

Hatasu Kobayashi

List of Publications by Citations

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

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 | Identification of RNF213 as a susceptibility gene for moyamoya disease and its possible role in vascular development. <i>PLoS ONE</i> , 2011 , 6, e22542 | 3.7 | 383 |
| 62 | 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> , 2011 , 89, 121-30 | 11 | 193 |
| 61 | 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> , 2013 , 438, 13-9 | 3.4 | 84 |
| 60 | Distribution of moyamoya disease susceptibility polymorphism p.R4810K in RNF213 in East and Southeast Asian populations. <i>Neurologia Medico-Chirurgica</i> , 2012 , 52, 299-303 | 2.6 | 84 |
| 59 | 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> , 2014 , 56, 28-38 | 2.3 | 73 |
| 58 | 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> , 2015 , 4, | 6 | 72 |
| 57 | A new horizon of moyamoya disease and associated health risks explored through RNF213. <i>Environmental Health and Preventive Medicine</i> , 2016 , 21, 55-70 | 4.2 | 69 |
| 56 | Biological Monitoring of Human Exposure to Neonicotinoids Using Urine Samples, and Neonicotinoid Excretion Kinetics. <i>PLoS ONE</i> , 2016 , 11, e0146335 | 3.7 | 67 |
| 55 | Proteomic identification of carbonylated proteins in the monkey hippocampus after ischemia-reperfusion. <i>Free Radical Biology and Medicine</i> , 2009 , 46, 1472-7 | 7.8 | 65 |
| 54 | Clinical features of SCA36: a novel spinocerebellar ataxia with motor neuron involvement (Asidan). <i>Neurology</i> , 2012 , 79, 333-41 | 6.5 | 61 |
| 53 | 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> , 2004 , 558, 111-20 | 3 | 61 |
| 52 | 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> , 2012 , 17, 292-8 | 4.2 | 55 |
| 51 | P.R4810K, a polymorphism of RNF213, the susceptibility gene for moyamoya disease, is associated with blood pressure. <i>Environmental Health and Preventive Medicine</i> , 2013 , 18, 121-9 | 4.2 | 50 |
| 50 | 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> , 2010 , 15, 94-104 | 4.2 | 48 |
| 49 | Inflammation-related DNA damage and expression of CD133 and Oct3/4 in cholangiocarcinoma patients with poor prognosis. <i>Free Radical Biology and Medicine</i> , 2013 , 65, 1464-1472 | 7.8 | 44 |
| 48 | Nitrative DNA damage and Oct3/4 expression in urinary bladder cancer with <i>Schistosoma haematobium</i> infection. <i>Biochemical and Biophysical Research Communications</i> , 2011 , 414, 344-9 | 3.4 | 41 |
| 47 | The moyamoya disease susceptibility variant RNF213 R4810K (rs112735431) induces genomic instability by mitotic abnormality. <i>Biochemical and Biophysical Research Communications</i> , 2013 , 439, 419-26 | 3.4 | 40 |

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| 46 | Ablation of Rnf213 retards progression of diabetes in the Akita mouse. <i>Biochemical and Biophysical Research Communications</i> , 2013 , 432, 519-25 | 3.4 | 40 |
| 45 | RNF213 Is Associated with Intracranial Aneurysms in the French-Canadian Population. <i>American Journal of Human Genetics</i> , 2016 , 99, 1072-1085 | 11 | 38 |
| 44 | Significant association of RNF213 p.R4810K, a moyamoya susceptibility variant, with coronary artery disease. <i>PLoS ONE</i> , 2017 , 12, e0175649 | 3.7 | 37 |
| 43 | Moyamoya Disease Susceptibility Variant RNF213 p.R4810K Increases the Risk of Ischemic Stroke Attributable to Large-Artery Atherosclerosis. <i>Circulation</i> , 2019 , 139, 295-298 | 16.7 | 37 |
| 42 | Temporal trend and age-dependent serum concentration of phenolic organohalogen contaminants in Japanese men during 1989-2010. <i>Environmental Pollution</i> , 2014 , 185, 228-33 | 9.3 | 31 |
| 41 | Rapid progression of unilateral moyamoya disease in a patient with a family history and an RNF213 risk variant. <i>Cerebrovascular Diseases</i> , 2013 , 36, 155-7 | 3.2 | 30 |
| 40 | Genetic study of intracranial aneurysms. <i>Stroke</i> , 2015 , 46, 620-6 | 6.7 | 28 |
| 39 | Infantile Pain Episodes Associated with Novel Nav1.9 Mutations in Familial Episodic Pain Syndrome in Japanese Families. <i>PLoS ONE</i> , 2016 , 11, e0154827 | 3.7 | 27 |
| 38 | Identification of MMP1 as a novel risk factor for intracranial aneurysms in ADPKD using iPSC models. <i>Scientific Reports</i> , 2016 , 6, 30013 | 4.9 | 26 |
| 37 | Cognitive and affective impairments of a novel SCA/MND crossroad mutation Asidan. <i>European Journal of Neurology</i> , 2012 , 19, 1070-8 | 6 | 26 |
| 36 | RNF213 Rare Variants in Slovakian and Czech Moyamoya Disease Patients. <i>PLoS ONE</i> , 2016 , 11, e0164759 | 9.7 | 26 |
| 35 | Genomewide association study identifies no major founder variant in Caucasian moyamoya disease. <i>Journal of Genetics</i> , 2013 , 92, 605-9 | 1.2 | 25 |
| 34 | Mutant KCNJ3 and KCNJ5 Potassium Channels as Novel Molecular Targets in Bradyarrhythmias and Atrial Fibrillation. <i>Circulation</i> , 2019 , 139, 2157-2169 | 16.7 | 23 |
| 33 | Dysregulation of RNF213 promotes cerebral hypoperfusion. <i>Scientific Reports</i> , 2018 , 8, 3607 | 4.9 | 22 |
| 32 | Significant Association of the RNF213 p.R4810K Polymorphism with Quasi-Moyamoya Disease. <i>Journal of Stroke and Cerebrovascular Diseases</i> , 2016 , 25, 2632-2636 | 2.8 | 22 |
| 31 | Prevalence of RNF213 p.R4810K Variant in Early-Onset Stroke With Intracranial Arterial Stenosis. <i>Stroke</i> , 2019 , 50, 1561-1563 | 6.7 | 20 |
| 30 | 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> , 2018 , 8, 2045894018778155 | 2.7 | 20 |
| 29 | 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> , 2015 , 20, 354-9 | 4.2 | 19 |

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| 28 | 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> , 2017 , 26, 1841-1847 | 2.8 | 18 |
| 27 | Mechanism of metal-mediated DNA damage and apoptosis induced by 6-hydroxydopamine in neuroblastoma SH-SY5Y cells. <i>Free Radical Research</i> , 2008 , 42, 651-60 | 4 | 17 |
| 26 | Combined linkage analysis and exome sequencing identifies novel genes for familial goiter. <i>Journal of Human Genetics</i> , 2013 , 58, 366-77 | 4.3 | 16 |
| 25 | Proteomic analysis of carbonylated proteins in the monkey substantia nigra after ischemia-reperfusion. <i>Free Radical Research</i> , 2014 , 48, 694-705 | 4 | 15 |
| 24 | 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> , 2016 , 21, 387-390 | 4.2 | 15 |
| 23 | Perfluorinated carboxylic acids discharged from the Yodo River Basin, Japan. <i>Chemosphere</i> , 2015 , 138, 81-8 | 8.4 | 12 |
| 22 | 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> , 2009 , 108, 397-407 | 6 | 12 |
| 21 | 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> , 2020 , 525, 668-674 | 3.4 | 11 |
| 20 | Early onset of moyamoya syndrome in a Down syndrome patient with the genetic variant RNF213 p.R4810K. <i>Brain and Development</i> , 2015 , 37, 822-4 | 2.2 | 10 |
| 19 | Mechanism of DNA damage and apoptosis induced by tetrahydropapaveroline, a metabolite of dopamine. <i>Neurochemical Research</i> , 2006 , 31, 523-32 | 4.6 | 10 |
| 18 | Polyphenols with Anti-Amyloid β Aggregation Show Potential Risk of Toxicity Via Pro-Oxidant Properties. <i>International Journal of Molecular Sciences</i> , 2020 , 21, | 6.3 | 10 |
| 17 | Damage to cellular and isolated DNA induced by a metabolite of aspirin. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2009 , 661, 93-100 | 3.3 | 7 |
| 16 | Familial episodic limb pain in kindreds with novel Nav1.9 mutations. <i>PLoS ONE</i> , 2018 , 13, e0208516 | 3.7 | 6 |
| 15 | Suppressed phosphorylation of collapsin response mediator protein-2 in the hippocampus of HCNP precursor transgenic mice. <i>Brain Research</i> , 2010 , 1355, 180-8 | 3.7 | 5 |
| 14 | Cell-specific overexpression of adiponectin receptor 1 does not improve diabetes mellitus in Akita mice. <i>PLoS ONE</i> , 2018 , 13, e0190863 | 3.7 | 5 |
| 13 | CD44v9 Induces Stem Cell-Like Phenotypes in Human Cholangiocarcinoma. <i>Frontiers in Cell and Developmental Biology</i> , 2020 , 8, 417 | 5.7 | 4 |
| 12 | Combination of RERG and ZNF671 methylation rates in circulating cell-free DNA: A novel biomarker for screening of nasopharyngeal carcinoma. <i>Cancer Science</i> , 2020 , 111, 2536-2545 | 6.9 | 4 |
| 11 | Rapid contralateral progression of focal cerebral arteriopathy distinguished from RNF213-related moyamoya disease and fibromuscular dysplasia. <i>Child's Nervous System</i> , 2017 , 33, 1405-1409 | 1.7 | 3 |

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| 10 | Importance of molecular diagnosis in the accurate diagnosis of systemic carnitine deficiency. <i>Journal of Genetics</i> , 2015 , 94, 147-50 | 1.2 | 3 |
| 9 | Comment on "Thyroid Cancer "Epidemic": A Socio-Environmental Health Problem Needs Collaborative Efforts". <i>Environmental Science & Technology</i> , 2020 , 54, 9713-9714 | 10.3 | 1 |
| 8 | Proteomic analysis of the monkey hippocampus for elucidating ischemic resistance. <i>Journal of Clinical Biochemistry and Nutrition</i> , 2020 , 67, 167-173 | 3.1 | 1 |
| 7 | 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> , 2022 , 873, 503420 | 3 | 0 |
| 6 | E44Q mutation in Na1.7 in a patient with infantile paroxysmal knee pain: electrophysiological analysis of voltage-dependent sodium current. <i>Heliyon</i> , 2021 , 7, e07396 | 3.6 | 0 |
| 5 | Suppression of RNF213, a susceptibility gene for moyamoya disease, inhibits endoplasmic reticulum stress through SEL1L upregulation.. <i>Biochemical and Biophysical Research Communications</i> , 2022 , 609, 62-68 | 3.4 | 0 |
| 4 | Copper-mediated DNA damage caused by purpurin, a natural anthraquinone.. <i>Genes and Environment</i> , 2022 , 44, 15 | 2.8 | 0 |
| 3 | 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> , 2017 , 69-78 | 0.3 | |
| 2 | RNF213 as a Susceptibility Gene for Moyamoya Disease has Multifunctional Roles in Biological Processes 2021 , 47-60 | | |
| 1 | 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> , 2022 , 31, 106481 | 2.8 | |