Magdalena Kaliszewska

List of Publications by Year
in descending order

11	115	1937685 4	1372567 10
papers	citations	h-index	g-index

11	11	11	239
all docs	docs citations	times ranked	citing authors

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