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Recommendations for the Management of Strokelike Episodes in Patients With Mitochondrial Encephalomyopathy, Lactic Acidosis, and Strokelike Episode

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#	Paper	IF	Citations
78	Mitochondrial diseases. <i>Nature Reviews Disease Primers</i> , 2016 , 2, 16080	51.1	585
77	Congenital Pediatric Dementia: a Case Study of Mitochondrial Myopathy, Encephalopathy, Lactic Acidosis, and Stroke-like Episodes (MELAS) Syndrome. <i>Journal of Pediatric Neuropsychology</i> , 2017 , 3, 96-105	0.7	
76	Increased cerebral blood flow as a predictor of episodes in MELAS using multimodal MRI. <i>Journal of Magnetic Resonance Imaging</i> , 2017 , 46, 915-918	5.6	6
75	Arginine and citrulline for the treatment of MELAS syndrome. <i>FIRE Forum for International Research in Education</i> , 2017 , 5,	1.4	24
74	Case 13-2017. A 41-Year-Old Man with Hearing Loss, Seizures, Weakness, and Cognitive Decline. <i>New England Journal of Medicine</i> , 2017 , 376, 1668-1678	59.2	1
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72	Therapies for mitochondrial diseases and current clinical trials. <i>Molecular Genetics and Metabolism</i> , 2017 , 122, 1-9	3.7	99
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70	Recognition, investigation and management of mitochondrial disease. <i>Archives of Disease in Childhood</i> , 2017 , 102, 1082-1090	2.2	24
69	Mitochondrial Encephalomyopathy, Lactic Acidosis, and Stroke (MELAS). 2017 ,		
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63	Metabolomes of mitochondrial diseases and inclusion body myositis patients: treatment targets and biomarkers. <i>EMBO Molecular Medicine</i> , 2018 , 10,	12	33
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61	Clinical syndromes associated with mtDNA mutations: where we stand after 30 years. <i>Essays in Biochemistry</i> , 2018 , 62, 235-254	7.6	19
60	Mitochondrial DNA mutation "m.3243A>G"-Heterogeneous clinical picture for cardiologists ("m.3243A>G": A phenotypic chameleon). <i>Congenital Heart Disease</i> , 2018 , 13, 671-677	3.1	11
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58	Dynamic derangement in amino acid profile during and after a stroke-like episode in adult-onset mitochondrial disease: a case report. <i>Journal of Medical Case Reports</i> , 2019 , 13, 313	1.2	4
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54	Diagnosis of adult-onset MELAS syndrome in a 63-year-old patient with suspected recurrent strokes - a case report. <i>BMC Neurology</i> , 2019 , 19, 91	3.1	10
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19	Adult-Onset Mitochondrial Encephalopathy, Lactic Acidosis, and Stroke-Like Episodes (MELAS) in a Patient Without Significant Family History.. <i>Cureus</i> , 2022 , 14, e21597	1.2	
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- 5 Currently available therapies in mitochondrial disease. **2023**, 189-206
- 4 Stroke in the young. **2023**, 36, 131-139
- 3 Clinical score for early diagnosis and treatment of stroke-like episodes in MELAS syndrome.
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