

# Fausto J Rodriguez

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1754797/publications.pdf>

Version: 2024-02-01

177  
papers

11,139  
citations

47006

47  
h-index

33894

99  
g-index

181  
all docs

181  
docs citations

181  
times ranked

15901  
citing authors

#	ARTICLE	IF	CITATIONS
1	RNA-seq highlights differential regulated pathways involved in cell cycle and inflammation in orbitofacial neurofibromas. <i>Brain Pathology</i> , 2022, 32, e13007.	4.1	2
2	Diffusion MRI is an early biomarker of overall survival benefit in IDH wild-type recurrent glioblastoma treated with immune checkpoint inhibitors. <i>Neuro-Oncology</i> , 2022, 24, 1020-1028.	1.2	12
3	Intracranial mesenchymal tumors with FET-CREB fusion are composed of at least two epigenetic subgroups distinct from meningioma and extracranial sarcomas. <i>Brain Pathology</i> , 2022, 32, e13037.	4.1	11
4	Mutational Landscape and Outcomes of Conjunctival Melanoma in 101 Patients. <i>Ophthalmology</i> , 2022, 129, 679-693.	5.2	16
5	Lynch syndrome caused by a novel deletion of the promoter and exons 1-13 of MLH1 gene. <i>Cancer Genetics</i> , 2022, 262-263, 91-94.	0.4	0
6	Clinical features and surgical outcomes of intracranial and spinal cord subependymomas. <i>Journal of Neurosurgery</i> , 2022, 137, 931-942.	1.6	3
7	Low-grade diffusely infiltrative tumour (LGDIT), SMARCB1-mutant: A clinical and histopathological distinct entity showing epigenetic similarity with ATRX-MYC. <i>Neuropathology and Applied Neurobiology</i> , 2022, 48, .	3.2	5
8	Molecular Biomarker Testing for the Diagnosis of Diffuse Gliomas. <i>Archives of Pathology and Laboratory Medicine</i> , 2022, 146, 547-574.	2.5	25
9	Detection of malignant peripheral nerve sheath tumors in patients with neurofibromatosis using aneuploidy and mutation identification in plasma. <i>ELife</i> , 2022, 11, .	6.0	4
10	Targeting farnesylation as a novel therapeutic approach in HRAS-mutant rhabdomyosarcoma. <i>Oncogene</i> , 2022, 41, 2973-2983.	5.9	9
11	Utility of targeted next-generation sequencing assay to detect 1p/19q co-deletion in formalin-fixed paraffin-embedded glioma specimens. <i>Human Pathology</i> , 2022, 126, 63-76.	2.0	5
12	Therapeutic Vulnerability to ATR Inhibition in Concurrent NF1 and ATRX-Deficient/ALT-Positive High-Grade Solid Tumors. <i>Cancers</i> , 2022, 14, 3015.	3.7	10
13	The WHO classification of tumors of the central nervous system—finally here, and welcome!. <i>Brain Pathology</i> , 2022, 32, .	4.1	2
14	Global microRNA profiling identified miR-10b-5p as a regulator of neurofibromatosis 1 (NF1)-glioma migration. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 96-107.	3.2	10
15	Biology and grading of pleomorphic xanthoastrocytoma—what have we learned about it?. <i>Brain Pathology</i> , 2021, 31, 20-32.	4.1	32
16	Intracranial mesenchymal tumor with FET-CREB fusion—A unifying diagnosis for the spectrum of intracranial myxoid mesenchymal tumors and angiomatoid fibrous histiocytoma-like neoplasms. <i>Brain Pathology</i> , 2021, 31, e12918.	4.1	44
17	Imaging of non-neurogenic peripheral nerve malignancy—a case series and systematic review. <i>Skeletal Radiology</i> , 2021, 50, 201-215.	2.0	5
18	GLI3 Is Associated With Neuronal Differentiation in SHH-Activated and WNT-Activated Medulloblastoma. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 129-136.	1.7	5

#	ARTICLE	IF	CITATIONS
19	Preoperative BMI Predicts Postoperative Weight Gain in Adult-onset Craniopharyngioma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2021, 106, 1603-1617.	3.6	8
20	Diagnostic Pathology of Tumors of Peripheral Nerve. <i>Neurosurgery</i> , 2021, 88, 443-456.	1.1	43
21	Chromosome 8 gain is associated with high-grade transformation in MPNST. <i>JCI Insight</i> , 2021, 6, .	5.0	23
22	Predictors of Postoperative Visual Outcome After Surgical Intervention for Craniopharyngiomas. <i>World Neurosurgery</i> , 2021, 148, e589-e599.	1.3	8
23	SMARCAL1 loss and alternative lengthening of telomeres (ALT) are enriched in giant cell glioblastoma. <i>Modern Pathology</i> , 2021, 34, 1810-1819.	5.5	8
24	Pectic Galactan Polysaccharide-Based Gene Delivery System for Targeting Neuroinflammation. <i>Advanced Functional Materials</i> , 2021, 31, 2100643.	14.9	4
25	Reimagining pilocytic astrocytomas in the context of pediatric low-grade gliomas. <i>Neuro-Oncology</i> , 2021, 23, 1634-1646.	1.2	19
26	Predicting BRAF V600E mutation in glioblastoma: utility of radiographic features. <i>Brain Tumor Pathology</i> , 2021, 38, 228-233.	1.7	9
27	Abstract 2251: Gene expression analysis by RNA-sequencing highlights differential regulated pathways involved in cell cycle and inflammation in orbitofacial neurofibromas. , 2021, , .		0
28	OUP accepted manuscript. <i>American Journal of Clinical Pathology</i> , 2021, , .	0.7	0
29	Conditional reprogramming culture conditions facilitate growth of lower-grade glioma models. <i>Neuro-Oncology</i> , 2021, 23, 770-782.	1.2	18
30	An update on the central nervous system manifestations of neurofibromatosis type 1. <i>Acta Neuropathologica</i> , 2020, 139, 625-641.	7.7	64
31	Intracranial cellular schwannomas: a clinicopathological study of 20 cases. <i>Histopathology</i> , 2020, 76, 275-282.	2.9	9
32	Assessing interobserver variability and accuracy in the histological diagnosis and classification of cutaneous neurofibromas. <i>Neuro-Oncology Advances</i> , 2020, 2, i117-i123.	0.7	3
33	Combined Inhibition of SHP2 and MEK Is Effective in Models of NF1-Deficient Malignant Peripheral Nerve Sheath Tumors. <i>Cancer Research</i> , 2020, 80, 5367-5379.	0.9	29
34	A Case of Metastatic Giant Cell Tumor of Soft Tissue of the Orbit Associated With <i>PALB2</i> Variant. <i>JAMA Ophthalmology</i> , 2020, 138, 1322.	2.5	1
35	Teaching NeuroImages: Intracranial DICER1-associated spindle cell sarcoma in a child. <i>Neurology</i> , 2020, 95, e2176-e2177.	1.1	1
36	Neurogenic Tumors of the Mediastinum. <i>Seminars in Diagnostic Pathology</i> , 2020, 37, 179-186.	1.5	17

#	ARTICLE	IF	CITATIONS
37	Low-Grade Gemistocytic Morphology in H3 G34R-Mutant Gliomas and Concurrent K27M Mutation: Clinicopathologic Findings. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 1038-1043.	1.7	3
38	Differential gene methylation and expression of HOX transcription factor family in orbitofacial neurofibroma. <i>Acta Neuropathologica Communications</i> , 2020, 8, 62.	5.2	7
39	Localized Hypertrophic Neuropathy as a Neoplastic Manifestation of KRAS-Mediated RASopathy. <i>Journal of Neuropathology and Experimental Neurology</i> , 2020, 79, 647-651.	1.7	6
40	Telomere length alterations and ATRX/DAXX loss in pituitary adenomas. <i>Modern Pathology</i> , 2020, 33, 1475-1481.	5.5	13
41	A clinically and genomically annotated nerve sheath tumor biospecimen repository. <i>Scientific Data</i> , 2020, 7, 184.	5.3	19
42	Response to letter to the editor: "All models are wrong; some models are useful". <i>Neuro-Oncology</i> , 2020, 22, 1406-1407.	1.2	0
43	Implications of new understandings of gliomas in children and adults with NF1: report of a consensus conference. <i>Neuro-Oncology</i> , 2020, 22, 773-784.	1.2	44
44	Astrocytic trans-Differentiation Completes a Multicellular Paracrine Feedback Loop Required for Medulloblastoma Tumor Growth. <i>Cell</i> , 2020, 180, 502-520.e19.	28.9	99
45	Intraneural perineurioma in neurofibromatosis type 2 with molecular analysis. , 2020, 39, 167-171.		8
46	MBCL-25. PILOT STUDY OF A SURGERY AND CHEMOTHERAPY-ONLY APPROACH IN THE UPFRONT THERAPY OF CHILDREN WITH WNT-POSITIVE STANDARD RISK MEDULLOBLASTOMA: UPDATED OUTCOMES. <i>Neuro-Oncology</i> , 2020, 22, iii393-iii394.	1.2	3
47	NFB-01. FUNCTIONAL CHARACTERIZATION OF ATRX LOSS IN NF1-ASSOCIATED GLIOMA AND MPNST. <i>Neuro-Oncology</i> , 2020, 22, iii417-iii418.	1.2	0
48	Histopathologic findings in malignant peripheral nerve sheath tumor predict response to radiotherapy and overall survival. <i>Neuro-Oncology Advances</i> , 2020, 2, vdaa131.	0.7	6
49	AI-Assisted <i>In Situ</i> Detection of Human Glioma Infiltration Using a Novel Computational Method for Optical Coherence Tomography. <i>Clinical Cancer Research</i> , 2019, 25, 6329-6338.	7.0	31
50	ATRX Mutations in Pineal Parenchymal Tumors of Intermediate Differentiation. <i>Journal of Neuropathology and Experimental Neurology</i> , 2019, 78, 703-708.	1.7	7
51	Somatostatin Receptor Ligand Therapy "A Potential Therapy for Neurocytoma. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 2395-2402.	3.6	7
52	Telomere alterations in neurofibromatosis type 1-associated solid tumors. <i>Acta Neuropathologica Communications</i> , 2019, 7, 139.	5.2	12
53	ADAM3A copy number gains occur in a subset of conjunctival squamous cell carcinoma and its high grade precursors. <i>Human Pathology</i> , 2019, 94, 92-97.	2.0	5
54	<i>Sleeping Beauty</i> Insertional Mutagenesis Reveals Important Genetic Drivers of Central Nervous System Embryonal Tumors. <i>Cancer Research</i> , 2019, 79, 905-917.	0.9	33

#	ARTICLE	IF	CITATIONS
55	Aquaporin-4 Expression Patterns in Glioblastoma Pre-Chemoradiation and at Time of Suspected Progression. <i>Cancer Investigation</i> , 2019, 37, 67-72.	1.3	4
56	Pathologic and molecular aspects of anaplasia in circumscribed gliomas and glioneuronal tumors. <i>Brain Tumor Pathology</i> , 2019, 36, 40-51.	1.7	9
57	MYD88 L265P mutation and CDKN2A loss are early mutational events in primary central nervous system diffuse large B-cell lymphomas. <i>Blood Advances</i> , 2019, 3, 375-383.	5.2	77
58	MEDU-34. PILOT STUDY OF A SURGERY AND CHEMOTHERAPY-ONLY APPROACH IN THE UPFRONT THERAPY OF CHILDREN WITH WNT-POSITIVE STANDARD RISK MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2019, 21, ii110-ii110.	1.2	10
59	Genomic Landscape of Intramedullary Spinal Cord Gliomas. <i>Scientific Reports</i> , 2019, 9, 18722.	3.3	28
60	Granular cell astrocytoma: an aggressive IDH-wildtype diffuse glioma with molecular genetic features of primary glioblastoma. <i>Brain Pathology</i> , 2019, 29, 193-204.	4.1	7
61	Alternative lengthening of telomeres, ATRX loss and H3K27M mutations in histologically defined pilocytic astrocytoma with anaplasia. <i>Brain Pathology</i> , 2019, 29, 126-140.	4.1	54
62	The consistency of neuropathological diagnoses in patients undergoing surgery for suspected recurrence of glioblastoma. <i>Journal of Neuro-Oncology</i> , 2019, 141, 347-354.	2.9	25
63	Inhibition of enhancer of zest homologue 2 is a potential therapeutic target for high-MYC medulloblastoma. <i>Neuropathology</i> , 2019, 39, 71-77.	1.2	8
64	Cytopathological Analysis in the Diagnosis of Corticotroph Adenomas: Technical Note. <i>Journal of Neurological Surgery, Part B: Skull Base</i> , 2019, 80, .	0.8	0
65	A recurrent kinase domain mutation in PRKCA defines chordoid glioma of the third ventricle. <i>Nature Communications</i> , 2018, 9, 810.	12.8	56
66	Hemophagocytic Lymphohistiocytosis in Adults with Intraocular Involvement: Clinicopathologic Features of 3 Cases. <i>Ocular Oncology and Pathology</i> , 2018, 4, 1-11.	1.0	9
67	DNA methylation-based classification of central nervous system tumours. <i>Nature</i> , 2018, 555, 469-474.	27.8	1,872
68	Anaplastic astrocytoma with piloid features, a novel molecular class of IDH wildtype glioma with recurrent MAPK pathway, CDKN2A/B and ATRX alterations. <i>Acta Neuropathologica</i> , 2018, 136, 273-291.	7.7	190
69	Recurrent copy number alterations in low-grade and anaplastic pleomorphic xanthoastrocytoma with and without BRAF V600E mutation. <i>Brain Pathology</i> , 2018, 28, 172-182.	4.1	64
70	GENE-42. THE GENOMIC LANDSCAPE OF TRIPLE-NEGATIVE GLIOBLASTOMA. <i>Neuro-Oncology</i> , 2018, 20, vi112-vi112.	1.2	0
71	PDCT-02. COMBINED INHIBITION OF MTORC1/C2 AND MEK PATHWAY IS SYNERGISTIC IN PRECLINICAL TESTING OF PEDIATRIC LOW-GRADE GLIOMA INCLUDING A NOVEL PATIENT-DERIVED NF1 PILOCYTIC ASTROCYTOMA CELL LINE. <i>Neuro-Oncology</i> , 2018, 20, vi200-vi201.	1.2	0
72	Central Nervous System-type Neuroepithelial Tumors and Tumor-like Proliferations Developing in the Gynecologic Tract and Pelvis. <i>American Journal of Surgical Pathology</i> , 2018, 42, 1429-1444.	3.7	18

#	ARTICLE	IF	CITATIONS
73	GENE-01. THE GENOMIC LANDSCAPE OF TRIPLE-NEGATIVE GLIOBLASTOMA. <i>Neuro-Oncology</i> , 2018, 20, vi102-vi103.	1.2	0
74	RARE-08. GRADING CONSIDERATIONS FOR MENINGEAL SOLITARY FIBROUS TUMOR/HEMANGIOPERICYTOMA. <i>Neuro-Oncology</i> , 2018, 20, vi237-vi238.	1.2	1
75	Anterior Cranial Fossa Calcifying Pseudoneoplasm of the Neuroaxis—Diagnosis Using a Transblepharoplasty Approach. <i>Journal of Neurological Surgery Reports</i> , 2018, 79, e75-e78.	0.6	5
76	Expression of renal cell markers and detection of 3p loss links endolymphatic sac tumor to renal cell carcinoma and warrants careful evaluation to avoid diagnostic pitfalls. <i>Acta Neuropathologica Communications</i> , 2018, 6, 107.	5.2	7
77	Malignant Peripheral Nerve Sheath Tumors Show Decreased Global DNA Methylation. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018, 77, 958-963.	1.7	9
78	The genomic landscape of TERT promoter wildtype-IDH wildtype glioblastoma. <i>Nature Communications</i> , 2018, 9, 2087.	12.8	124
79	Neuropathology Education Using Social Media. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018, 77, 454-460.	1.7	13
80	Subependymal giant cell astrocytoma-like astrocytoma: a neoplasm with a distinct phenotype and frequent neurofibromatosis type-1-association. <i>Modern Pathology</i> , 2018, 31, 1787-1800.	5.5	24
81	MicroRNA (miR) 125b regulates cell growth and invasion in pediatric low grade glioma. <i>Scientific Reports</i> , 2018, 8, 12506.	3.3	30
82	A multiprotein supercomplex controlling oncogenic signalling in lymphoma. <i>Nature</i> , 2018, 560, 387-391.	27.8	276
83	MYD88 L265P mutation and CDKN2A loss as early mutational events in primary central nervous system lymphomas. <i>Journal of Clinical Oncology</i> , 2018, 36, e14041-e14041.	1.6	1
84	Epstein-Barr virus-associated smooth muscle tumor of the cavernous sinus: a delayed complication of allogeneic peripheral blood stem cell transplantation: case report. <i>Journal of Neurosurgery</i> , 2017, 126, 1479-1483.	1.6	12
85	Clinicopathological features of peripheral nerve sheath tumors involving the eye and ocular adnexa. <i>Human Pathology</i> , 2017, 63, 70-78.	2.0	22
86	miRNA Regulation in Gliomas: Usual Suspects in Glial Tumorigenesis and Evolving Clinical Applications. <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 246-254.	1.7	25
87	Comparative volumetric analysis of the extent of resection of molecularly and histologically distinct low grade gliomas and its role on survival. <i>Journal of Neuro-Oncology</i> , 2017, 134, 65-74.	2.9	46
88	Differential neuronal susceptibility and apoptosis in congenital Zika virus infection. <i>Annals of Neurology</i> , 2017, 82, 121-127.	5.3	31
89	Absence of Cytomegalovirus in Glioblastoma and Other High-grade Gliomas by Real-time PCR, Immunohistochemistry, and In Situ Hybridization. <i>Clinical Cancer Research</i> , 2017, 23, 3150-3157.	7.0	52
90	Immunohistochemical analysis of H3K27me3 demonstrates global reduction in group-A childhood posterior fossa ependymoma and is a powerful predictor of outcome. <i>Acta Neuropathologica</i> , 2017, 134, 705-714.	7.7	168

#	ARTICLE	IF	CITATIONS
91	The efficacy of lapatinib and nilotinib in combination with radiation therapy in a model of NF2 associated peripheral schwannoma. <i>Journal of Neuro-Oncology</i> , 2017, 135, 47-56.	2.9	10
92	Late post-treatment radiographic changes 3 years following chemoradiation for glioma: the importance of histopathology. <i>CNS Oncology</i> , 2017, 6, 195-201.	3.0	4
93	HIF-1 $\alpha$ - Targeting Acriflavine Provides Long Term Survival and Radiological Tumor Response in Brain Cancer Therapy. <i>Scientific Reports</i> , 2017, 7, 14978.	3.3	62
94	MicroRNA profiling of low-grade glial and glioneuronal tumors shows an independent role for cluster 14q32.31 member miR-487b. <i>Modern Pathology</i> , 2017, 30, 204-216.	5.5	37
95	Low-grade Schwann cell neoplasms with leptomeningeal dissemination: clinicopathologic and autopsy findings. <i>Human Pathology</i> , 2017, 60, 121-128.	2.0	6
96	Differential Neuronal Susceptibility and Apoptosis in Congenital Zika Virus Infection. <i>Open Forum Infectious Diseases</i> , 2017, 4, S56-S56.	0.9	0
97	PDTB-11. DISRUPTING THE EPIGENETIC MODIFIER HMGA2 IN LETHAL PEDIATRIC AND ADULT GLIOMAS INHIBITS INVASION, GROWTH AND TUMORIGENICITY. <i>Neuro-Oncology</i> , 2016, 18, vi152-vi152.	1.2	0
98	MPTH-48. INTER-INTERPRETER CONSISTENCY IN REVIEWING HISTOPATHOLOGY FROM PATIENTS WITH GLIOBLASTOMA AND RADIOGRAPHIC PROGRESSION FOLLOWING STANDARD RADIATION AND TEMOZOLOMIDE. <i>Neuro-Oncology</i> , 2016, 18, vi116-vi116.	1.2	0
99	Molecular Analysis of Pediatric Oligodendrogliomas Highlights Genetic Differences with Adult Counterparts and Other Pediatric Gliomas. <i>Brain Pathology</i> , 2016, 26, 206-214.	4.1	25
100	Compound gonadotrophic pituitary adenoma and rhabdomyosarcoma. <i>Histopathology</i> , 2016, 68, 1111-1114.	2.9	4
101	Clinicopathological Features of Ophthalmic Neoplasms Arising in the Setting of Xeroderma Pigmentosum. <i>Ocular Oncology and Pathology</i> , 2016, 2, 112-121.	1.0	10
102	Zika Virus Infection with Prolonged Maternal Viremia and Fetal Brain Abnormalities. <i>New England Journal of Medicine</i> , 2016, 374, 2142-2151.	27.0	754
103	CSF1 Overexpression Promotes High-Grade Glioma Formation without Impacting the Polarization Status of Glioma-Associated Microglia and Macrophages. <i>Cancer Research</i> , 2016, 76, 2552-2560.	0.9	69
104	Recent Advances on the Molecular Pathology of Glial Neoplasms in Children and Adults. <i>Journal of Molecular Diagnostics</i> , 2016, 18, 620-634.	2.8	42
105	N for nucleus in neurofibromin: new role for an old tumor suppressor?. <i>Journal of Neurochemistry</i> , 2016, 136, 11-12.	3.9	0
106	Expanded Endonasal Endoscopic Approach for Resection of an Infraselar Craniopharyngioma. <i>World Neurosurgery</i> , 2016, 95, 618.e7-618.e12.	1.3	9
107	Distinct patterns of primary and motile cilia in Rathke's cleft cysts and craniopharyngioma subtypes. <i>Modern Pathology</i> , 2016, 29, 1446-1459.	5.5	15
108	Frequent alternative lengthening of telomeres and ATRX loss in adult NF1-associated diffuse and high-grade astrocytomas. <i>Acta Neuropathologica</i> , 2016, 132, 761-763.	7.7	23

#	ARTICLE	IF	CITATIONS
109	Next-generation sequencing in neuropathologic diagnosis of infections of the nervous system. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2016, 3, e251.	6.0	142
110	Migration Phenotype of Brain-Cancer Cells Predicts Patient Outcomes. <i>Cell Reports</i> , 2016, 15, 2616-2624.	6.4	63
111	Histologically benign, clinically aggressive: Progressive non- $\epsilon$ optic pathway pilocytic astrocytomas in adults with NF1. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 1455-1461.	1.2	16
112	MYB-QKI rearrangements in angiocentric glioma drive tumorigenicity through a tripartite mechanism. <i>Nature Genetics</i> , 2016, 48, 273-282.	21.4	214
113	Diffusion tensor imaging suggests extrapontine extension of pediatric diffuse intrinsic pontine gliomas. <i>European Journal of Radiology</i> , 2016, 85, 700-706.	2.6	10
114	Pathologic deposition of non-amyloid immunoglobulin in the brain leading to mass effect and neurological deficits. <i>Journal of Clinical Neuroscience</i> , 2016, 30, 143-145.	1.5	3
115	Inter-interpreter reliability of neuropathological assessment of disease status after early repeat resection for suspected recurrent glioblastoma. <i>Journal of Clinical Oncology</i> , 2016, 34, 2050-2050.	1.6	0
116	Clinicopathologic implications of NF1 gene alterations in diffuse gliomas. <i>Human Pathology</i> , 2015, 46, 1323-1330.	2.0	25
117	Pleomorphic Xanthoastrocytoma: Natural History and Long-Term Follow-Up. <i>Brain Pathology</i> , 2015, 25, 575-586.	4.1	188
118	Src family kinases differentially influence glioma growth and motility. <i>Molecular Oncology</i> , 2015, 9, 1783-1798.	4.6	52
119	Detection of human brain cancer infiltration ex vivo and in vivo using quantitative optical coherence tomography. <i>Science Translational Medicine</i> , 2015, 7, 292ra100.	12.4	247
120	High rate of concurrent BRAF-KIAA1549 gene fusion and 1p deletion in disseminated oligodendroglioma-like leptomeningeal neoplasms (DOLN). <i>Acta Neuropathologica</i> , 2015, 129, 609-610.	7.7	95
121	A clinicopathologic study of diencephalic pediatric low-grade gliomas with BRAF V600 mutation. <i>Acta Neuropathologica</i> , 2015, 130, 575-585.	7.7	50
122	Genetic Profiling by Single-Nucleotide Polymorphism-Based Array Analysis Defines Three Distinct Subtypes of Orbital Meningioma. <i>Brain Pathology</i> , 2015, 25, 193-201.	4.1	19
123	Sleeping Beauty Mouse Models Identify Candidate Genes Involved in Gliomagenesis. <i>PLoS ONE</i> , 2014, 9, e113489.	2.5	21
124	Increased 5-hydroxymethylcytosine and decreased 5-methylcytosine are indicators of global epigenetic dysregulation in diffuse intrinsic pontine glioma. <i>Acta Neuropathologica Communications</i> , 2014, 2, 59.	5.2	35
125	Clinicopathologic Features of Pediatric Oligodendrogliomas. <i>American Journal of Surgical Pathology</i> , 2014, 38, 1058-1070.	3.7	57
126	Genetic and pathologic evolution of early secondary gliosarcoma. <i>Brain Tumor Pathology</i> , 2014, 31, 40-46.	1.7	10



#	ARTICLE	IF	CITATIONS
127	Exome sequencing identifies BRAF mutations in papillary craniopharyngiomas. <i>Nature Genetics</i> , 2014, 46, 161-165.	21.4	408
128	Cytologic features in vitreous preparations of patients with suspicion of intraocular lymphoma. <i>Diagnostic Cytopathology</i> , 2014, 42, 37-44.	1.0	28
129	Incidental parenchymal magnetic resonance imaging findings in the brains of patients with neurofibromatosis type 2. <i>NeuroImage: Clinical</i> , 2014, 4, 258-265.	2.7	11
130	Pleomorphic xanthoastrocytoma: report of two cases with unconventional clinical presentations. , 2014, 33, 380-387.		16
131	An 80-year experience with optic nerve glioma cases at the Armed Forces Institute of Pathology: evolution from museum to molecular evaluation suggests possible interventions in the cellular senescence and microglial pathways (an American Ophthalmological Society thesis). <i>Transactions of the American Ophthalmological Society</i> , 2014, 112, 11-25.	1.4	2
132	Molecular and Morphologic Correlates of the Alternative Lengthening of Telomeres Phenotype in High-Grade Astrocytomas. <i>Brain Pathology</i> , 2013, 23, 237-243.	4.1	73
133	Immunohistochemistry is highly sensitive and specific for detection of BRAF V600E mutation in pleomorphic xanthoastrocytoma. <i>Acta Neuropathologica Communications</i> , 2013, 1, 20.	5.2	52
134	cMYC expression in infiltrating gliomas: associations with IDH1 mutations, clinicopathologic features and outcome. <i>Journal of Neuro-Oncology</i> , 2013, 115, 249-259.	2.9	28
135	MicroRNA profiling in pediatric pilocytic astrocytoma reveals biologically relevant targets, including PBX3, NFIB, and METAP2. <i>Neuro-Oncology</i> , 2013, 15, 69-82.	1.2	56
136	Cellular pleomorphism in papillary tumors of the pineal region. <i>Brain Tumor Pathology</i> , 2013, 30, 93-98.	1.7	10
137	Exomic Sequencing of Four Rare Central Nervous System Tumor Types. <i>Oncotarget</i> , 2013, 4, 572-583.	1.8	69
138	<i>BRAF</i> Alterations in Primary Glial and Glioneuronal Neoplasms of the Central Nervous System With Identification of 2 Novel KIAA1549. <i>Journal of Neuropathology and Experimental Neurology</i> , 2012, 71, 66-72.	1.7	147
139	<i>BRAF</i> Duplications and MAPK Pathway Activation Are Frequent in Gliomas of the Optic Nerve Proper. <i>Journal of Neuropathology and Experimental Neurology</i> , 2012, 71, 789-795.	1.7	59
140	Diagnostic neuropathology of tumors of the central nervous system. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2012, 104, 77-107.	1.8	6
141	Disseminated oligodendroglial-like leptomeningeal tumor of childhood: a distinctive clinicopathologic entity. <i>Acta Neuropathologica</i> , 2012, 124, 627-641.	7.7	143
142	Frequent <i>ATRX</i> , <i>CIC</i> , <i>FUBP1</i> and <i>IDH1</i> mutations refine the classification of malignant gliomas. <i>Oncotarget</i> , 2012, 3, 709-722.	1.8	532
143	Crystal-storing histiocytosis: An unusual relapsing inflammatory CNS disorder. <i>Multiple Sclerosis and Related Disorders</i> , 2012, 1, 95-99.	2.0	8
144	Genetic predisposition to peripheral nerve neoplasia: diagnostic criteria and pathogenesis of neurofibromatoses, Carney complex, and related syndromes. <i>Acta Neuropathologica</i> , 2012, 123, 349-367.	7.7	74

#	ARTICLE	IF	CITATIONS
145	Pathology of peripheral nerve sheath tumors: diagnostic overview and update on selected diagnostic problems. <i>Acta Neuropathologica</i> , 2012, 123, 295-319.	7.7	525
146	Peripheral nerve sheath tumors: the elegant chapter in surgical neuropathology. <i>Acta Neuropathologica</i> , 2012, 123, 293-294.	7.7	7
147	Neoplastic cells are a rare component in human glioblastoma microvasculature. <i>Oncotarget</i> , 2012, 3, 98-106.	1.8	79
148	Neurofibromatosis-1 Heterozygosity Increases Microglia in a Spatially and Temporally Restricted Pattern Relevant to Mouse Optic Glioma Formation and Growth. <i>Journal of Neuropathology and Experimental Neurology</i> , 2011, 70, 51-62.	1.7	110
149	PI3K/AKT pathway alterations are associated with clinically aggressive and histologically anaplastic subsets of pilocytic astrocytoma. <i>Acta Neuropathologica</i> , 2011, 121, 407-420.	7.7	118
150	Superficial neurofibromas in the setting of schwannomatosis: nosologic implications. <i>Acta Neuropathologica</i> , 2011, 121, 663-668.	7.7	8
151	Altered Telomeres in Tumors with <i>ATR</i> X and <i>DAXX</i> Mutations. <i>Science</i> , 2011, 333, 425-425.	12.6	891
152	Anaplasia in Pilocytic Astrocytoma Predicts Aggressive Behavior. <i>American Journal of Surgical Pathology</i> , 2010, 34, 147-160.	3.7	120
153	Oligodendroglial tumors: diagnostic and molecular pathology. <i>Seminars in Diagnostic Pathology</i> , 2010, 27, 136-145.	1.5	17
154	Phenotypic variations in NF1-associated low grade astrocytomas: possible role for increased mTOR activation in a subset. <i>International Journal of Clinical and Experimental Pathology</i> , 2010, 4, 43-57.	0.5	19
155	Interphase Cytogenetics for 1p19q and t(1;19)(q10;p10) may Distinguish Prognostically Relevant Subgroups in Extraventricular Neurocytoma. <i>Brain Pathology</i> , 2009, 19, 623-629.	4.1	58
156	Ectopic Low-grade Adrenocortical Carcinoma in the Spinal Region. <i>American Journal of Surgical Pathology</i> , 2009, 33, 142-148.	3.7	26
157	Malignant Peripheral Nerve Sheath Tumors of Cranial Nerves and Intracranial Contents. <i>American Journal of Surgical Pathology</i> , 2009, 33, 325-338.	3.7	127
158	Ependymoma and intraparenchymal calcifying pseudoneoplasm of the neural axis: incidental collision or unique reactive phenomenon?. <i>Acta Neuropathologica</i> , 2008, 115, 363-366.	7.7	48
159	Epithelial and pseudoepithelial differentiation in glioblastoma and gliosarcoma. <i>Cancer</i> , 2008, 113, 2779-2789.	4.1	78
160	Immunoglobulin derived depositions in the nervous system: novel mass spectrometry application for protein characterization in formalin-fixed tissues. <i>Laboratory Investigation</i> , 2008, 88, 1024-1037.	3.7	103
161	MGMT Immunohistochemical Expression and Promoter Methylation in Human Glioblastoma. <i>Applied Immunohistochemistry and Molecular Morphology</i> , 2008, 16, 59-65.	1.2	105
162	Gliomas in Neurofibromatosis Type 1: A Clinicopathologic Study of 100 Patients. <i>Journal of Neuropathology and Experimental Neurology</i> , 2008, 67, 240-249.	1.7	162

#	ARTICLE	IF	CITATIONS
163	Gene Expression Profiling of NF-1-Associated and Sporadic Pilocytic Astrocytoma Identifies Aldehyde Dehydrogenase 1 Family Member L1 (ALDH1L1) as an Underexpressed Candidate Biomarker in Aggressive Subtypes. <i>Journal of Neuropathology and Experimental Neurology</i> , 2008, 67, 1194-1204.	1.7	43
164	Ependymal Tumors With Sarcomatous Change (â€œEpendymosarcomaâ€): A Clinicopathologic and Molecular Cytogenetic Study. <i>American Journal of Surgical Pathology</i> , 2008, 32, 699-709.	3.7	32
165	Galectin-3 Expression is Ubiquitous in Tumors of the Sellar Region, Nervous System, and Mimics. <i>American Journal of Surgical Pathology</i> , 2008, 32, 1344-1352.	3.7	21
166	Experimental gliomas in mice using the Sleeping Beauty (SB) transposon system: neuropathologic aspects. <i>FASEB Journal</i> , 2008, 22, 172.4.	0.5	1
167	Massive Sellar and Parasellar Schwannoma. <i>Archives of Neurology</i> , 2007, 64, 1198.	4.5	7
168	Gliosarcoma Arising in Oligodendroglial Tumors (â€œOligosarcomaâ€). <i>American Journal of Surgical Pathology</i> , 2007, 31, 351-362.	3.7	50
169	The Spectrum of Malignancy in Craniopharyngioma. <i>American Journal of Surgical Pathology</i> , 2007, 31, 1020-1028.	3.7	70
170	Pulmonary Chondroma: A Tumor Associated With Carney Triad and Different From Pulmonary Hamartoma. <i>American Journal of Surgical Pathology</i> , 2007, 31, 1844-1853.	3.7	72
171	Histopathologic grading of adult medulloblastomas. <i>Cancer</i> , 2007, 109, 2557-2565.	4.1	34
172	Low grade malignant peripheral nerve sheath tumor with smooth muscle differentiation. <i>Acta Neuropathologica</i> , 2007, 113, 705-709.	7.7	24
173	Gliosarcoma arising in ependymomas (â€œependymosarcomaâ€): a clinicopathologic study. <i>FASEB Journal</i> , 2007, 21, A395.	0.5	0
174	Anaplastic medulloblastoma with granular cell change. <i>Acta Neuropathologica</i> , 2006, 113, 95-99.	7.7	7
175	Unusual malignant glioneuronal tumors of the cerebrum of adults: a clinicopathologic study of three cases. <i>Acta Neuropathologica</i> , 2006, 112, 727-737.	7.7	21
176	Venous congestive myelopathy: a mimic of neoplasia. <i>Modern Pathology</i> , 2005, 18, 710-718.	5.5	34
177	Fine-needle aspiration cytology findings from a case of pancreatic heterotopia at the gastroesophageal junction. <i>Diagnostic Cytopathology</i> , 2004, 31, 175-179.	1.0	34