

Anders Oldfors

List of Publications by Year in descending order

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Version: 2024-02-01

110
papers

8,102
citations

136950

32
h-index

48315

88
g-index

144
all docs

144
docs citations

144
times ranked

9612
citing authors

#	ARTICLE	IF	CITATIONS
1	Premature ageing in mice expressing defective mitochondrial DNA polymerase. <i>Nature</i> , 2004, 429, 417-423.	27.8	2,318
2	Mitochondrial transcription factor A is necessary for mtDNA maintenance and embryogenesis in mice. <i>Nature Genetics</i> , 1998, 18, 231-236.	21.4	1,377
3	Impaired insulin secretion and β -cell loss in tissue-specific knockout mice with mitochondrial diabetes. <i>Nature Genetics</i> , 2000, 26, 336-340.	21.4	417
4	Dilated cardiomyopathy and atrioventricular conduction blocks induced by heart-specific inactivation of mitochondrial DNA gene expression. <i>Nature Genetics</i> , 1999, 21, 133-137.	21.4	393
5	The incidence of mitochondrial encephalomyopathies in childhood: Clinical features and morphological, biochemical, and DNA abnormalities. <i>Annals of Neurology</i> , 2001, 49, 377-383.	5.3	311
6	Ageing muscle: clonal expansions of mitochondrial DNA point mutations and deletions cause focal impairment of mitochondrial function. <i>Neuromuscular Disorders</i> , 2002, 12, 484-493.	0.6	180
7	Leber's hereditary optic neuropathy and complex I deficiency in muscle. <i>Annals of Neurology</i> , 1991, 30, 701-708.	5.3	155
8	A single mouse gene encodes the mitochondrial transcription factor A and a testis-specific nuclear HMG-box protein. <i>Nature Genetics</i> , 1996, 13, 296-302.	21.4	145
9	Polyglucosan body myopathy caused by defective ubiquitin ligase RBCK1. <i>Annals of Neurology</i> , 2013, 74, 914-919.	5.3	132
10	Mitochondrial encephalomyopathies in childhood. I. Biochemical and morphologic investigations. <i>Journal of Pediatrics</i> , 1991, 119, 242-250.	1.8	128
11	Cardiomyopathy and Exercise Intolerance in Muscle Glycogen Storage Disease O. <i>New England Journal of Medicine</i> , 2007, 357, 1507-1514.	27.0	123
12	Hereditary myosin myopathies. <i>Neuromuscular Disorders</i> , 2007, 17, 355-367.	0.6	117
13	Mitochondrial DNA Deletions in Muscle Fibers in Inclusion Body Myositis. <i>Journal of Neuropathology and Experimental Neurology</i> , 1995, 54, 581-587.	1.7	103
14	Clonal expansion of mitochondrial DNA with multiple deletions in autosomal dominant progressive external ophthalmoplegia. <i>Annals of Neurology</i> , 1996, 40, 707-713.	5.3	96
15	Glycogenin-1 Deficiency and Inactivated Priming of Glycogen Synthesis. <i>New England Journal of Medicine</i> , 2010, 362, 1203-1210.	27.0	95
16	Whole exome sequencing reveals mutations in <i>NARS2</i> and <i>PARS2</i> , encoding the mitochondrial asparaginyl-tRNA synthetase and prolyl-tRNA synthetase, in patients with Alpers syndrome. <i>Molecular Genetics & Genomic Medicine</i> , 2015, 3, 59-68.	1.2	87
17	Down-Regulation of Mitochondrial Transcription Factor a During Spermatogenesis in Humans. <i>Human Molecular Genetics</i> , 1997, 6, 185-1991.	2.9	75
18	Congenital myasthenic syndrome with tubular aggregates caused by GFPT1 mutations. <i>Journal of Neurology</i> , 2012, 259, 838-850.	3.6	72

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19	A new muscle glycogen storage disease associated with glycogeninâ€1 deficiency. <i>Annals of Neurology</i> , 2014, 76, 891-898.	5.3	72
20	Mitochondrial ATP-Synthase Deficiency in a Child with 3-Methylglutaconic Aciduria. <i>Pediatric Research</i> , 1992, 32, 731-736.	2.3	65
21	Polyglucosan storage myopathies. <i>Molecular Aspects of Medicine</i> , 2015, 46, 85-100.	6.4	63
22	Pathogenetic aspects of the A8344G mutation of mitochondrial DNA associated with MERRF syndrome and multiple symmetric lipomas. <i>Muscle and Nerve</i> , 1995, 18, S102-S106.	2.2	59
23	Restricted use of T cell receptor V genes in endomysial infiltrates of patients with inflammatory myopathies. <i>European Journal of Immunology</i> , 1994, 24, 2659-2663.	2.9	56
24	Analysis of multiple mitochondrial DNA deletions in inclusion body myositis. <i>Human Mutation</i> , 1997, 10, 381-386.	2.5	53
25	New cardiac and skeletal protein aggregate myopathy associated with combined MuRF1 and MuRF3 mutations. <i>Human Molecular Genetics</i> , 2015, 24, 3638-3650.	2.9	51
26	Mitochondrial (Mt)Dna Changes in Tissue May Not be Reflected by Depletion of Mtdna in Peripheral Blood Mononuclear Cells in HIV-Infected Patients. <i>Antiviral Therapy</i> , 2006, 11, 601-608.	1.0	48
27	New insights in the field of muscle glycogenoses. <i>Current Opinion in Neurology</i> , 2013, 26, 544-553.	3.6	44
28	Accurate mapping of mitochondrial DNA deletions and duplications using deep sequencing. <i>PLoS Genetics</i> , 2020, 16, e1009242.	3.5	41
29	Mitochondrial Encephalomyopathies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2003, 62, 217-227.	1.7	38
30	De novo mutation in the mitochondrial ATP synthase subunit 6 gene (T8993G) with rapid segregation resulting in Leigh syndrome in the offspring. <i>Human Genetics</i> , 1995, 96, 290-4.	3.8	35
31	Cardiomyopathy with lethal arrhythmias associated with inactivation of KLHL24. <i>Human Molecular Genetics</i> , 2019, 28, 1919-1929.	2.9	35
32	Xâ€linked myotubular myopathy: a linkage study. <i>Clinical Genetics</i> , 1990, 37, 335-340.	2.0	34
33	Recessive myosin myopathy with external ophthalmoplegia associated with MYH2 mutations. <i>European Journal of Human Genetics</i> , 2014, 22, 801-808.	2.8	33
34	Early onset cardiomyopathy in females with Danon disease. <i>Neuromuscular Disorders</i> , 2015, 25, 493-501.	0.6	32
35	Cardiomyopathy as presenting sign of glycogeninâ€1 deficiencyâ€”report of three cases and review of the literature. <i>Journal of Inherited Metabolic Disease</i> , 2017, 40, 139-149.	3.6	32
36	<i>TANGO2</i> deficiency as a cause of neurodevelopmental delay with indirect effects on mitochondrial energy metabolism. <i>Journal of Inherited Metabolic Disease</i> , 2019, 42, 898-908.	3.6	32

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37	Mitochondrial pathology in inclusion body myositis. <i>Neuromuscular Disorders</i> , 2015, 25, 281-288.	0.6	31
38	Broad phenotypic variability in patients with complex I deficiency due to mutations in <i>NDUFS1</i> and <i>NDUFV1</i> . <i>Mitochondrion</i> , 2015, 21, 33-40.	3.4	30
39	A novel complex neurological phenotype due to a homozygous mutation in <i>FDX2</i> . <i>Brain</i> , 2018, 141, 2289-2298.	7.6	29
40	Muscle pathology and whole-body MRI in a polyglucosan myopathy associated with a novel glycogenin-1 mutation. <i>Neuromuscular Disorders</i> , 2015, 25, 780-785.	0.6	28
41	Anticipation of autosomal dominant progressive external ophthalmoplegia with hypogonadism. , 1996, 19, 1561-1569.		27
42	<i>B3GALNT2</i> is a gene associated with congenital muscular dystrophy with brain malformations. <i>European Journal of Human Genetics</i> , 2014, 22, 707-710.	2.8	27
43	1st ENMC European meeting: The EURO-NMD pathology working group Recommended Standards for Muscle Pathology Amsterdam, The Netherlands, 7 December 2018. <i>Neuromuscular Disorders</i> , 2019, 29, 483-485.	0.6	27
44	Upregulation of Fas/Fas ligand in inclusion body myositis. <i>Annals of Neurology</i> , 1998, 43, 127-130.	5.3	25
45	Pathogenic Variants in the Myosin Chaperone <i>UNC-45B</i> Cause Progressive Myopathy with Eccentric Cores. <i>American Journal of Human Genetics</i> , 2020, 107, 1078-1095.	6.2	24
46	<i>RBCK1</i> -related disease: A rare multisystem disorder with polyglucosan storage, autoinflammation, recurrent infections, skeletal, and cardiac myopathy—Four additional patients and a review of the current literature. <i>Journal of Inherited Metabolic Disease</i> , 2020, 43, 1002-1013.	3.6	23
47	Phenotypic and genotypic variability in Alpers syndrome. <i>European Journal of Paediatric Neurology</i> , 2012, 16, 379-389.	1.6	22
48	Loss of Sarcomeric Scaffolding as a Common Baseline Histopathologic Lesion in Titin-Related Myopathies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018, 77, 1101-1114.	1.7	22
49	A novel heteroplasmic point mutation in the mitochondrial <i>tRNA^{Lys}</i> gene in a sporadic case of mitochondrial encephalomyopathy: De novo mutation and no transmission to the offspring. <i>Human Mutation</i> , 1999, 13, 203-209.	2.5	21
50	Muscle pathology in Vici syndrome—A case study with a novel mutation in <i>EPG5</i> and a summary of the literature. <i>Neuromuscular Disorders</i> , 2017, 27, 771-776.	0.6	21
51	Parvovirus B19 in Endomyocardial Biopsy of Patients With Idiopathic Dilated Cardiomyopathy: Foe or Bystander?. <i>Journal of Cardiac Failure</i> , 2019, 25, 60-63.	1.7	21
52	Thick Filament Diseases. <i>Advances in Experimental Medicine and Biology</i> , 2008, 642, 78-91.	1.6	21
53	Fatal mitochondrial myopathy, lactic acidosis, and complex I deficiency associated with a heteroplasmic A→G mutation at position 3251 in the mitochondrial <i>tRNA^{Leu}(UUR)</i> gene. <i>Human Genetics</i> , 1996, 97, 269-273.	3.8	20
54	Glycogen pathways in disease: new developments in a classical field of medical genetics. <i>Journal of Inherited Metabolic Disease</i> , 2015, 38, 483-487.	3.6	20

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55	An AARS variant as the likely cause of Swedish type hereditary diffuse leukoencephalopathy with spheroids. <i>Acta Neuropathologica Communications</i> , 2019, 7, 188.	5.2	19
56	Carey-Fineman-Ziter syndrome with mutations in the myomaker gene and muscle fiber hypertrophy. <i>Neurology: Genetics</i> , 2018, 4, e254.	1.9	18
57	Hereditary myopathy with early respiratory failure is associated with misfolding of the titin fibronectin III 119 subdomain. <i>Neuromuscular Disorders</i> , 2014, 24, 373-379.	0.6	17
58	Leukoencephalopathy due to Complex II Deficiency and Bi-Allelic SDHB Mutations: Further Cases and Implications for Genetic Counselling. <i>JIMD Reports</i> , 2016, 33, 69-77.	1.5	17
59	Start codon mutation of GYG1 causing late-onset polyglucosan body myopathy with nemaline rods. <i>Journal of Neurology</i> , 2016, 263, 2133-2135.	3.6	17
60	A new early-onset neuromuscular disorder associated with kyphoscoliosis peptidase (KY) deficiency. <i>European Journal of Human Genetics</i> , 2016, 24, 1771-1777.	2.8	17
61	Deep morphological analysis of muscle biopsies from type III glycogenesis (GSDIII), debranching enzyme deficiency, revealed stereotyped vacuolar myopathy and autophagy impairment. <i>Acta Neuropathologica Communications</i> , 2019, 7, 167.	5.2	17
62	Mitochondrial complex IV deficiency caused by a novel frameshift variant in MT-CO2 associated with myopathy and perturbed acylcarnitine profile. <i>European Journal of Human Genetics</i> , 2019, 27, 331-335.	2.8	17
63	Glycogenin is Dispensable for Glycogen Synthesis in Human Muscle, and Glycogenin Deficiency Causes Polyglucosan Storage. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, 557-566.	3.6	17
64	Mitochondrial DNA variants in inclusion body myositis characterized by deep sequencing. <i>Brain Pathology</i> , 2021, 31, e12931.	4.1	17
65	Hypoxic cardiac fibroblasts from failing human hearts decrease cardiomyocyte beating frequency in an ALOX15 dependent manner. <i>PLoS ONE</i> , 2018, 13, e0202693.	2.5	16
66	Diagnosis, management, and outcome of cardiac sarcoidosis and giant cell myocarditis: a Swedish single center experience. <i>BMC Cardiovascular Disorders</i> , 2022, 22, 192.	1.7	16
67	Diagnosis, pathogenesis and treatment of inclusion body myositis. <i>Current Opinion in Neurology</i> , 2005, 18, 497-503.	3.6	15
68	A novel dynamin-2 gene mutation associated with a late-onset centronuclear myopathy with necklace fibres. <i>Neuromuscular Disorders</i> , 2015, 25, 345-348.	0.6	15
69	Loss of supervillin causes myopathy with myofibrillar disorganization and autophagic vacuoles. <i>Brain</i> , 2020, 143, 2406-2420.	7.6	15
70	A novel MYH2 mutation in family members presenting with congenital myopathy, ophthalmoplegia and facial weakness. <i>Journal of Neurology</i> , 2016, 263, 1427-1433.	3.6	14
71	Some DNM2 mutations cause extremely severe congenital myopathy and phenocopy myotubular myopathy. <i>Acta Neuropathologica Communications</i> , 2018, 6, 93.	5.2	14
72	Long-term follow-up and characteristic pathological findings in severe nemaline myopathy due to LMOD3 mutations. <i>Neuromuscular Disorders</i> , 2019, 29, 108-113.	0.6	14

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73	The phenotypic variability and natural history of NARS2 associated disease. <i>European Journal of Paediatric Neurology</i> , 2021, 31, 31-37.	1.6	14
74	Proteomic characterisation of polyglucosan bodies in skeletal muscle in RBCK1 deficiency. <i>Neuropathology and Applied Neurobiology</i> , 2022, 48, .	3.2	13
75	LCâ€MS/MS characterization of combined glycogenin-1 and glycogenin-2 enzymatic activities reveals their self-glucosylation preferences. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2014, 1844, 398-405.	2.3	11
76	The Atrioventricular Junction: A Potential Niche Region for Progenitor Cells in the Adult Human Heart. <i>Stem Cells and Development</i> , 2019, 28, 1078-1088.	2.1	11
77	Progressive external ophthalmoplegia associated with novel <i>MT</i> and <i>TN</i> mutations. <i>Acta Neurologica Scandinavica</i> , 2021, 143, 103-108.	2.1	11
78	Cardiac arrest in Wilson's disease after curative liver transplantation: a life-threatening complication of myocardial copper excess?. <i>ESC Heart Failure</i> , 2019, 6, 228-231.	3.1	10
79	Epidemiology, Survival, and Clinical Characteristics of Inclusion Body Myositis. <i>Annals of Neurology</i> , 2022, 92, 201-212.	5.3	10
80	Histopathological changes in skeletal muscle associated with chronic ischaemia. <i>Apmis</i> , 2016, 124, 935-941.	2.0	9
81	A novel neuromuscular form of glycogen storage disease type IV with arthrogryposis, spinal stiffness and rare polyglucosan bodies in muscle. <i>Neuromuscular Disorders</i> , 2016, 26, 681-687.	0.6	9
82	Novel myopathy in a newborn with Shwachmanâ€Diamond syndrome and review of neonatal presentation. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 1155-1164.	1.2	9
83	Is Glycogenin Essential for Glycogen Synthesis?. <i>Cell Metabolism</i> , 2017, 26, 12-14.	16.2	9
84	COX deficiency and leukoencephalopathy due to a novel homozygous <i>APOPT1/COA8</i> mutation. <i>Neurology: Genetics</i> , 2020, 6, e464.	1.9	9
85	Functional characterization of <i>GYG1</i> variants in two patients with myopathy and glycogenin-1 deficiency. <i>Neuromuscular Disorders</i> , 2019, 29, 951-960.	0.6	8
86	Deep sequencing of mitochondrial DNA and characterization of a novel <i>POLG</i> mutation in a patient with arPEO. <i>Neurology: Genetics</i> , 2020, 6, e391.	1.9	8
87	Association between muscle strength, histopathology, and magnetic resonance imaging in sporadic inclusion body myositis. <i>Acta Neurologica Scandinavica</i> , 2019, 139, 177-182.	2.1	7
88	The incidence of mitochondrial encephalomyopathies in childhood: Clinical features and morphological, biochemical, and DNA abnormalities. <i>Annals of Neurology</i> , 2001, 49, 377-383.	5.3	7
89	Grand paternal inheritance of X-linked myotubular myopathy due to mosaicism, and identification of necklace fibers in an asymptomatic male. <i>Neuromuscular Disorders</i> , 2017, 27, 843-847.	0.6	6
90	Paranodal myelin retraction in protein deficient and normal rats. <i>Acta Neuropathologica</i> , 1977, 40, 249-252.	7.7	5

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91	THE EFFECT OF Î²-ADRENERGIC BLOCKADE ON THE MYOPATHIC CHANGES IN EXPERIMENTAL HYPERTHYROIDISM IN RATS. <i>Acta Pathologica, Microbiologica, Et Immunologica Scandinavica Section A, Pathology</i> , 1986, 94A, 91-99.	0.3	5
92	Danon disease presenting with early onset of hypertrophic cardiomyopathy and peripheral pigmentary retinal dystrophy in a female with a <i>de novo</i> novel mosaic mutation in the <i>LAMP2</i> gene. <i>Ophthalmic Genetics</i> , 2019, 40, 227-236.	1.2	5
93	Somatostatin Receptor Positron Emission Tomography/Computed Tomography in Giant Cell Myocarditis: A Promising Approach to Molecular Myocardial Inflammation Imaging. <i>Circulation: Cardiovascular Imaging</i> , 2022, 15, CIRCIMAGING121013551.	2.6	5
94	The effect of undernutrition on spinal ganglion nerve cell size. <i>Acta Neuropathologica</i> , 1977, 39, 75-80.	7.7	4
95	Oxygen consumption in platelets as an adjunct diagnostic method for pediatric mitochondrial disease. <i>Pediatric Research</i> , 2018, 83, 455-465.	2.3	4
96	251st ENMC international workshop: Polyglucosan storage myopathies 13-15 December 2019, Hoofddorp, the Netherlands. <i>Neuromuscular Disorders</i> , 2021, 31, 466-477.	0.6	4
97	The localization of amyloid precursor protein to ependymal cilia in vertebrates and its role in ciliogenesis and brain development in zebrafish. <i>Scientific Reports</i> , 2021, 11, 19115.	3.3	4
98	Tissue distribution and disease manifestations of the tRNA ^{Lys} A?G(8344) mitochondrial DNA mutation in a case of myoclonus epilepsy and ragged red fibres. <i>Acta Neuropathologica</i> , 1995, 90, 328-333.	7.7	4
99	Expression pattern of mitochondrial respiratory chain enzymes in skeletal muscle of patients with mitochondrial myopathy associated with the homoplasmic m.14674T>C variant. <i>Brain Pathology</i> , 2021, , e13038.	4.1	4
100	Duchenne muscular dystrophy and spinal muscular atrophy type I segregating in the same family. <i>Clinical Genetics</i> , 1994, 45, 97-103.	2.0	2
101	Functional analysis of a novel POLÎ³A mutation associated with a severe perinatal mitochondrial encephalomyopathy. <i>Neuromuscular Disorders</i> , 2021, 31, 348-358.	0.6	2
102	Cardiac involvement in immune-mediated necrotizing myopathy: insights from CMR and somatostatin receptor PET/CT. <i>European Heart Journal Cardiovascular Imaging</i> , 2022, 23, e237-e237.	1.2	2
103	Changes in specific gravity as a sign of disturbed brain maturation in protein-deprived rats. <i>Acta Neurologica Scandinavica</i> , 2009, 73, 76-77.	2.1	1
104	Tubular aggregates in congenital myasthenic syndrome. <i>Neuromuscular Disorders</i> , 2018, 28, 174-175.	0.6	1
105	Tropomyosin mutations responsible for muscle weakness in inherited skeletal muscle diseases. , 2008, , 20-21.		0
106	Neuromuscular Disorders Special Issue 2021 Marking the 90th birthday of Victor Dubowitz, Founding Editor-in-Chief. <i>Neuromuscular Disorders</i> , 2021, 31, 919-920.	0.6	0
107	Accurate mapping of mitochondrial DNA deletions and duplications using deep sequencing. , 2020, 16, e1009242.		0
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109	Accurate mapping of mitochondrial DNA deletions and duplications using deep sequencing. , 2020, 16, e1009242.		0
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