

Laura Kytvuori

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.

20 papers	110 citations	5 h-index	10 g-index
22 ext. papers	152 ext. citations	4 avg, IF	2.24 L-index

#	Paper	IF	Citations
20	Biallelic expansion in RFC1 as a rare cause of Parkinson's disease.. <i>Npj Parkinson's Disease</i> , 2022 , 8, 6	9.7	3
19	Molecular epidemiology of hereditary ataxia in Finland. <i>BMC Neurology</i> , 2021 , 21, 382	3.1	2
18	Association of mitochondrial DNA haplogroups J and K with low response in exercise training among Finnish military conscripts. <i>BMC Genomics</i> , 2021 , 22, 75	4.5	0
17	Mutation Analysis of the Genes Associated with Parkinson's Disease in a Finnish Cohort of Early-Onset Dementia. <i>Journal of Alzheimer's Disease</i> , 2020 , 76, 955-965	4.3	
16	Mitochondrial DNA variation in sudden cardiac death: a population-based study. <i>International Journal of Legal Medicine</i> , 2020 , 134, 39-44	3.1	1
15	Mutation Analysis of the Genes Linked to Early Onset Alzheimer's Disease and Frontotemporal Lobar Degeneration. <i>Journal of Alzheimer's Disease</i> , 2019 , 69, 775-782	4.3	2
14	Analysis of functional variants in mitochondrial DNA of Finnish athletes. <i>BMC Genomics</i> , 2019 , 20, 784	4.5	2
13	Finnish Parkinson's disease study integrating protein-protein interaction network data with exome sequencing analysis. <i>Scientific Reports</i> , 2019 , 9, 18865	4.9	3
12	Mutation m.15923A>G in the MT-TT gene causes mild myopathy - case report of an adult-onset phenotype. <i>BMC Neurology</i> , 2018 , 18, 149	3.1	3
11	Molecular Epidemiology of Charcot-Marie-Tooth Disease in Northern Ostrobothnia, Finland: A Population-Based Study. <i>Neuroepidemiology</i> , 2017 , 49, 34-39	5.4	14
10	A nonsynonymous mutation in the WFS1 gene in a Finnish family with age-related hearing impairment. <i>Hearing Research</i> , 2017 , 355, 97-101	3.9	7
9	A novel MTTT mutation m.15933G > A revealed in analysis of mitochondrial DNA in patients with suspected mitochondrial disease. <i>BMC Medical Genetics</i> , 2017 , 18, 14	2.1	2
8	Case report: a novel frameshift mutation in the mitochondrial cytochrome c oxidase II gene causing mitochondrial disorder. <i>BMC Neurology</i> , 2017 , 17, 96	3.1	4
7	The m.7510T>C mutation: Hearing impairment and a complex neurologic phenotype. <i>Brain and Behavior</i> , 2017 , 7, e00859	3.4	3
6	A novel mutation m.8561C>G in MT-ATP6/8 causing a mitochondrial syndrome with ataxia, peripheral neuropathy, diabetes mellitus, and hypergonadotropic hypogonadism. <i>Journal of Neurology</i> , 2016 , 263, 2188-2195	5.5	32
5	Effects of pathogenic mutations in membrane subunits of mitochondrial Complex I on redox activity and proton translocation studied by modeling in Escherichia coli. <i>Mitochondrion</i> , 2015 , 22, 23-30	4.9	1
4	Novel mitofusin 2 splice-site mutation causes Charcot-Marie-Tooth disease type 2 with prominent sensory dysfunction. <i>Neuromuscular Disorders</i> , 2014 , 24, 360-4	2.9	3

3	WFS1 mutations in hearing-impaired children. <i>International Journal of Audiology</i> , 2014 , 53, 446-51	2.6	3
2	Juvenile parkinsonism, hypogonadism and Leigh-like MRI changes in a patient with m.4296G>A mutation in mitochondrial DNA. <i>Mitochondrion</i> , 2013 , 13, 83-6	4.9	18
1	WFS1 variants in Finnish patients with diabetes mellitus, sensorineural hearing impairment or optic atrophy, and in suicide victims. <i>Journal of Human Genetics</i> , 2013 , 58, 495-500	4.3	7