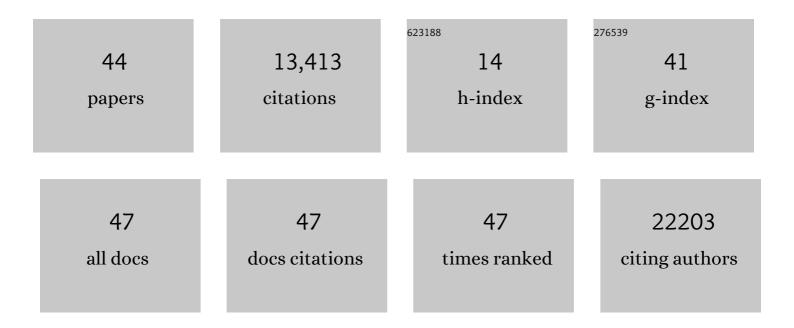
Kazuharu Misawa

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	MAFFT: a novel method for rapid multiple sequence alignment based on fast Fourier transform. Nucleic Acids Research, 2002, 30, 3059-3066.	6.5	12,537
2	Molecular systematics of Volvocales (Chlorophyceae, Chlorophyta) based on exhaustive 18S rRNA phylogenetic analyses. Molecular Phylogenetics and Evolution, 2008, 48, 281-291.	1.2	160
3	Origin and Evolution of the Colonial Volvocales (Chlorophyceae) as Inferred from Multiple, Chloroplast Gene Sequences. Molecular Phylogenetics and Evolution, 2000, 17, 256-268.	1.2	135
4	Phylogenetic positions of Glaucophyta, green plants (Archaeplastida) and Haptophyta (Chromalveolata) as deduced from slowly evolving nuclear genes. Molecular Phylogenetics and Evolution, 2009, 53, 872-880.	1.2	62
5	Phylogeny of Primary Photosynthetic Eukaryotes as Deduced from Slowly Evolving Nuclear Genes. Molecular Biology and Evolution, 2007, 24, 1592-1595.	3.5	55
6	A multi-ethnic meta-analysis identifies novel genes, including ACSL5, associated with amyotrophic lateral sclerosis. Communications Biology, 2020, 3, 526.	2.0	49
7	Revisiting the Glires concept—phylogenetic analysis of nuclear sequences. Molecular Phylogenetics and Evolution, 2003, 28, 320-327.	1.2	38
8	Cyanobacterial contribution to the genomes of the plastid-lacking protists. BMC Evolutionary Biology, 2009, 9, 197.	3.2	38
9	Reanalysis of Murphy et al.?s Data Gives Various Mammalian Phylogenies and Suggests Overcredibility of Bayesian Trees. Journal of Molecular Evolution, 2003, 57, S290-S296.	0.8	36
10	Genomic structure of nitric oxide synthase in the terrestrial slug is highly conserved. Gene, 2008, 415, 74-81.	1.0	28
11	The Universal Trend of Amino Acid Gain–Loss is Caused by CpG Hypermutability. Journal of Molecular Evolution, 2008, 67, 334-342.	0.8	27
12	Estimation of the Amount of DNA Polymorphism When the Neutral Mutation Rate Varies Among Sites. Genetics, 1997, 147, 1959-1964.	1.2	27
13	Evaluation of the effect of CpG hypermutability on human codon substitution. Gene, 2009, 431, 18-22.	1.0	19
14	Hypouricemia and Urate Transporters. Biomedicines, 2022, 10, 652.	1.4	17
15	Origins of a cyanobacterial 6-phosphogluconate dehydrogenase in plastid-lacking eukaryotes. BMC Evolutionary Biology, 2008, 8, 151.	3.2	16
16	STR-realigner: a realignment method for short tandem repeat regions. BMC Genomics, 2016, 17, 991.	1.2	15
17	Contribution of Rare Variants of the <i>SLC22A12</i> Gene to the Missing Heritability of Serum Urate Levels. Genetics, 2020, 214, 1079-1090.	1.2	15
18	Effect of Group Selection on the Evolution of Altruistic Behavior. Journal of Theoretical Biology, 2003, 220, 55-66.	0.8	13

Kazuharu Misawa

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19	Construction of full-length Japanese reference panel of class I HLA genes with single-molecule, real-time sequencing. Pharmacogenomics Journal, 2019, 19, 136-146.	0.9	12
20	Relationship between amino acid composition and gene expression in the mouse genome. BMC Research Notes, 2011, 4, 20.	0.6	11
21	A codon substitution model that incorporates the effect of the GC contents, the gene density and the density of CpG islands of human chromosomes. BMC Genomics, 2011, 12, 397.	1.2	10
22	RF: A method for filtering short reads with tandem repeats for genome mapping. Genomics, 2013, 102, 35-37.	1.3	9
23	Construction of JRG (Japanese reference genome) with single-molecule real-time sequencing. Human Genome Variation, 2019, 6, 27.	0.4	9
24	A Simple Method for Classifying Genes and a Bootstrap Test for Classifications. Molecular Biology and Evolution, 2000, 17, 1879-1884.	3.5	8
25	AP-SKAT: highly-efficient genome-wide rare variant association test. BMC Genomics, 2016, 17, 745.	1.2	8
26	Influence of the 3'-UTR-length of mKIAA cDNAs and their Sequence Features to the mRNA Expression Level in the Brain. DNA Research, 2005, 12, 181-189.	1.5	7
27	New correction algorithms for multiple comparisons in case–control multilocus association studies based on haplotypes and diplotype configurations. Journal of Human Genetics, 2008, 53, 789-801.	1.1	7
28	GeneWaltz–A new method for reducing the false positives of gene finding. BioData Mining, 2010, 3, 6.	2.2	7
29	The amount and pattern of DNA polymorphism under the neutral mutation hypothesis. Genetica, 1998, 102/103, 103-107.	0.5	6
30	ParaHaplo: A program package for haplotype-based whole-genome association study using parallel computing. Source Code for Biology and Medicine, 2009, 4, 7.	1.7	5
31	AQUATIC PLANT SPECIATION AFFECTED BY DIVERSIFYING SELECTION OF ORGANELLE DNA REGIONS ¹ . Journal of Phycology, 2011, 47, 999-1008.	1.0	5
32	The amount and pattern of DNA polymorphism under the neutral mutation hypothesis. Contemporary Issues in Genetics and Evolution, 1998, , 103-107.	0.9	5
33	ParaHaplo 3.0: A program package for imputation and a haplotype-based whole-genome association study using hybrid parallel computing. Source Code for Biology and Medicine, 2011, 6, 10.	1.7	3
34	Genomic Heritabilities and Correlations of 17 Traits Related to Obesity and Associated Conditions in the Japanese Population. G3: Genes, Genomes, Genetics, 2020, 10, 2221-2228.	0.8	3
35	Antiemetic effects of baclofen in a shrew model of postoperative nausea and vomiting: Wholeâ€ŧranscriptome analysis in the nucleus of the solitary tract. CNS Neuroscience and Therapeutics, 2022, , .	1.9	3
36	New Weighting Methods for Phylogenetic Tree Reconstruction Using Multiple Loci. Journal of Molecular Evolution, 2012, 75, 1-10.	0.8	2

Kazuharu Misawa

#	Article	IF	CITATIONS
37	Multiple Sequence Alignments: The Next Generation. Seibutsu Butsuri, 2006, 46, 312-317.	0.0	1
38	ParaHaplo 2.0: a program package for haplotype-estimation and haplotype-based whole-genome association study using parallel computing. Source Code for Biology and Medicine, 2010, 5, 5.	1.7	1
39	PAFFT: A new homology search algorithm for third-generation sequencers. Genomics, 2015, 106, 265-267.	1.3	1
40	Short tandem repeats in the human, cow, mouse, chicken, and lizard genomes are concentrated in the terminal regions of chromosomes. Gene Reports, 2016, 4, 280-285.	0.4	1
41	Population-scale whole genome sequencing identifies 271 highly polymorphic short tandem repeats from Japanese population. Heliyon, 2018, 4, e00625.	1.4	1
42	Is the population of Sado Island genetically close to the population of western Japan?. Human Genome Variation, 2019, 6, 26.	0.4	0
43	Estimating Human Point Mutation Rates from Codon Substitution Rates. , 0, , .		0
44	Genotype Value Decomposition: Simple Methods for the Computation of Kernel Statistics. Genetics & Genomics Next, 0, , 2100066.	0.8	0