Laura Kytvuori

List of Publications by Citations

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	A novel mutation m.8561C>G in MT-ATP6/8 causing a mitochondrial syndrome with ataxia, peripheral neuropathy, diabetes mellitus, and hypergonadotropic hypogonadism. *Journal of Neurology*, **2016**, 263, 2188-2195	5.5	32
19	Juvenile parkinsonism, hypogonadism and Leigh-like MRI changes in a patient with m.4296G>A mutation in mitochondrial DNA. *Mitochondrion*, **2013**, 13, 83-6	4.9	18
18	Molecular Epidemiology of Charcot-Marie-Tooth Disease in Northern Ostrobothnia, Finland: A Population-Based Study. *Neuroepidemiology*, **2017**, 49, 34-39	5.4	14
17	A nonsynonymous mutation in the WFS1 gene in a Finnish family with age-related hearing impairment. *Hearing Research*, **2017**, 355, 97-101	3.9	7
16	WFS1 variants in Finnish patients with diabetes mellitus, sensorineural hearing impairment or optic atrophy, and in suicide victims. *Journal of Human Genetics*, **2013**, 58, 495-500	4.3	7
15	Case report: a novel frameshift mutation in the mitochondrial cytochrome c oxidase II gene causing mitochondrial disorder. *BMC Neurology*, **2017**, 17, 96	3.1	4
14	Novel mitofusin 2 splice-site mutation causes Charcot-Marie-Tooth disease type 2 with prominent sensory dysfunction. *Neuromuscular Disorders*, **2014**, 24, 360-4	2.9	3
13	The m.7510T>C mutation: Hearing impairment and a complex neurologic phenotype. *Brain and Behavior*, **2017**, 7, e00859	3.4	3
12	WFS1 mutations in hearing-impaired children. *International Journal of Audiology*, **2014**, 53, 446-51	2.6	3
11	Biallelic expansion in RFC1 as a rare cause of Parkinsonâs disease.. *Npj Parkinsonâs Disease*, **2022**, 8, 6	9.7	3
10	Finnish Parkinsonâs disease study integrating protein-protein interaction network data with exome sequencing analysis. *Scientific Reports*, **2019**, 9, 18865	4.9	3
9	Mutation m.15923A>G in the MT-TT gene causes mild myopathy - case report of an adult-onset phenotype. *BMC Neurology*, **2018**, 18, 149	3.1	3
8	Mutation Analysis of the Genes Linked to Early Onset Alzheimerâs Disease and Frontotemporal Lobar Degeneration. *Journal of Alzheimerâs Disease*, **2019**, 69, 775-782	4.3	2
7	Analysis of functional variants in mitochondrial DNA of Finnish athletes. *BMC Genomics*, **2019**, 20, 784	4.5	2
6	A novel MTTT mutation m.15933G > A revealed in analysis of mitochondrial DNA in patients with suspected mitochondrial disease. *BMC Medical Genetics*, **2017**, 18, 14	2.1	2
5	Molecular epidemiology of hereditary ataxia in Finland. *BMC Neurology*, **2021**, 21, 382	3.1	2
4	Effects of pathogenic mutations in membrane subunits of mitochondrial Complex I on redox activity and proton translocation studied by modeling in Escherichia coli. *Mitochondrion*, **2015**, 22, 23-30	4.9	1

3	Mitochondrial DNA variation in sudden cardiac death: a population-based study. *International Journal of Legal Medicine*, **2020**, 134, 39-44	3.1	1
2	Association of mitochondrial DNA haplogroups J and K with low response in exercise training among Finnish military conscripts. *BMC Genomics*, **2021**, 22, 75	4.5	0
1	Mutation Analysis of the Genes Associated with Parkinsonas Disease in a Finnish Cohort of Early-Onset Dementia. *Journal of Alzheimer's Disease*, **2020**, 76, 955-965	4.3	