Home Blood Pressure Measurements in Hypertensive Patients: A Randomized Crossover Study. Internal Medicine, 2021, 60, 2181-2188.	0.3	4

#	Article	IF	CITATIONS
410	Lifestyle Score and Genetic Factors With Hypertension and Blood Pressure Among Adults in Rural China. Frontiers in Public Health, 2021, 9, 687174.	1.3	9
411	ATP2B1 genotypes rs2070759 and rs2681472 polymorphisms and risk of hypertension in Saudi population. Nucleosides, Nucleotides and Nucleic Acids, 2021, 40, 1075-1089.	0.4	2
412	The Aldehyde Dehydrogenase ALDH2*2 Allele, Associated with Alcohol Drinking Behavior, Dates Back to Prehistoric Times. Biomolecules, 2021, 11, 1376.	1.8	3
413	Identifying the predictive effectiveness of a genetic risk score for incident hypertension using machine learning methods among populations in rural China. Hypertension Research, 2021, 44, 1483-1491.	1.5	10
414	Heritability and genomeâ€wide association study of blood pressure in Chinese adult twins. Molecular Genetics & Chinese adult twins.	0.6	8
415	Molecular Diversity of Plasma Membrane Ca2+ Transporting ATPases: Their Function Under Normal and Pathological Conditions. Advances in Experimental Medicine and Biology, 2020, 1131, 93-129.	0.8	17
416	Epidemiology of Hypertension in Children. , 2016, , 1907-1950.		1
417	Multiancestry genome-wide association study of 520,000 subjects identifies 32 loci associated with stroke and stroke subtypes. Nature Genetics, 2018, 50, 524-537.	9.4	1,124
421	Genetic modifiers of hypertension in soluble guanylate cyclase α1–deficient mice. Journal of Clinical Investigation, 2012, 122, 2316-2325.	3.9	28
422	2018 Chinese Guidelines for Prevention and Treatment of Hypertension-A report of the Revision Committee of Chinese Guidelines for Prevention and Treatment of Hypertension. Journal of Geriatric Cardiology, 2019, 16, 182-241.	0.2	380
423	Rare variants in fox-1 homolog A (RBFOX1) are associated with lower blood pressure. PLoS Genetics, 2017, 13, e1006678.	1.5	18
424	Single-trait and multi-trait genome-wide association analyses identify novel loci for blood pressure in African-ancestry populations. PLoS Genetics, 2017, 13, e1006728.	1.5	88
425	Positional cloning of quantitative trait nucleotides for blood pressure and cardiac QT-interval by targeted CRISPR/Cas9 editing of a novel long non-coding RNA. PLoS Genetics, 2017, 13, e1006961.	1.5	26
426	Identification of IGF1, SLC4A4, WWOX, and SFMBT1 as Hypertension Susceptibility Genes in Han Chinese with a Genome-Wide Gene-Based Association Study. PLoS ONE, 2012, 7, e32907.	1.1	51
427	STK39 Polymorphism Is Associated with Essential Hypertension: A Systematic Review and Meta-Analysis. PLoS ONE, 2013, 8, e59584.	1.1	23
428	Prospective Associations of Coronary Heart Disease Loci in African Americans Using the MetaboChip: The PAGE Study. PLoS ONE, 2014, 9, e113203.	1.1	27
429	A Genome-Wide Association Study Uncovers a Genetic Locus Associated with Thoracic-to-Hip Ratio in Koreans. PLoS ONE, 2015, 10, e0145220.	1.1	10
430	Genome-wide study of resistant hypertension identified from electronic health records. PLoS ONE, 2017, 12, e0171745.	1.1	36

#	Article	IF	CITATIONS
431	Identification of polymorphisms in $12q24.1$, <i>ACAD10</i> , and <ibrap< i=""> as novel genetic determinants of blood pressure in Japanese by exome-wide association studies. Oncotarget, 2017, 8, 43068-43079.</ibrap<>	0.8	13
432	Could ALDH2*2 be the reason for low incidence and mortality of ovarian cancer for East Asia women?. Oncotarget, 2018, 9, 12503-12512.	0.8	7
433	Aldehyde dehydrogenase 2 inhibits inflammatory response and regulates atherosclerotic plaque. Oncotarget, 2016, 7, 35562-35576.	0.8	43
434	Association of a Missense ALDH2 Single Nucleotide Polymorphism (Glu504Lys) With Benign Prostate Hyperplasia in a Korean Population. International Neurourology Journal, 2013, 17, 168.	0.5	5
435	KAREBrowser: SNP database of Korea Association REsource Project. BMB Reports, 2012, 45, 47-50.	1,1	7
436	Replication of genome-wide association studies on asthma and allergic diseases in Korean adult population. BMB Reports, 2012, 45, 305-310.	1.1	10
437	EvoSNP-DB: A database of genetic diversity in East Asian populations. BMB Reports, 2013, 46, 416-421.	1.1	1
438	Genome-wide Survey of Copy Number Variants Associated with Blood Pressure and Body Mass Index in a Korean Population. Genomics and Informatics, 2011, 9, 152-160.	0.4	2
439	Decreases in <i>Casz1 </i> mRNA by an siRNA Complex Do not Alter Blood Pressure in Mice. Genomics and Informatics, 2012, 10, 40.	0.4	6
440	Association Analysis of Reactive Oxygen Species-Hypertension Genes Discovered by Literature Mining. Genomics and Informatics, 2012, 10, 244.	0.4	2
441	Genome-wide analyses of multiple obesity-related cytokines and hormones informs biology of cardiometabolic traits. Genome Medicine, 2021, 13, 156.	3.6	6
442	CYP17A1–ATP2B1 SNPs and Gene–Gene and Gene–Environment Interactions on Essential Hypertension. Frontiers in Cardiovascular Medicine, 2021, 8, 720884.	1.1	5
443	Meta-Analysis of Joint Test of SNP and SNP-Environment Interaction with Heterogeneity. Human Heredity, 2021, , 1-9.	0.4	1
444	Genome-wide association studies: Where we are heading?. World Journal of Medical Genetics, $2011, 1, 23$.	1.0	3
445	Genetics and Hypertension: Which Information for Clinical Practice., 2012,, 439-452.		0
447	Blood Pressure Genomics. , 2015, , 685-694.		0
450	Mechanism and Pathophysiology. Nephrology Self-assessment Program: NephSAP, 2020, 19, 43-57.	3.0	0
452	Current knowledge in hypertension genetics: mosaic theory, candidate genes and genome-wide association studies. Arterial Hypertension (Russian Federation), 2020, 26, 490-500.	0.1	1

#	Article	IF	CITATIONS
453	Genetics of Hypertension: The Human and Veterinary Perspectives. , 2020, , 145-168.		0
454	Effects of ATP2B1 Variants on the Systolic and Diastolic Blood Pressure according to the Degree of Obesity in the South Korean Population. Korean Journal of Clinical Laboratory Science, 2020, 52, 45-52.	0.1	0
455	A Genome-Wide Association Study of a Korean Population Identifies Genetic Susceptibility to Hypertension Based on Sex-Specific Differences. Genes, 2021, 12, 1804.	1.0	3
456	The alleles of AGT and HIF1A gene affect the risk of hypertension in plateau residents. Experimental Biology and Medicine, 2022, 247, 237-245.	1.1	1
457	Fetal Undernutrition Programming, Sympathetic Nerve Activity, and Arterial Hypertension Development. Frontiers in Physiology, 2021, 12, 704819.	1.3	10
458	Effects of Bushen-Jiangya granules on blood pressure and pharmacogenomic evaluation in low-to-medium-risk hypertensive patients: study protocol for a randomized double-blind controlled trial. Trials, 2022, 23, 37.	0.7	0
459	Human Gut Microbes Associated with Systolic Blood Pressure. International Journal of Hypertension, 2022, 2022, 1-12.	0.5	1
460	Additive Effects of Drinking Habits and a Susceptible Genetic Polymorphism on Cholesterol Efflux Capacity. Journal of Atherosclerosis and Thrombosis, 2023, 30, 23-38.	0.9	5
461	Lysine-specific demethylase 1 as a corepressor of mineralocorticoid receptor. Hypertension Research, 2022, 45, 641-649.	1.5	5
462	é'å°¿è,½å•–体C与å¿fè¡€ç®;疾病ç"ç©¶è¿å±•. Scientia Sinica Vitae, 2022, , .	0.1	0
463	Cataloging the potential SNPs (single nucleotide polymorphisms) associated with quantitative traits, viz. BMI (body mass index), IQ (intelligence quotient) and BP (blood pressure): an updated review. Egyptian Journal of Medical Human Genetics, 2022, 23, .	0.5	5
464	Macrophage ALDH2 (Aldehyde Dehydrogenase 2) Stabilizing Rac2 Is Required for Efferocytosis Internalization and Reduction of Atherosclerosis Development. Arteriosclerosis, Thrombosis, and Vascular Biology, 2022, 42, 700-716.	1.1	17
465	Aldehyde dehydrogenase 2 and PARP1 interaction modulates hepatic HDL biogenesis by LXRα-mediated ABCA1 expression. JCI Insight, 2022, 7, .	2.3	3
466	Shared genetic etiology and antagonistic relationship of plasma renin activity and systolic blood pressure in a Korean cohorts. Genomics, 2022, 114, 110334.	1.3	0
467	Association between pulse pressure, systolic blood pressure and the risk of rapid decline of kidney function among general population without hypertension: results from the China health and retirement longitudinal study (CHARLS). Journal of Translational Medicine, 2021, 19, 512.	1.8	8
479	Interactive effects of the lowâ€carbohydrate diet score and genetic risk score on Hypoâ€HDLâ€cholesterolemia among Korean adults: A crossâ€sectional analysis from the Ansan and Ansung Study of the Korean Genome and Epidemiology Study. Food Science and Nutrition, 2022, 10, 3106-3116.	1.5	1
480	Associations of Genome-Wide Polygenic Risk Score and Risk Factors With Hypertension in a Japanese Population. Circulation Genomic and Precision Medicine, 2022, 15, .	1.6	6
481	Kidney omics in hypertension: from statistical associations to biological mechanisms and clinical applications. Kidney International, 2022, 102, 492-505.	2.6	11

#	Article	IF	CITATIONS
482	A multi-ethnic polygenic risk score is associated with hypertension prevalence and progression throughout adulthood. Nature Communications, 2022, 13 , .	5.8	27
483	Insights From a Large-Scale Whole-Genome Sequencing Study of Systolic Blood Pressure, Diastolic Blood Pressure, and Hypertension. Hypertension, 2022, 79, 1656-1667.	1.3	12
484	Epigenetic and integrative cross-omics analyses of cerebral white matter hyperintensities on MRI. Brain, 2023, 146, 492-506.	3.7	12
485	The association of AGER and ALDH2 gene polymorphisms with diabetic retinopathy. European Journal of Ophthalmology, 0, , 112067212211262.	0.7	0
486	ALDH7A1 rs12514417 polymorphism may increase ischemic stroke risk in alcohol-exposed individuals. Nutrition and Metabolism, 2022, 19, .	1.3	2
487	The contribution of common and rare genetic variants to variation in metabolic traits in 288,137 East Asians. Nature Communications, 2022, 13, .	5.8	13
488	Association of ALDH2 Genotypes and Alcohol Intake with Dietary Patterns: The Bunkyo Health Study. Nutrients, 2022, 14, 4830.	1.7	1
489	CASZ1: a promising factor modulating aldosterone biosynthesis and mineralocorticoid receptor activity. Hypertension Research, 0 , , .	1.5	2
490	Identifying susceptibility genes for essential hypertension by transcriptome-wide association study. Biochemistry and Biophysics Reports, 2022, 32, 101387.	0.7	1
491	Aldehyde dehydrogenase in solid tumors and other diseases: Potential biomarkers and therapeutic targets. MedComm, 2023, 4, .	3.1	9
492	Meta-Analysis and Multivariate GWAS Analyses in 80,950 Individuals of African Ancestry Identify Novel Variants Associated with Blood Pressure Traits. International Journal of Molecular Sciences, 2023, 24, 2164.	1.8	0
493	The role of aldehyde dehydrogenase 2 in cardiovascular disease. Nature Reviews Cardiology, 2023, 20, 495-509.	6.1	16
494	A candidate locus in the renalase gene and susceptibility to blood pressure responses to the dietary salt. Journal of Hypertension, 2023, 41, 723-732.	0.3	1
495	Hypertension as a Novel Link for Shared Heritability in Age at Menarche and Cardiometabolic Traits. Journal of Clinical Endocrinology and Metabolism, 2023, 108, 2389-2399.	1.8	1
496	Cold-induced vasodilation response in a Japanese cohort: insights from cold-water immersion and genome-wide association studies. Journal of Physiological Anthropology, 2023, 42, .	1.0	0
498	Comparison of cardiovascular disease risk factors among FiLWHEL (2014–2016), NNS (2013) and KNHANES (2013–2015) women. BMC Women's Health, 2023, 23, .	0.8	0
499	Aldehyde Dehydrogenase 2 Ameliorates LPS-Induced Acute Kidney Injury through Detoxification of 4-HNE and Suppression of the MAPK Pathway. Journal of Immunology Research, 2023, 2023, 1-12.	0.9	2
500	Summary statistics-based association test for identifying the pleiotropic effects with set of genetic variants. Bioinformatics, 2023, 39, .	1.8	2

#	Article	IF	CITATIONS
507	ALDH2 polymorphism rs671 and alcohol consumption: possible explanatory factors for race/ethnic differences in bone density. Osteoporosis International, $0, \dots$	1.3	0
513	Genetics of Hypertension and Heart Failure. Updates in Hypertension and Cardiovascular Protection, 2023, , 37-51.	0.1	0