

# N L Sheremet

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9382474/publications.pdf>

Version: 2024-02-01

19  
papers

231  
citations

1162367

8  
h-index

1058022

14  
g-index

44  
all docs

44  
docs citations

44  
times ranked

263  
citing authors

#	ARTICLE	IF	CITATIONS
1	Impaired complex I repair causes recessive Leber's hereditary optic neuropathy. <i>Journal of Clinical Investigation</i> , 2021, 131, .	3.9	89
2	Plasma FGF21 and GDF15 are elevated in different inherited metabolic diseases and are not diagnostic for mitochondrial disorders. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 918-933.	1.7	30
3	Three rare pathogenic mtDNA substitutions in LHON patients with low heteroplasmy. <i>Mitochondrion</i> , 2020, 50, 139-144.	1.6	18
4	Previously unclassified mutation of mtDNA m.3472T>C: Evidence of pathogenicity in Leber's hereditary optic neuropathy. <i>Biochemistry (Moscow)</i> , 2016, 81, 748-754.	0.7	8
5	ETIOLOGICAL STRUCTURE OF NONGLAUCOMATOUS OPTIC NEUROPATHIES. <i>The Siberian Scientific Medical Journal</i> , 2018, , .	0.1	4
6	High-Resolution Respirometry in Diagnostics of Mitochondrial Diseases Caused by Mitochondrial Complex I Deficiency. <i>Biochemistry (Moscow) Supplement Series B: Biomedical Chemistry</i> , 2018, 12, 43-49.	0.2	2
7	Damage to the retinal pigment epithelium in the experiment. <i>Modern Technologies in Ophthalmology</i> , 2019, , 215-217.	0.0	2
8	Leber's Hereditary Optic Neuropathy with Neurological Abnormalities. Case Report. <i>Oftalmologiya</i> , 2021, 18, 753-757.	0.2	2
9	Clinical features of vision improvement in patients with Leber hereditary optic neuropathy. <i>Point of View East &amp; West</i> , 2020, , 47-49.	0.0	2
10	Comparative study on aging, UV treatment, and radiation on cataract formation. <i>Biophysics (Russian) Tj ETQq0 0 0 rgBT /Overlock 10 T</i>	0.2	1
11	Clinical features of toxic optic neuropathies. <i>Kazan Medical Journal</i> , 2017, 98, 400-403.	0.1	1
12	Metabolic alterations in Leber Optic Neuropathy patients. <i>Modern Technologies in Ophthalmology</i> , 2019, , 212-214.	0.0	1
13	Optomyelitis associated with the presence of antibodies to myelin oligodendrocyte glycoprotein. Case report. <i>Consilium Medicum</i> , 2022, 24, 132-136.	0.1	1
14	The Role of Mitophagy in Hereditary Optic Neuropathies. Literature Review. <i>Oftalmologiya</i> , 2021, 18, 646-653.	0.2	0
15	Phenotype-genotype correlations in patients with inherited retinal diseases with [L541P; A1038V] mutation in the ABCA4 gene. <i>Point of View East &amp; West</i> , 2018, , 51-54.	0.0	0
16	The role of acquired and hereditary thrombophilia in the development of optical neuropathies in young patients. <i>Point of View East &amp; West</i> , 2018, , 80-83.	0.0	0
17	Predicting the clinical course of inherited retinal diseases in patients with p.G1961E mutation in the ABCA4 gene. <i>Modern Technologies in Ophthalmology</i> , 2019, , 63-67.	0.0	0
18	Clinical manifestations of hereditary retinal diseases in patients with rare mutations in the ABCA4 gen. <i>Point of View East &amp; West</i> , 2020, , 54-57.	0.0	0

#	ARTICLE	IF	CITATIONS
19	The importance of OCT angiography and Doppler ultrasound methods in diagnosis of optic neuropathies. Vestnik Oftalmologii, 2022, 138, 132.	0.1	0