

# Magdalena Kaliszewska

## List of Publications by Year in descending order

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Version: 2024-02-01

11  
papers

115  
citations

1937685

4  
h-index

1372567

10  
g-index

11  
all docs

11  
docs citations

11  
times ranked

239  
citing authors

#	ARTICLE	IF	CITATIONS
1	Comprehensive non-invasive assessment of electrocardiographic abnormalities and cardiac arrhythmias in patients with genetically confirmed mitochondrial diseases. <i>Journal of Electrocardiology</i> , 2021, 65, 136-142.	0.9	1
2	Progressive External Ophthalmoplegia in Polish Patients – From Clinical Evaluation to Genetic Confirmation. <i>Genes</i> , 2021, 12, 54.	2.4	1
3	Whole exome sequencing identifies a homozygous POLG2 missense variant in an adult patient presenting with optic atrophy, movement disorders, premature ovarian failure and mitochondrial DNA depletion. <i>European Journal of Medical Genetics</i> , 2020, 63, 103821.	1.3	5
4	Answer to Finsterer about “Multisystem presentation of a homozygous POLG2 variant”. <i>European Journal of Medical Genetics</i> , 2020, 63, 103900.	1.3	0
5	The frequency of mitochondrial polymerase gamma related disorders in a large Polish population cohort. <i>Mitochondrion</i> , 2019, 47, 179-187.	3.4	4
6	Nuclear genes involved in mitochondrial diseases caused by instability of mitochondrial DNA. <i>Journal of Applied Genetics</i> , 2018, 59, 43-57.	1.9	62
7	Identification of the first in Poland CACNA1A gene mutation in familial hemiplegic migraine. Case report. <i>Neurologia i Neurochirurgia Polska</i> , 2017, 51, 184-189.	1.2	2
8	Mitochondrial DNA levels in Huntington disease leukocytes and dermal fibroblasts. <i>Metabolic Brain Disease</i> , 2017, 32, 1237-1247.	2.9	19
9	Yeast model analysis of novel polymerase gamma variants found in patients with autosomal recessive mitochondrial disease. <i>Human Genetics</i> , 2015, 134, 951-966.	3.8	17
10	Mitochondrial encephalomyopathy: Towards diagnosis. A case report. <i>Neurologia i Neurochirurgia Polska</i> , 2014, 48, 76-80.	1.2	1
11	Mitochondrial cytopathies: clinical, morphological and genetic characteristics. <i>Neurologia i Neurochirurgia Polska</i> , 2009, 43, 216-27.	1.2	3