Giovanna Cenacchi

List of Publications by Year
in descending order

150	4,803	81839	118793
papers	citations	39 h-index	62 g-index

155	155	155	6507
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A new acrylic resin formulation: a useful tool for histological, ultrastructural, and immunocytochemical investigations.. Journal of Histochemistry and Cytochemistry, 1992, 40, 1799-1804.	1.3	190
2	Reversibility of GERD Ultrastructural Alterations and Relief of Symptoms After Omeprazole Treatment. American Journal of Gastroenterology, 2005, 100, 537-542.	0.2	190
3	Extraventricular Neoplasms with Neurocytoma Features. American Journal of Surgical Pathology, 1997, 21, 206-212.	2.1	176
4	`Spindle Cell Oncocytoma' of the Adenohypophysis. American Journal of Surgical Pathology, 2002, 26, 1048-1055.	2.1	160
5	Peripheral neuropathy associated with primary Sjogren's syndrome.. Journal of Neurology, Neurosurgery and Psychiatry, 1994, 57, 983-986.	0.9	123
6	Dilated intercellular spaces as a marker of oesophageal damage: comparative results in gastro-oesophageal reflux disease with or without bile reflux. Alimentary Pharmacology and Therapeutics, 2003, 18, 525-532.	1.9	122
7	The genetic and metabolic signature of oncocytic transformation implicates HIF1Î± destabilization. Human Molecular Genetics, 2010, 19, 1019-1032.	1.4	113
8	Molecular mechanisms of CD99-induced caspase-independent cell death and cellâ€“cell adhesion in Ewing's sarcoma cells: actin and zyxin as key intracellular mediators. Oncogene, 2004, 23, 5664-5674.	2.6	108
9	Amyloid deposition as a cause of atrial remodelling in persistent valvular atrial fibrillation. European Heart Journal, 2004, 25, 1237-1241.	1.0	101
10	Induction of chromosomal instability in colonic cells by the human polyomavirus JC virus. Cancer Research, 2003, 63, 7256-62.	0.4	97
11	A novel deletion in the GTPase domain of OPA1 causes defects in mitochondrial morphology and distribution, but not in function. Human Molecular Genetics, 2008, 17, 3291-3302.	1.4	91
12	Ultrastructural changes in dysferlinopathy support defective membrane repair mechanism. Journal of Clinical Pathology, 2005, 58, 190-195.	1.0	90
13	Basement Membrane Production by Hepatocytes in Chronic Liver Disease. Hepatology, 1984, 4, 1167-1172.	3.6	89
14	Liver transplantation for mitochondrial neurogastrointestinal encephalomyopathy. Annals of Neurology, 2016, 80, 448-455.	2.8	81
15	Emerging Tumor Entities and Variants of CNS Neoplasms. Journal of Neuropathology and Experimental Neurology, 2004, 63, 185-192.	0.9	78
16	Chordoid Glioma of the Third Ventricle. American Journal of Surgical Pathology, 2001, 25, 401-405.	2.1	75
17	TMEM199 Deficiency Is a Disorder of Golgi Homeostasis Characterized by Elevated Aminotransferases, Alkaline Phosphatase, and Cholesterol and Abnormal Glycosylation. American Journal of Human Genetics, 2016, 98, 322-330.	2.6	73
18	Medullocytoma (Lipidized Medulloblastoma). American Journal of Surgical Pathology, 1996, 20, 656-664.	2.1	73

#	Article	IF	Citations
19	Supratentorial Cortical Ependymoma: Report of Three Cases. Neurosurgery, 2005, 57, E192-E192.	0.6	65
20	Lipomatous Meningioma. American Journal of Surgical Pathology, 2001, 25, 769-775.	2.1	64
21	Review: Danon disease: Review of natural history and recent advances. Neuropathology and Applied Neurobiology, 2020, 46, 303-322.	1.8	62
22	Brain ischemic injury in COVIDâ€19â€infected patients: a series of 10 postâ€mortem cases. Brain Pathology, 2021, 31, 205-210.	2.1	61
23	Bacteria Sorting by Field-Flow Fractionation. Application to Whole-CellEscherichia coliVaccine Strains. Analytical Chemistry, 2002, 74, 4895-4904.	3.2	59
24	Differentiation of Mesenchymal Stem Cells Derived from Pancreatic Islets and Bone Marrow into Islet-Like Cell Phenotype. PLoS ONE, 2011, 6, e28175.	1.1	59
25	Pituicytoma: Ultrastructural Evidence of a Possible Origin from Folliculo-Stellate Cells of the Adenohypophysis. Ultrastructural Pathology, 2001, 25, 309-312.	0.4	58
26	Somatic complex I disruptive mitochondrial DNA mutations are modifiers of tumorigenesis that correlate with low genomic instability in pituitary adenomas. Human Molecular Genetics, 2013, 22, 226-238.	1.4	55
27	A Mutation in theCASQ1Gene Causes a Vacuolar Myopathy with Accumulation of Sarcoplasmic Reticulum Protein Aggregates. Human Mutation, 2014, 35, 1163-1170.	1.1	53
28	Satellite cell characterization from aging human muscle. Neurological Research, 2010, 32, 63-72.	0.6	50
29	True and false aneurysms in BehÃ§et's disease: Case report with ultrastructural observations. Journal of Vascular Surgery, 1993, 17, 762-767.	0.6	49
30	Immunohistochemical and Ultrastructural Investigation of Neural Differentiation in Ewing Sarcoma/PNET of Bone and Soft Tissues. Ultrastructural Pathology, 2001, 25, 219-225.	0.4	49
31	A novel Notch3 gene mutation not involving a cysteine residue in an Italian family with CADASIL. Neurology, 2004, 63, 561-564.	1.5	48
32	Homozygous <scp>NOTCH</scp> 3 null mutation and impaired <scp>NOTCH</scp> 3 signaling in recessive earlyâ€onset arteriopathy and cavitating leukoencephalopathy. EMBO Molecular Medicine, 2015, 7, 848-858.	3.3	48
33	Peculiar combinations of individually non-pathogenic missense mitochondrial DNA variants cause low penetrance Leberâ€™s hereditary optic neuropathy. PLoS Genetics, 2018, 14, e1007210.	1.5	47
34	Mitochondrial neurogastrointestinal encephalomyopathy (MNGIE): Position paper on diagnosis, prognosis, and treatment by the <scp>MNGIE</scp> International Network. Journal of Inherited Metabolic Disease, 2021, 44, 376-387.	1.7	47
35	MYH7 gene mutation in myosin storage myopathy and scapulo-peroneal myopathy. Neuromuscular Disorders, 2007, 17, 321-329.	0.3	46
36	Development and evaluation of a decellularized membrane from human dermis. Journal of Tissue Engineering and Regenerative Medicine, 2014, 8, 325-336.	1.3	44

#	Article	IF	Citations
37	Aberrant Compartment Formation by HSPB2 Mislocalizes Lamin A and Compromises Nuclear Integrity and Function. Cell Reports, 2017, 20, 2100-2115.	2.9	43
38	Liver as a Source for Thymidine Phosphorylase Replacement in Mitochondrial Neurogastrointestinal Encephalomyopathy. PLoS ONE, 2014, 9, e96692.	1.1	42
39	Primary coenzyme Q10 deficiency presenting as fatal neonatal multiorgan failure. European Journal of Human Genetics, 2015, 23, 1254-1258.	1.4	42
40	Androgen-dependent impairment of myogenesis in spinal and bulbar muscular atrophy. Acta Neuropathologica, 2013, 126, 109-121.	3.9	41
41	CD99 suppresses osteosarcoma cell migration through inhibition of ROCK2 activity. Oncogene, 2014, 33, 1912-1921.	2.6	41
42	Mesenchymal Stem Cells in Renal Function Recovery after Acute Kidney Injury: Use of a Differentiating Agent in a Rat Model. Cell Transplantation, 2011, 20, 1193-1208.	1.2	40
43	Bi-allelic mutations in TRAPPC2L result in a neurodevelopmental disorder and have an impact on RAB11 in fibroblasts. Journal of Medical Genetics, 2018, 55, 753-764.	1.5	39
44	The effects of palladium nanoparticles on the renal function of female Wistar rats. Nanotoxicology, 2015, 9, 843-851.	1.6	38
45	Ultrastructural Characterization of Oligodendroglial-like Cells in Central Nervous System Tumors. Ultrastructural Pathology, 1996, 20, 537-547.	0.4	37
46	Skeletal Muscle Satellite Cells in Amyotrophic Lateral Sclerosis. Ultrastructural Pathology, 2014, 38, 295-302.	0.4	37
47	Nonrandom gain of chromosome 7 in central neurocytoma: A chromosomal analysis and fluorescence in situ hybridization study. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 1997, 430, 47-51.	1.4	36
48	A clinically complex form of dominant optic atrophy (OPA8) maps on chromosome 16. Human Molecular Genetics, 2011, 20, 1893-1905.	1.4	36
49	Medullospheres from DAOY, UW228 and ONS-76 Cells: Increased Stem Cell Population and Proteomic Modifications. PLoS ONE, 2013, 8, e63748.	1.1	35
50	Magnetic Labelling of Mesenchymal Stem Cells with Iron-Doped Hydroxyapatite Nanoparticles as Tool for Cell Therapy. Journal of Biomedical Nanotechnology, 2016, 12, 909-921.	0.5	34
51	Ependymoma with neuropil-like islands: a case report with diagnostic and histogenetic implications. Acta Neuropathologica, 2005, 109, 231-234.	3.9	33
52	A mutation in PAK3 with a dual molecular effect deregulates the RAS/MAPK pathway and drives an X-linked syndromic phenotype. Human Molecular Genetics, 2014, 23, 3607-3617.	1.4	33
53	Gamma rays induce a p53-independent mitochondrial biogenesis that is counter-regulated by HIF1Î±. Cell Death and Disease, 2013, 4, e663-e663.	2.7	31
54	Ultrastructural and Immunohistochemical Contribution to the Histogenesis of Human Cardiac Myxoma. Ultrastructural Pathology, 1988, 12, 221-233.	0.4	30

#	Article	IF	Citations
55	Lipolysis and lipophagy in lipid storage myopathies. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2016, 1862, 1367-1373.	1.8	30
56	Functional expression of calciumâ€permeable canonical transient receptor potential 4â€containing channels promotes migration of medulloblastoma cells. Journal of Physiology, 2017, 595, 5525-5544.	1.3	30
57	Subpopulations of dermal skin fibroblasts secrete distinct extracellular matrix: implications for using skin substitutes in the clinic. British Journal of Dermatology, 2018, 179, 381-393.	1.4	30
58	Lipoastrocytoma: a rare low-grade astrocytoma variant of pediatric age. Acta Neuropathologica, 2002, 103, 152-156.	3.9	29
59	Clinical and ultrastructural spectrum of diffuse lung disease associated with surfactant protein C mutations. European Journal of Human Genetics, 2015, 23, 1033-1041.	1.4	29
60	Relationship between connective tissue cells and fibronectin in a sequential model of experimental hepatic fibrosis. Vigiliae Christianae, 1983, 43, 75-84.	0.1	28
61	Sleep and cardiovascular phenotype in middleâ€aged hypocretinâ€deficient narcoleptic mice. Journal of Sleep Research, 2014, 23, 98-106.	1.7	28
62	One-step esterification of nanocellulose in a BrÃ¸nsted acid ionic liquid for delivery to glioblastoma cancer cells. New Journal of Chemistry, 2018, 42, 5237-5242.	1.4	28
63	Update on polyglucosan storage diseases. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2019, 475, 671-686.	1.4	28
64	Validity of internal expression of the major histocompatibility complex class I in the diagnosis of inflammatory myopathies. Journal of Clinical Pathology, 2012, 65, 14-19.	1.0	26
65	The cellular component in the parietal infiltrate of inflammatory abdominal aortic aneurysms (IAAA). European Journal of Vascular Surgery, 1991, 5, 65-70.	0.9	25
66	Biochemical and ultrastructural evidence of endoplasmic reticulum stress in LGMD2I. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2007, 451, 1047-1055.	1.4	23
67	Comparison of muscle ultrastructure in myasthenia gravis with anti-MuSK and anti-AChR antibodies. Journal of Neurology, 2011, 258, 746-752.	1.8	23
68	Progression on metastatic neuroendocrine carcinoma from a recurrent prolactinoma: a case report. Journal of Clinical Pathology, 2002, 55, 148-151.	1.0	23
69	Hypertrophic inflammatory neuropathy involving bilateral brachial plexus. World Neurosurgery, 1999, 52, 458-465.	1.3	22
70	Variable phenotypic expression of chylomicron retention disease in a kindred carrying a mutation of the Sara2 gene. Metabolism: Clinical and Experimental, 2010, 59, 463-467.	1.5	21
71	Content and turnover of extracellular matrix protein in human â€œNon-specificâ€•and inflammatory abdominal aortic aneurysms. European Journal of Vascular Surgery, 1993, 7, 546-553.	0.9	20
72	Mixed Tumors, Myoepitheliomas, and Oncocytomas of the Soft Tissues Are Likely Members of the Same Family: A Clinicopathologic and Ultrastructural Study. Ultrastructural Pathology, 2001, 25, 399-418.	0.4	20

#	Article	IF	Citations
73	Ultrastructural changes in <scp>LGMD1F</scp>. Neuropathology, 2013, 33, 276-280.	0.7	20
74	Renal Thrombotic Microangiopathy in Concurrent COVID-19 Vaccination and Infection. Pathogens, 2021, 10, 1045.	1.2	20
75	Clinico-pathological Evaluation of Ciliary Dyskinesia: Diagnostic Role of Electron Microscopy. Ultrastructural Pathology, 2003, 27, 243-252.	0.4	19
76	Proteomic analysis of extracellular vesicles from medullospheres reveals a role for iron in the cancer progression of medulloblastoma. Molecular and Cellular Therapies, 2015, 3, 8.	0.2	19
77	Cerebral Mitochondrial Microangiopathy Leads to Leukoencephalopathy in Mitochondrial Neurogastrointestinal Encephalopathy. American Journal of Neuroradiology, 2018, 39, 427-434.	1.2	18
78	Melanotic schwannoma of the sympathetic ganglia: a histologic, immunohistochemical and ultrastructural study. Journal of Neuro-Oncology, 1997, 35, 149-152.	1.4	17
79	Renal calcium phosphate and oxalate deposition in prolonged vitamin B6 deficiency: studies on a rat model of urolithiasis. BJU International, 2002, 89, 571-575.	1.3	17
80	Radiobiologic response of medulloblastoma cell lines: involvement of Î²-catenin?. Journal of Neuro-Oncology, 2008, 90, 243-251.	1.4	17
81	Liver transplantation in mitochondrial neurogastrointestinal encephalomyopathy (MNGIE): clinical long-term follow-up and pathogenic implications. Journal of Neurology, 2020, 267, 3702-3710.	1.8	17
82	Peroneal muscular atrophy with hereditary spastic paraparesis (HMSN V) is pathologically heterogeneous. Acta Neuropathologica, 1992, 83, 196-201.	3.9	16
83	Human cytomegalovirus nuclear and cytoplasmic dense bodies. Archives of Virology, 1992, 123, 193-207.	0.9	16
84	Small heat-shock protein HSPB3 promotes myogenesis by regulating the lamin B receptor. Cell Death and Disease, 2021, 12, 452.	2.7	16
85	Transport and MHD studies at highTein FTU tokamak. Plasma Physics and Controlled Fusion, 1999, 41, B351-B365.	0.9	15
86	The Problematic Issue of Kufs Disease Diagnosis as Performed on Rectal Biopsies: A Case Report. Ultrastructural Pathology, 2004, 28, 43-48.	0.4	14
87	Lithium induces mortality in medulloblastoma cell lines. International Journal of Oncology, 2010, 37, 745-52.	1.4	14
88	The clinical spectrum of CASQ1-related myopathy. Neurology, 2018, 91, e1629-e1641.	1.5	14
89	The use of an acellular matrix derived from human dermis for the treatment of full-thickness skin wounds. Cell and Tissue Banking, 2019, 20, 183-192.	0.5	14
90	Effect of omeprazole on symptoms and ultrastructural esophageal damage in acid bile reflux. World Journal of Gastroenterology, 2005, 11, 1876.	1.4	14

#	Article	IF	Citations
91	Endoscopic appearance of GERD: Putative role of cell proliferation. Digestive and Liver Disease, 2007, 39, 713-719.	0.4	13
92	ITA-MNGIE: an Italian regional and national survey for mitochondrial neuro-gastro-intestinal encephalomyopathy. Neurological Sciences, 2016, 37, 1149-1151.	0.9	13
93	Not All Granular Cell Tumors Show Schwann Cell Differentiation: A Granular Cell Leiomyosarcoma of the Thumb, a Case Report. American Journal of Dermatopathology, 1999, 21, 307-309.	0.3	13
94	Early-onset dementia with prolonged occipital seizures. Neurology, 2008, 71, 1709-1712.	1.5	12
95	Wnt activation affects proliferation, invasiveness and radiosensitivity in medulloblastoma. Journal of Neuro-Oncology, 2015, 121, 119-127.	1.4	12
96	Morphological study of TNPO3 and SRSF1 interaction during myogenesis by combining confocal, structured illumination and electron microscopy analysis. Molecular and Cellular Biochemistry, 2021, 476, 1797-1811.	1.4	12
97	Esophageal cell proliferation in gastroesophageal reflux disease: Clinical-morphological data before and after pantoprazole. World Journal of Gastroenterology, 2009, 15, 936.	1.4	12
98	COVID-19 and the Brain: The Neuropathological Italian Experience on 33 Adult Autopsies. Biomolecules, 2022, 12, 629.	1.8	12
99	Secretory meningioma of the middle ear: A light microscopic, immunohistochemical and ultrastructural study of one case. Neuropathology, 2008, 28, 69-73.	0.7	11
100	First evidence of a pathogenic insertion in the NOTCH3 gene causing CADASIL. Journal of Neurology, Neurosurgery and Psychiatry, 2008, 79, 108-110.	0.9	11
101	Response of human chondrocytes and mesenchymal stromal cells to a decellularized human dermis. BMC Musculoskeletal Disorders, 2013, 14, 12.	0.8	11
102	Familial polyglucosan body myopathy with unusual phenotype. Neuropathology and Applied Neurobiology, 2015, 41, 385-390.	1.8	11
103	Transportin 3 (TNPO3) and related proteins in limb girdle muscular dystrophy D2 muscle biopsies: A morphological study and pathogenetic hypothesis. Neuromuscular Disorders, 2020, 30, 685-692.	0.3	11
104	XAV939-Mediated ARTD Activity Inhibition in Human MB Cell Lines. PLoS ONE, 2015, 10, e0124149.	1.1	11
105	Gastrointestinal Autonomic Nerve Tumor (Plexosarcoma). Acta Cytologica, 1998, 42, 1189-1194.	0.7	10
106	Glucocorticoid receptor antagonization propels endogenous cardiomyocyte proliferation and cardiac regeneration. , 2022, 1, 617-633.		10
107	A new polychrome stain and simultaneous methods of histological, histochemical and immunohistochemical stainings performed on semithin sections of Bioacryl-embedded human tissues. The Histochemical Journal, 1993, 25, 670-677.	0.6	9
108	Mucin-Secreting Cellular Ependymoma: A Light and Electron Microscopy Study. Ultrastructural Pathology, 1999, 23, 319-323.	0.4	9

#	Article	IF	Citations
109	ECRH at high heating power density in FTU tokamak. Fusion Engineering and Design, 2001, 53, 301-308.	1.0	9
110	Intracellular Distribution ofÎ²-Catenin in Human Medulloblastoma Cell Lines with Different Degree of Neuronal Differentiation. Ultrastructural Pathology, 2007, 31, 33-44.	0.4	9
111	A novel T137A SOD1 mutation in an Italian family with two subjects affected by amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2011, 12, 385-388.	2.3	9
112	Evidence of enteric angiopathy and neuromuscular hypoxia in patients with mitochondrial neurogastrointestinal encephalomyopathy. American Journal of Physiology - Renal Physiology, 2021, 320, G768-G779.	1.6	9
113	Follicular baso-squamous melanocytic tumour of the skin. Histopathology, 2002, 41, 337-341.	1.6	8
114	Spontaneously occurring intramural coronary arteriosclerosis in regularly slaughtered veal calves and beef cattle: aÂscreening study about prevalence and histopathological features. Journal of Veterinary Cardiology, 2018, 20, 55-63.	0.3	8
115	X-Linked Duchenne-Type Muscular Dystrophy in Jack Russell Terrier Associated with a Partial Deletion of the Canine DMD Gene. Genes, 2020, 11, 1175.	1.0	8
116	Correspondence. Cytopathology, 2000, 11, 193-196.	0.4	7
117	Cell proliferation and ultrastructural changes of the duodenal mucosa of patients affected by familial adenomatous polyposis. Human Pathology, 2004, 35, 622-626.	1.1	7
118	HIV Enteropathy: Undescribed Ultrastructural Changes of Duodenal Mucosa and Their Regression After Triple Antiviral Therapy. A Case Report. Digestive Diseases and Sciences, 2005, 50, 617-622.	1.1	7
119	The Role of Ultrastructural Examination in Storage Diseases. Ultrastructural Pathology, 2010, 34, 243-251.	0.4	7
120	The empowerment of translational research: lessons from laminopathies. Orphanet Journal of Rare Diseases, 2012, 7, 37.	1.2	7
121	Different renal phenotypes in related adult males with Fabry disease with the same classic genotype. Molecular Genetics & Genomic Medicine, 2017, 5, 438-442.	0.6	6
122	Melanin in human vestibular organs: what do we know now? An ultrastructural study and review of the literature. Hearing, Balance and Communication, 2018, 16, 101-107.	0.1	6
123	Histological and ultrastructural evaluation of human decellularized matrix as a hernia repair device. Ultrastructural Pathology, 2018, 42, 32-38.	0.4	5
124	Meningeal Sarcoma with Rhabdomyoblastic Differentiation. Neurosurgery, 1992, 30, 782-785.	0.6	5
125	An Italian case of CADASIL with mutation CGC-TCG in codon 1006, exon 19 Notch3 gene. Neurological Sciences, 2004, 24, 401-406.	0.9	4
126	Myocardial Adenomatoid Tumor in Eight Cattle: Evidence for Mesothelial Origin of Bovine Myocardial Epithelial Inclusions. Veterinary Pathology, 2009, 46, 897-903.	0.8	4

#	Article	IF	Citations
127	Sclerosing Paraganglioma of the Carotid Body: A Potential Pitfall of Malignancy. Head and Neck Pathology, 2015, 9, 300-304.	1.3	4
128	Human immunodeficiency virus type 1 antigen detection in endomyocardial biopsy: an immunomorphological study. Microbiologica, 1990, 13, 145-9.	0.2	4
129	Case 2: Calcium Pyrophosphate Dihydrate Microcrystal-associated Arthropathy. Ultrastructural Pathology, 1986, 10, 395-400.	0.4	3
130	Human cytomegalovirus structural components: intracellular and intraviral localization of p38. Virus Research, 1991, 19, 189-198.	1.1	3
131	Human immunodeficiency virus (HIV) infection of cell cultures. An ultrastructural and immunocytochemical approach.. Cytotechnology, 1991, 5, 82-83.	0.7	3
132	Ultrastructural evidence of ependymal differentiation in a genetically proven atypical teratoid/rhabdoid tumor. Child's Nervous System, 2009, 25, 1627-1631.	0.6	3
133	Juvenile dermatomyositis: A report of three cases. Ultrastructural Pathology, 2016, 40, 83-85.	0.4	3
134	In vitro activity of a partially purified and characterized bark extract ofCastanea sativa Mill. (ENCÂ®) againstChlamydiaspp. Ultrastructural Pathology, 2017, 41, 147-153.	0.4	3
135	The clinical and molecular spectrum of autosomal dominant limb-girdle muscular dystrophies focusing on transportinopathy. Expert Opinion on Orphan Drugs, 2019, 7, 223-232.	0.5	3
136	Potential role of Anitschkow cells in cardiovascular disease in human and veterinary medicine: A review of the literature. Journal of Veterinary Medicine Series C: Anatomia Histologia Embryologia, 2019, 48, 201-206.	0.3	3
137	In situ hybridization at the ultrastructural level: localization of cytomegalovirus DNA using digoxigenin labelled probes. Journal of Submicroscopic Cytology and Pathology, 1993, 25, 341-5.	0.3	3
138	LGMD D2 TNPO3-Related: From Clinical Spectrum to Pathogenetic Mechanism. Frontiers in Neurology, 2022, 13, 840683.	1.1	3
139	Case 1: Primary Metabolic Cardiomyopathy Mimicking an Ischemic Heart Disease. Ultrastructural Pathology, 1986, 10, 387-394.	0.4	2
140	Hereditary sensory and autonomic neuropathy with ataxia and late onset. Clinical Neurology and Neurosurgery, 1994, 96, 191-196.	0.6	2
141	Erosions or not in GORD? The potential role of oesophageal cell proliferation. Gut, 2005, 54, 887-888.	6.1	2
142	Cardiac delayed enhancement distribution in extralysosomial glycogen storage disease. Clinical Imaging, 2008, 32, 474-476.	0.8	2
143	Activity of synthetic peptides against Chlamydia. Biopolymers, 2017, 108, e23032.	1.2	2
144	The role of transmission electron microscopy in vacuole-associated myopathies. Ultrastructural Pathology, 2017, 41, 88-90.	0.4	1

#	Article	IF	Citations
145	Medullocytoma and Glioneurocytoma: Related Tumors?. American Journal of Surgical Pathology, 1997, 21, 615-616.	2.1	1
146	Communications to the Editor and Clinical Case Reports. Vascular Surgery, 1986, 20, 181-187.	0.3	0
147	Vestibular findings in patients with schwannoma of the 8th cranial nerve: a survey of nine cases and review of the literature. Hearing, Balance and Communication, 2018, 16, 166-172.	0.1	0
148	A rare case of intracranial extra-axial ependymoma. Ultrastructural Pathology, 2019, 43, 216-219.	0.4	0
149	A Pathogenic Galactosidase A Mutation Coexisting With an MYBPC3 Mutation in a Female Patient With Hypertrophic Cardiomyopathy. Canadian Journal of Cardiology, 2020, 36, 1554.e1-1554.e3.	0.8	0
150	Ultrastructural and Immunohistochemical Diagnosis of a Neonatal Herpes Simplex Virus Infection Presenting as Fulminant Hepatitis: A Case Report. Advances in Experimental Medicine and Biology, 2021, , .	0.8	0