## Vinh S Le

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6731346/publications.pdf

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25 6,452 8 17
papers citations h-index g-index

26 26 26 9623 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	UFBoot2: Improving the Ultrafast Bootstrap Approximation. Molecular Biology and Evolution, 2018, 35, 518-522.	3.5	5,798
2	POY version 4: phylogenetic analysis using dynamic homologies. Cladistics, 2010, 26, 72-85.	1.5	186
3	IQPNNI: Moving Fast Through Tree Space and Stopping in Time. Molecular Biology and Evolution, 2004, 21, 1565-1571.	3.5	146
4	MPBoot: fast phylogenetic maximum parsimony tree inference and bootstrap approximation. BMC Evolutionary Biology, 2018, 18, 11.	3.2	129
5	FLU, an amino acid substitution model for influenza proteins. BMC Evolutionary Biology, 2010, 10, 99.	3.2	53
6	QMaker: Fast and Accurate Method to Estimate Empirical Models of Protein Evolution. Systematic Biology, 2021, 70, 1046-1060.	2.7	39
7	A Vietnamese human genetic variation database. Human Mutation, 2019, 40, 1664-1675.	1.1	36
8	FastMG: a simple, fast, and accurate maximum likelihood procedure to estimate amino acid replacement rate matrices from large data sets. BMC Bioinformatics, 2014, 15, 341.	1.2	16
9	nQMaker: Estimating Time Nonreversible Amino Acid Substitution Models. Systematic Biology, 2022, 71, 1110-1123.	2.7	9
10	FLAVI: An Amino Acid Substitution Model for Flaviviruses. Journal of Molecular Evolution, 2020, 88, 445-452.	0.8	8
11	Pairwise alignment with rearrangements. Genome Informatics, 2006, 17, 141-51.	0.4	8
12	Random Tree-Puzzle Leads to the Yule–Harding Distribution. Molecular Biology and Evolution, 2011, 28, 873-877.	3.5	6
13	A novel de�1/2novo variant of LAMA2 contributes to merosin deficient congenital muscular dystrophy type 1A: Case report. Biomedical Reports, 2020, 12, 46-50.	0.9	5
14	Whole genome analysis of a Vietnamese trio. Journal of Biosciences, 2015, 40, 113-124.	0.5	4
15	mPartition: A Model-Based Method for Partitioning Alignments. Journal of Molecular Evolution, 2020, 88, 641-652.	0.8	3
16	Response to: A commentary on "A Vietnamese human genetic variation database― Human Mutation, 2020, 41, 1461-1462.	1.1	2
17	De novo homozygous variant of the SCN1A gene in a patient with severe Dravet syndrome complicated by acute encephalopathy. Neurogenetics, 2021, 22, 133-136.	0.7	2
18	A Hybrid Approach to Optimize the Number of Recombinations in Ancestral Recombination Graphs. , 2019, , .		1

#	Article	IF	CITATIONS
19	pQMaker: empirically estimating amino acid substitution models in a parallel environment. , 2020, , .		1
20	A Fast and Efficient Method for Estimating Amino Acid Substitution Models. , 2011, , .		0
21	MVRM: A Hybrid Approach to Predict siRNA Efficacy. , 2015, , .		O
22	Genomedics: Whole exome analysis system for clinical studies. , 2017, , .		0
23	Building minimum recombination ancestral recombination graphs for whole genomes. , 2017, , .		O
24	Exploring the Kinh Vietnamese genomic database for the polymorphisms of the P450 genes towards precision public health. Annals of Human Biology, 2022, 49, 152-155.	0.4	0
25	Response to: A commentary on "A Vietnamese human genetic variation database― , 0, , .		0