

# Hatasu Kobayashi

## List of Publications by Year in Descending Order

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

63

papers

2,174

citations

26

h-index

46

g-index

66

ext. papers

2,557

ext. citations

5

avg, IF

4.17

L-index

#	Paper	IF	Citations
63	Mechanism of reactive oxygen species generation and oxidative DNA damage induced by acrylohydroxamic acid, a putative metabolite of acrylamide.. <i>Mutation Research - Genetic Toxicology and Environmental Mutagenesis</i> , <b>2022</b> , 873, 503420	3	0
62	Suppression of RNF213, a susceptibility gene for moyamoya disease, inhibits endoplasmic reticulum stress through SEL1L upregulation.. <i>Biochemical and Biophysical Research Communications</i> , <b>2022</b> , 609, 62-68	3.4	0
61	Characterization of Moyamoya and Middle Cerebral Artery Diseases by Carotid Canal Diameter and RNF213 p.R4810K Genotype.. <i>Journal of Stroke and Cerebrovascular Diseases</i> , <b>2022</b> , 31, 106481	2.8	
60	Copper-mediated DNA damage caused by purpurin, a natural anthraquinone.. <i>Genes and Environment</i> , <b>2022</b> , 44, 15	2.8	0
59	E44Q mutation in Na1.7 in a patient with infantile paroxysmal knee pain: electrophysiological analysis of voltage-dependent sodium current. <i>Heliyon</i> , <b>2021</b> , 7, e07396	3.6	0
58	RNF213 as a Susceptibility Gene for Moyamoya Disease has Multifunctional Roles in Biological Processes <b>2021</b> , 47-60		
57	CD44v9 Induces Stem Cell-Like Phenotypes in Human Cholangiocarcinoma. <i>Frontiers in Cell and Developmental Biology</i> , <b>2020</b> , 8, 417	5.7	4
56	Moyamoya disease patient mutations in the RING domain of RNF213 reduce its ubiquitin ligase activity and enhance NFB activation and apoptosis in an AAA+ domain-dependent manner. <i>Biochemical and Biophysical Research Communications</i> , <b>2020</b> , 525, 668-674	3.4	11
55	Comment on "Thyroid Cancer "Epidemic": A Socio-Environmental Health Problem Needs Collaborative Efforts". <i>Environmental Science &amp; Technology</i> , <b>2020</b> , 54, 9713-9714	10.3	1
54	Combination of RERG and ZNF671 methylation rates in circulating cell-free DNA: A novel biomarker for screening of nasopharyngeal carcinoma. <i>Cancer Science</i> , <b>2020</b> , 111, 2536-2545	6.9	4
53	Polyphenols with Anti-Amyloid Aggregation Show Potential Risk of Toxicity Via Pro-Oxidant Properties. <i>International Journal of Molecular Sciences</i> , <b>2020</b> , 21,	6.3	10
52	Proteomic analysis of the monkey hippocampus for elucidating ischemic resistance. <i>Journal of Clinical Biochemistry and Nutrition</i> , <b>2020</b> , 67, 167-173	3.1	1
51	Prevalence of RNF213 p.R4810K Variant in Early-Onset Stroke With Intracranial Arterial Stenosis. <i>Stroke</i> , <b>2019</b> , 50, 1561-1563	6.7	20
50	Mutant KCNJ3 and KCNJ5 Potassium Channels as Novel Molecular Targets in Bradyarrhythmias and Atrial Fibrillation. <i>Circulation</i> , <b>2019</b> , 139, 2157-2169	16.7	23
49	Moyamoya Disease Susceptibility Variant RNF213 p.R4810K Increases the Risk of Ischemic Stroke Attributable to Large-Artery Atherosclerosis. <i>Circulation</i> , <b>2019</b> , 139, 295-298	16.7	37
48	Dysregulation of RNF213 promotes cerebral hypoperfusion. <i>Scientific Reports</i> , <b>2018</b> , 8, 3607	4.9	22
47	Rare variants in RNF213, a susceptibility gene for moyamoya disease, are found in patients with pulmonary hypertension and aggravate hypoxia-induced pulmonary hypertension in mice. <i>Pulmonary Circulation</i> , <b>2018</b> , 8, 2045894018778155	2.7	20

46	Cell-specific overexpression of adiponectin receptor 1 does not improve diabetes mellitus in Akita mice. <i>PLoS ONE</i> , <b>2018</b> , 13, e0190863	3.7	5
45	Familial episodic limb pain in kindreds with novel Nav1.9 mutations. <i>PLoS ONE</i> , <b>2018</b> , 13, e0208516	3.7	6
44	RNF213 p.R4810K Variant and Intracranial Arterial Stenosis or Occlusion in Relatives of Patients with Moyamoya Disease. <i>Journal of Stroke and Cerebrovascular Diseases</i> , <b>2017</b> , 26, 1841-1847	2.8	18
43	Rapid contralateral progression of focal cerebral arteriopathy distinguished from RNF213-related moyamoya disease and fibromuscular dysplasia. <i>Child's Nervous System</i> , <b>2017</b> , 33, 1405-1409	1.7	3
42	Significant association of RNF213 p.R4810K, a moyamoya susceptibility variant, with coronary artery disease. <i>PLoS ONE</i> , <b>2017</b> , 12, e0175649	3.7	37
41	Pathological Investigation on RNF213: Animal Models Knockout and Transgenic Mice in Diabetes and Signal Transduction. <i>Current Topics in Environmental Health and Preventive Medicine</i> , <b>2017</b> , 69-78	0.3	
40	Significant Association of the RNF213 p.R4810K Polymorphism with Quasi-Moyamoya Disease. <i>Journal of Stroke and Cerebrovascular Diseases</i> , <b>2016</b> , 25, 2632-2636	2.8	22
39	Identification of MMP1 as a novel risk factor for intracranial aneurysms in ADPKD using iPSC models. <i>Scientific Reports</i> , <b>2016</b> , 6, 30013	4.9	26
38	RNF213 Is Associated with Intracranial Aneurysms in the French-Canadian Population. <i>American Journal of Human Genetics</i> , <b>2016</b> , 99, 1072-1085	11	38
37	A new horizon of moyamoya disease and associated health risks explored through RNF213. <i>Environmental Health and Preventive Medicine</i> , <b>2016</b> , 21, 55-70	4.2	69
36	Biological Monitoring of Human Exposure to Neonicotinoids Using Urine Samples, and Neonicotinoid Excretion Kinetics. <i>PLoS ONE</i> , <b>2016</b> , 11, e0146335	3.7	67
35	Infantile Pain Episodes Associated with Novel Nav1.9 Mutations in Familial Episodic Pain Syndrome in Japanese Families. <i>PLoS ONE</i> , <b>2016</b> , 11, e0154827	3.7	27
34	RNF213 Rare Variants in Slovakian and Czech Moyamoya Disease Patients. <i>PLoS ONE</i> , <b>2016</b> , 11, e0164759	3.7	26
33	Frequency of RNF213 p.R4810K, a susceptibility variant for moyamoya disease, and health characteristics of carriers in the Japanese population. <i>Environmental Health and Preventive Medicine</i> , <b>2016</b> , 21, 387-390	4.2	15
32	Genetic study of intracranial aneurysms. <i>Stroke</i> , <b>2015</b> , 46, 620-6	6.7	28
31	Whole-exome sequencing reveals genetic variants associated with chronic kidney disease characterized by tubulointerstitial damages in North Central Region, Sri Lanka. <i>Environmental Health and Preventive Medicine</i> , <b>2015</b> , 20, 354-9	4.2	19
30	Perfluorinated carboxylic acids discharged from the Yodo River Basin, Japan. <i>Chemosphere</i> , <b>2015</b> , 138, 81-8	8.4	12
29	Importance of molecular diagnosis in the accurate diagnosis of systemic carnitine deficiency. <i>Journal of Genetics</i> , <b>2015</b> , 94, 147-50	1.2	3

28	Early onset of moyamoya syndrome in a Down syndrome patient with the genetic variant RNF213 p.R4810K. <i>Brain and Development</i> , <b>2015</b> , 37, 822-4	2.2	10
27	Biochemical and Functional Characterization of RNF213 (Mysterin) R4810K, a Susceptibility Mutation of Moyamoya Disease, in Angiogenesis In Vitro and In Vivo. <i>Journal of the American Heart Association</i> , <b>2015</b> , 4,	6	72
26	Temporal trend and age-dependent serum concentration of phenolic organohalogen contaminants in Japanese men during 1989-2010. <i>Environmental Pollution</i> , <b>2014</b> , 185, 228-33	9.3	31
25	Proteomic analysis of carbonylated proteins in the monkey substantia nigra after ischemia-reperfusion. <i>Free Radical Research</i> , <b>2014</b> , 48, 694-705	4	15
24	An integrative study of the genetic, social and environmental determinants of chronic kidney disease characterized by tubulointerstitial damages in the North Central Region of Sri Lanka. <i>Journal of Occupational Health</i> , <b>2014</b> , 56, 28-38	2.3	73
23	P.R4810K, a polymorphism of RNF213, the susceptibility gene for moyamoya disease, is associated with blood pressure. <i>Environmental Health and Preventive Medicine</i> , <b>2013</b> , 18, 121-9	4.2	50
22	Downregulation of Securin by the variant RNF213 R4810K (rs112735431, G>A) reduces angiogenic activity of induced pluripotent stem cell-derived vascular endothelial cells from moyamoya patients. <i>Biochemical and Biophysical Research Communications</i> , <b>2013</b> , 438, 13-9	3.4	84
21	Genomewide association study identifies no major founder variant in Caucasian moyamoya disease. <i>Journal of Genetics</i> , <b>2013</b> , 92, 605-9	1.2	25
20	The moyamoya disease susceptibility variant RNF213 R4810K (rs112735431) induces genomic instability by mitotic abnormality. <i>Biochemical and Biophysical Research Communications</i> , <b>2013</b> , 439, 419-24	3.4	40
19	Ablation of Rnf213 retards progression of diabetes in the Akita mouse. <i>Biochemical and Biophysical Research Communications</i> , <b>2013</b> , 432, 519-25	3.4	40
18	Inflammation-related DNA damage and expression of CD133 and Oct3/4 in cholangiocarcinoma patients with poor prognosis. <i>Free Radical Biology and Medicine</i> , <b>2013</b> , 65, 1464-1472	7.8	44
17	Combined linkage analysis and exome sequencing identifies novel genes for familial goiter. <i>Journal of Human Genetics</i> , <b>2013</b> , 58, 366-77	4.3	16
16	Rapid progression of unilateral moyamoya disease in a patient with a family history and an RNF213 risk variant. <i>Cerebrovascular Diseases</i> , <b>2013</b> , 36, 155-7	3.2	30
15	Cognitive and affective impairments of a novel SCA/MND crossroad mutation Asidan. <i>European Journal of Neurology</i> , <b>2012</b> , 19, 1070-8	6	26
14	Clinical features of SCA36: a novel spinocerebellar ataxia with motor neuron involvement (Asidan). <i>Neurology</i> , <b>2012</b> , 79, 333-41	6.5	61
13	Preliminary assessment of ecological exposure of adult residents in Fukushima Prefecture to radioactive cesium through ingestion and inhalation. <i>Environmental Health and Preventive Medicine</i> , <b>2012</b> , 17, 292-8	4.2	55
12	Distribution of moyamoya disease susceptibility polymorphism p.R4810K in RNF213 in East and Southeast Asian populations. <i>Neurologia Medico-Chirurgica</i> , <b>2012</b> , 52, 299-303	2.6	84
11	Nitrative DNA damage and Oct3/4 expression in urinary bladder cancer with <i>Schistosoma haematobium</i> infection. <i>Biochemical and Biophysical Research Communications</i> , <b>2011</b> , 414, 344-9	3.4	41

10	Expansion of intronic GGCCTG hexanucleotide repeat in NOP56 causes SCA36, a type of spinocerebellar ataxia accompanied by motor neuron involvement. <i>American Journal of Human Genetics</i> , <b>2011</b> , 89, 121-30	11	193
9	Identification of RNF213 as a susceptibility gene for moyamoya disease and its possible role in vascular development. <i>PLoS ONE</i> , <b>2011</b> , 6, e22542	3-7	383
8	A rare Asian founder polymorphism of Raptor may explain the high prevalence of Moyamoya disease among East Asians and its low prevalence among Caucasians. <i>Environmental Health and Preventive Medicine</i> , <b>2010</b> , 15, 94-104	4-2	48
7	Suppressed phosphorylation of collapsin response mediator protein-2 in the hippocampus of HCNP precursor transgenic mice. <i>Brain Research</i> , <b>2010</b> , 1355, 180-8	3-7	5
6	Damage to cellular and isolated DNA induced by a metabolite of aspirin. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , <b>2009</b> , 661, 93-100	3-3	7
5	Proteomic identification of carbonylated proteins in the monkey hippocampus after ischemia-reperfusion. <i>Free Radical Biology and Medicine</i> , <b>2009</b> , 46, 1472-7	7-8	65
4	The mechanisms of oxidative DNA damage and apoptosis induced by norsalsolinol, an endogenous tetrahydroisoquinoline derivative associated with Parkinson's disease. <i>Journal of Neurochemistry</i> , <b>2009</b> , 108, 397-407	6	12
3	Mechanism of metal-mediated DNA damage and apoptosis induced by 6-hydroxydopamine in neuroblastoma SH-SY5Y cells. <i>Free Radical Research</i> , <b>2008</b> , 42, 651-60	4	17
2	Mechanism of DNA damage and apoptosis induced by tetrahydropapaveroline, a metabolite of dopamine. <i>Neurochemical Research</i> , <b>2006</b> , 31, 523-32	4-6	10
1	Metal-mediated oxidative damage to cellular and isolated DNA by gallic acid, a metabolite of antioxidant propyl gallate. <i>Mutation Research - Genetic Toxicology and Environmental Mutagenesis</i> , <b>2004</b> , 558, 111-20	3	61