Robert Y L Zee

List of Publications by Year
in descending order

33	1,346	331538	414303
papers	citations	21 h-index	32 g-index

34	34	34	2587
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Gene Variation of Endoplasmic Reticulum Aminopeptidases 1 and 2, and Risk of Blood Pressure Progression and Incident Hypertension among 17,255 Initially Healthy Women. International Journal of Genomics, 2018, 2018, 1-9.	0.8	12
2	Islet amyloid polypeptide gene variation (IAPP) and the risk of incident type 2 diabetes mellitus: The women's genome health study. Clinica Chimica Acta, 2011, 412, 785-787.	0.5	3
3	Genetic variants in eleven telomere-associated genes and the risk of incident cardio/cerebrovascular disease: The Women's Genome Health Study. Clinica Chimica Acta, 2011, 412, 199-202.	0.5	23
4	Mitochondrial uncoupling protein gene cluster variation (UCP2â€“UCP3) and the risk of incident type 2 diabetes mellitus: The Women's Genome Health Study. Atherosclerosis, 2011, 214, 107-109.	0.4	12
5	Genetic variants of 11 telomere-pathway gene loci and the risk of incident type 2 diabetes mellitus: The Women's Genome Health Study. Atherosclerosis, 2011, 218, 144-146.	0.4	28
6	Mean leukocyte telomere length shortening and type 2 diabetes mellitus: a case-control study. Translational Research, 2010, 155, 166-169.	2.2	152
7	An Evaluation of Candidate Genes of Inflammation and Thrombosis in Relation to the Risk of Venous Thromboembolism. Circulation: Cardiovascular Genetics, 2009, 2, 57-62.	5.1	36
8	Mean Telomere Length and Risk of Incident Colorectal Carcinoma: A Prospective, Nested Case-Control Approach. Cancer Epidemiology Biomarkers and Prevention, 2009, 18, 2280-2282.	1.1	52
9	Genetic risk factors in recurrent venous thromboembolism: A multilocus, population-based, prospective approach. Clinica Chimica Acta, 2009, 402, 189-192.	0.5	18
10	Association of shorter mean telomere length with risk of incident myocardial infarction: A prospective, nested caseâ€“control approach. Clinica Chimica Acta, 2009, 403, 139-141.	0.5	77
11	Mean telomere length and risk of incident venous thromboembolism: A prospective, nested caseâ€“control approach. Clinica Chimica Acta, 2009, 406, 148-150.	0.5	6
12	Purinergic receptor P2Y, G-protein coupled, 12 gene variants and risk of incident ischemic stroke, myocardial infarction, and venous thromboembolism. Atherosclerosis, 2008, 197, 694-699.	0.4	40
13	Genetic variants within the interleukin-1 gene cluster, and risk of incident myocardial infarction, and ischemic stroke: A nested case-control approach. Atherosclerosis, 2008, 201, 124-129.	0.4	26
14	Association of reninâ€“angiotensin and endothelial nitric oxide synthase gene polymorphisms with blood pressure progression and incident hypertension: prospective cohort study. Journal of Hypertension, 2008, 26, 1780-1786.	0.3	37
15	Homocysteine, 5,10-Methylenetetrahydrofolate Reductase 677C>T Polymorphism, Nutrient Intake, and Incident Cardiovascular Disease in 24 968 Initially Healthy Women. Clinical Chemistry, 2007, 53, 845-851.	1.5	62
16	Natriuretic Peptide Precursor A Gene Polymorphisms and Risk of Blood Pressure Progression and Incident Hypertension. Hypertension, 2007, 50, 1114-1119.	1.3	33
17	Intercellular Adhesion Molecule 1 (ICAM1) Lys56Met and Gly241Arg Gene Variants, Plasma-Soluble ICAM1 Concentrations, and Risk of Incident Cardiovascular Events in 23 014 Initially Healthy White Women. Stroke, 2007, 38, 3152-3157.	1.0	12
18	Complement factor H Y402H gene polymorphism, C-reactive protein, and risk of incident myocardial infarction, ischaemic stroke, and venous thromboembolism: A nested caseâ€“control study. Atherosclerosis, 2006, 187, 332-335.	0.4	46

#	Article	IF	Citations
19	Genetic Variants of Arachidonate 5-Lipoxygenaseâ€“Activating Protein, and Risk of Incident Myocardial Infarction and Ischemic Stroke. Stroke, 2006, 37, 2007-2011.	1.0	95
20	Polymorphisms of the Phosphodiesterase 4D, cAMP-Specific (PDE4D) Gene and Risk of Ischemic Stroke. Stroke, 2006, 37, 2012-2017.	1.0	50
21	Polymorphism in the Î² 2 -Adrenergic Receptor and Lipoprotein Lipase Genes as Risk Determinants for Idiopathic Venous Thromboembolism. Circulation, 2006, 113, 2193-2200.	1.6	39
22	Haplotype Analysis of the Î²2 Adrenergic Receptor Gene and Risk of Myocardial Infarction in Humans. Genetics, 2005, 169, 1583-1587.	1.2	28
23	Toll-like Receptor 4 Asp299Gly Gene Polymorphism and Risk of Atherothrombosis. Stroke, 2005, 36, 154-157.	1.0	69
24	Tryptophanyl-tRNA synthetase gene polymorphisms and risk of incident myocardial infarction. Atherosclerosis, 2005, 181, 137-141.	0.4	2
25	Prospective Evaluation of the Alcohol Dehydrogenase Î³1/Î³2 Gene Polymorphism and Risk of Stroke. Stroke, 2004, 35, e39-42.	1.0	7
26	TP53 haplotype-based analysis and incidence of post-angioplasty restenosis. Human Genetics, 2004, 114, 386-390.	1.8	6
27	Threonine for alanine substitution in the eotaxin (CCL11) gene and the risk of incident myocardial infarction. Atherosclerosis, 2004, 175, 91-94.	0.4	49
28	C-reactive protein gene polymorphisms and the incidence of post-angioplasty restenosis. Atherosclerosis, 2004, 176, 393-396.	0.4	19
29	IL-1 cluster genes and occurrence of post-percutaneous transluminal coronary angioplasty restenosis: a prospective, angiography-based evaluation. Atherosclerosis, 2003, 171, 259-264.	0.4	9
30	Polymorphism in the P-selectin and interleukin-4 genes as determinants of stroke: a population-based, prospective genetic analysis. Human Molecular Genetics, 2003, 13, 389-396.	1.4	85
31	Polymorphism in the human C-reactive protein (CRP) gene, plasma concentrations of CRP, and the risk of future arterial thrombosis. Atherosclerosis, 2002, 162, 217-219.	0.4	191
32	A Prospective Evaluation of the CD14 and CD18 Gene Polymorphisms and Risk of Stroke. Stroke, 2002, 33, 892-895.	1.0	22
33	A prospective evaluation of the heat shock protein 70 gene polymorphisms and the risk of stroke. Thrombosis and Haemostasis, 2002, 87, 622-5.	1.8	0