

Arie Perry

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47,111
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L-index

#	Paper	IF	Citations
419	The 2016 World Health Organization Classification of Tumors of the Central Nervous System: a summary. <i>Acta Neuropathologica</i> , 2016 , 131, 803-20	14.3	8580
418	Glioma Groups Based on 1p/19q, IDH, and TERT Promoter Mutations in Tumors. <i>New England Journal of Medicine</i> , 2015 , 372, 2499-508	59.2	1181
417	DNA methylation-based classification of central nervous system tumours. <i>Nature</i> , 2018 , 555, 469-474	50.4	992
416	Alterations of chromosome arms 1p and 19q as predictors of survival in oligodendrogliomas, astrocytomas, and mixed oligoastrocytomas. <i>Journal of Clinical Oncology</i> , 2000 , 18, 636-45	2.2	899
415	The 2021 WHO Classification of Tumors of the Central Nervous System: a summary. <i>Neuro-Oncology</i> , 2021 , 23, 1231-1251	1	708
414	Meningioma grading: an analysis of histologic parameters. <i>American Journal of Surgical Pathology</i> , 1997 , 21, 1455-65	6.7	500
413	New Brain Tumor Entities Emerge from Molecular Classification of CNS-PNETs. <i>Cell</i> , 2016 , 164, 1060-1073	36.2	483
412	Primary brain tumours in adults. <i>Lancet, The</i> , 2018 , 392, 432-446	40	469
411	Genome sequencing of SHH medulloblastoma predicts genotype-related response to smoothed inhibition. <i>Cancer Cell</i> , 2014 , 25, 393-405	24.3	469
410	International Society Of Neuropathology--Haarlem consensus guidelines for nervous system tumor classification and grading. <i>Brain Pathology</i> , 2014 , 24, 429-35	6	408
409	"Malignancy" in meningiomas: a clinicopathologic study of 116 patients, with grading implications. <i>Cancer</i> , 1999 , 85, 2046-56	6.4	406
408	Pathology of peripheral nerve sheath tumors: diagnostic overview and update on selected diagnostic problems. <i>Acta Neuropathologica</i> , 2012 , 123, 295-319	14.3	401
407	Medulloblastoma: clinicopathological correlates of SHH, WNT, and non-SHH/WNT molecular subgroups. <i>Acta Neuropathologica</i> , 2011 , 121, 381-96	14.3	376
406	Malignancy in meningiomas. <i>Cancer</i> , 1999 , 85, 2046-2056	6.4	374
405	cIMPACT-NOW update 3: recommended diagnostic criteria for "Diffuse astrocytic glioma, IDH-wildtype, with molecular features of glioblastoma, WHO grade IV". <i>Acta Neuropathologica</i> , 2018 , 136, 805-810	14.3	367
404	Histological classification and molecular genetics of meningiomas. <i>Lancet Neurology, The</i> , 2006 , 5, 1045-54	54.1	351
403	Notch1 and notch2 have opposite effects on embryonal brain tumor growth. <i>Cancer Research</i> , 2004 , 64, 7787-93	10.1	339

402	Localization of common deletion regions on 1p and 19q in human gliomas and their association with histological subtype. <i>Oncogene</i> , 1999 , 18, 4144-52	9.2	311
401	Evaluating pediatric brain tumor cellularity with diffusion-tensor imaging. <i>American Journal of Roentgenology</i> , 2001 , 177, 449-54	5.4	307
400	Pathological classification and molecular genetics of meningiomas. <i>Journal of Neuro-Oncology</i> , 2010 , 99, 379-91	4.8	289
399	Orally administered colony stimulating factor 1 receptor inhibitor PLX3397 in recurrent glioblastoma: an Ivy Foundation Early Phase Clinical Trials Consortium phase II study. <i>Neuro-Oncology</i> , 2016 , 18, 557-64	1	284
398	Diffuse Midline Gliomas with Histone H3-K27M Mutation: A Series of 47 Cases Assessing the Spectrum of Morphologic Variation and Associated Genetic Alterations. <i>Brain Pathology</i> , 2016 , 26, 569-80	6	243
397	Primarily resected meningiomas: outcome and prognostic factors in 581 Mayo Clinic patients, 1978 through 1988. <i>Mayo Clinic Proceedings</i> , 1998 , 73, 936-42	6.4	232
396	Integrated analysis of pediatric glioblastoma reveals a subset of biologically favorable tumors with associated molecular prognostic markers. <i>Acta Neuropathologica</i> , 2015 , 129, 669-78	14.3	220
395	Molecular pathogenesis of meningiomas. <i>Journal of Neuro-Oncology</i> , 2004 , 70, 183-202	4.8	220
394	Rituximab-associated progressive multifocal leukoencephalopathy in rheumatoid arthritis. <i>Archives of Neurology</i> , 2011 , 68, 1156-64		215
393	Genetic alterations in uncommon low-grade neuroepithelial tumors: BRAF, FGFR1, and MYB mutations occur at high frequency and align with morphology. <i>Acta Neuropathologica</i> , 2016 , 131, 833-45	14.3	209
392	Optic nerve glioma in mice requires astrocyte Nf1 gene inactivation and Nf1 brain heterozygosity. <i>Cancer Research</i> , 2003 , 63, 8573-7	10.1	199
391	Immunogenomics of Hypermutated Glioblastoma: A Patient with Germline POLE Deficiency Treated with Checkpoint Blockade Immunotherapy. <i>Cancer Discovery</i> , 2016 , 6, 1230-1236	24.4	196
390	cIMPACT-NOW update 6: new entity and diagnostic principle recommendations of the cIMPACT-Utrecht meeting on future CNS tumor classification and grading. <i>Brain Pathology</i> , 2020 , 30, 844-856	6	196
389	Cancer therapy-associated CNS neuropathology: an update and review of the literature. <i>Acta Neuropathologica</i> , 2006 , 111, 197-212	14.3	194
388	Adult infiltrating gliomas with WHO 2016 integrated diagnosis: additional prognostic roles of ATRX and TERT. <i>Acta Neuropathologica</i> , 2017 , 133, 1001-1016	14.3	185
387	A phase I trial of the MEK inhibitor selumetinib (AZD6244) in pediatric patients with recurrent or refractory low-grade glioma: a Pediatric Brain Tumor Consortium (PBTC) study. <i>Neuro-Oncology</i> , 2017 , 19, 1135-1144	1	180
386	The immunophenotypic spectrum of meningeal hemangiopericytoma: a comparison with fibrous meningioma and solitary fibrous tumor of meninges. <i>American Journal of Surgical Pathology</i> , 1997 , 21, 1354-60	6.7	180
385	The prognostic significance of MIB-1, p53, and DNA flow cytometry in completely resected primary meningiomas 1998 , 82, 2262-2269		178

384	Adult medulloblastoma comprises three major molecular variants. <i>Journal of Clinical Oncology</i> , 2011 , 29, 2717-23	2.2	176
383	Soft-tissue perineurioma. Evidence for an abnormality of chromosome 22, criteria for diagnosis, and review of the literature. <i>American Journal of Surgical Pathology</i> , 1997 , 21, 164-73	6.7	176
382	Integrative genomic analysis identifies NDRG2 as a candidate tumor suppressor gene frequently inactivated in clinically aggressive meningioma. <i>Cancer Research</i> , 2005 , 65, 7121-6	10.1	174
381	Large-scale molecular comparison of human schwann cells to malignant peripheral nerve sheath tumor cell lines and tissues. <i>Cancer Research</i> , 2006 , 66, 2584-91	10.1	171
380	cIMPACT-NOW update 5: recommended grading criteria and terminologies for IDH-mutant astrocytomas. <i>Acta Neuropathologica</i> , 2020 , 139, 603-608	14.3	170
379	Histopathologic evaluation of atypical neurofibromatous tumors and their transformation into malignant peripheral nerve sheath tumor in patients with neurofibromatosis 1-a consensus overview. <i>Human Pathology</i> , 2017 , 67, 1-10	3.7	169
378	Prognostic value of 1p, 19q, 9p, 10q, and EGFR-FISH analyses in recurrent oligodendrogliomas. <i>Journal of Neuropathology and Experimental Neurology</i> , 2004 , 63, 314-22	3.1	169
377	Pituitary blastoma: a pathognomonic feature of germ-line DICER1 mutations. <i>Acta Neuropathologica</i> , 2014 , 128, 111-22	14.3	160
376	Gene expression profile identifies tyrosine kinase c-Met as a targetable mediator of antiangiogenic therapy resistance. <i>Clinical Cancer Research</i> , 2013 , 19, 1773-83	12.9	160
375	"Rhabdoid" meningioma: an aggressive variant. <i>American Journal of Surgical Pathology</i> , 1998 , 22, 1482-90	6.7	160
374	Spectrum and prevalence of genetic predisposition in medulloblastoma: a retrospective genetic study and prospective validation in a clinical trial cohort. <i>Lancet Oncology</i> , 2018 , 19, 785-798	21.7	159
373	Aggressive phenotypic and genotypic features in pediatric and NF2-associated meningiomas: a clinicopathologic study of 53 cases. <i>Journal of Neuropathology and Experimental Neurology</i> , 2001 , 60, 994-1003	3.1	156
372	Glioblastoma. <i>Archives of Pathology and Laboratory Medicine</i> , 2007 , 131, 397-406	5	155
371	Embryonal tumor with abundant neuropil and true rosettes (ETANTR), ependymoblastoma, and medulloepithelioma share molecular similarity and comprise a single clinicopathological entity. <i>Acta Neuropathologica</i> , 2014 , 128, 279-89	14.3	152
370	Distinct genetic signatures among pilocytic astrocytomas relate to their brain region origin. <i>Cancer Research</i> , 2007 , 67, 890-900	10.1	148
369	Molecular diagnosis of Ewing sarcoma/primitive neuroectodermal tumor in routinely processed tissue: a comparison of two FISH strategies and RT-PCR in malignant round cell tumors. <i>Modern Pathology</i> , 2006 , 19, 1-8	9.8	144
368	Distinct neural stem cell populations give rise to disparate brain tumors in response to N-MYC. <i>Cancer Cell</i> , 2012 , 21, 601-613	24.3	141
367	Significance of necrosis in grading of oligodendroglial neoplasms: a clinicopathologic and genetic study of newly diagnosed high-grade gliomas. <i>Journal of Clinical Oncology</i> , 2006 , 24, 5419-26	2.2	140

366	Gliomas in neurofibromatosis type 1: a clinicopathologic study of 100 patients. <i>Journal of Neuropathology and Experimental Neurology</i> , 2008 , 67, 240-9	3.1	137
365	Oligodendrogliomas with neurocytic differentiation. A report of 4 cases with diagnostic and histogenetic implications. <i>Journal of Neuropathology and Experimental Neurology</i> , 2002 , 61, 947-55	3.1	136
364	Merlin, DAL-1, and progesterone receptor expression in clinicopathologic subsets of meningioma: a correlative immunohistochemical study of 175 cases. <i>Journal of Neuropathology and Experimental Neurology</i> , 2000 , 59, 872-9	3.1	132
363	Malignant gliomas with primitive neuroectodermal tumor-like components: a clinicopathologic and genetic study of 53 cases. <i>Brain Pathology</i> , 2009 , 19, 81-90	6	129
362	Oligodendroglioma: pathology, molecular mechanisms and markers. <i>Acta Neuropathologica</i> , 2015 , 129, 809-27	14.3	128
361	Transcriptomic versus chromosomal prognostic markers and clinical outcome in uveal melanoma. <i>Clinical Cancer Research</i> , 2007 , 13, 1466-71	12.9	127
360	INI1 expression is retained in composite rhabdoid tumors, including rhabdoid meningiomas. <i>Modern Pathology</i> , 2005 , 18, 951-8	9.8	126
359	Preclinical cancer therapy in a mouse model of neurofibromatosis-1 optic glioma. <i>Cancer Research</i> , 2008 , 68, 1520-8	10.1	120
358	Intermediate-risk meningioma: initial outcomes from NRG Oncology RTOG 0539. <i>Journal of Neurosurgery</i> , 2018 , 129, 35-47	3.2	119
357	Targeted next-generation sequencing of pediatric neuro-oncology patients improves diagnosis, identifies pathogenic germline mutations, and directs targeted therapy. <i>Neuro-Oncology</i> , 2017 , 19, 699-709	7.0	118
356	Histologic classification of gliomas. <i>Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn</i> , 2016 , 134, 71-95	3	118
355	Clinicopathologic and genetic profile of intracranial marginal zone lymphoma: a primary low-grade CNS lymphoma that mimics meningioma. <i>Journal of Clinical Oncology</i> , 2005 , 23, 5718-27	2.2	118
354	Loss of neurofibromatosis 1 (NF1) gene expression in NF1-associated pilocytic astrocytomas. <i>Neuropathology and Applied Neurobiology</i> , 2000 , 26, 361-7	5.2	118
353	Blocking CXCR4-mediated cyclic AMP suppression inhibits brain tumor growth in vivo. <i>Cancer Research</i> , 2007 , 67, 651-8	10.1	116
352	Chromosome 1p and 14q FISH analysis in clinicopathologic subsets of meningioma: diagnostic and prognostic implications. <i>Journal of Neuropathology and Experimental Neurology</i> , 2001 , 60, 628-36	3.1	116
351	Composite pleomorphic xanthoastrocytoma and ganglioglioma: report of four cases and review of the literature. <i>American Journal of Surgical Pathology</i> , 1997 , 21, 763-71	6.7	114
350	Small cell astrocytoma: an aggressive variant that is clinicopathologically and genetically distinct from anaplastic oligodendroglioma. <i>Cancer</i> , 2004 , 101, 2318-26	6.4	113
349	Clinicopathologic aspects of 1p/19q loss and the diagnosis of oligodendroglioma. <i>Archives of Pathology and Laboratory Medicine</i> , 2007 , 131, 242-51	5	113

348	A role for chromosome 9p21 deletions in the malignant progression of meningiomas and the prognosis of anaplastic meningiomas. <i>Brain Pathology</i> , 2002 , 12, 183-90	6	112
347	CNS-PNETs with C19MC amplification and/or LIN28 expression comprise a distinct histogenetic diagnostic and therapeutic entity. <i>Acta Neuropathologica</i> , 2014 , 128, 291-303	14.3	111
346	Methylation-based classification of benign and malignant peripheral nerve sheath tumors. <i>Acta Neuropathologica</i> , 2016 , 131, 877-87	14.3	110
345	NF1 deletions in S-100 protein-positive and negative cells of sporadic and neurofibromatosis 1 (NF1)-associated plexiform neurofibromas and malignant peripheral nerve sheath tumors. <i>American Journal of Pathology</i> , 2001 , 159, 57-61	5.8	110
344	Ancillary FISH analysis for 1p and 19q status: preliminary observations in 287 gliomas and oligodendroglioma mimics. <i>Frontiers in Bioscience - Landmark</i> , 2003 , 8, a1-9	2.8	109
343	Disseminated oligodendroglial-like leptomeningeal tumor of childhood: a distinctive clinicopathologic entity. <i>Acta Neuropathologica</i> , 2012 , 124, 627-41	14.3	107
342	Germ-line and somatic DICER1 mutations in pineoblastoma. <i>Acta Neuropathologica</i> , 2014 , 128, 583-95	14.3	103
341	Morphologic and immunohistochemical features of malignant peripheral nerve sheath tumors and cellular schwannomas. <i>Modern Pathology</i> , 2015 , 28, 187-200	9.8	97
340	A sensitive and specific histopathologic prognostic marker for H3F3A K27M mutant pediatric glioblastomas. <i>Acta Neuropathologica</i> , 2014 , 128, 743-53	14.3	96
339	Molecular characterization of human meningiomas by gene expression profiling using high-density oligonucleotide microarrays. <i>American Journal of Pathology</i> , 2002 , 161, 665-72	5.8	96
338	Immunohistochemical analysis supports a role for INI1/SMARCB1 in hereditary forms of schwannomas, but not in solitary, sporadic schwannomas. <i>Brain Pathology</i> , 2008 , 18, 517-9	6	95
337	Differential NF1, p16, and EGFR patterns by interphase cytogenetics (FISH) in malignant peripheral nerve sheath tumor (MPNST) and morphologically similar spindle cell neoplasms. <i>Journal of Neuropathology and Experimental Neurology</i> , 2002 , 61, 702-9	3.1	95
336	Targeted inhibition of cyclic AMP phosphodiesterase-4 promotes brain tumor regression. <i>Clinical Cancer Research</i> , 2008 , 14, 7717-25	12.9	94
335	Spatiotemporal differences in CXCL12 expression and cyclic AMP underlie the unique pattern of optic glioma growth in neurofibromatosis type 1. <i>Cancer Research</i> , 2007 , 67, 8588-95	10.1	92
334	CDKN2A loss is associated with shortened overall survival in lower-grade (World Health Organization Grades II-III) astrocytomas. <i>Journal of Neuropathology and Experimental Neurology</i> , 2015 , 74, 442-52	3.1	91
333	Chromosome 22q deletions in atypical teratoid/rhabdoid tumors in adults. <i>Brain Pathology</i> , 2005 , 15, 23-8	6	90
332	A role for fluorescence in situ hybridization detection of chromosome 22q dosage in distinguishing atypical teratoid/rhabdoid tumors from medulloblastoma/central primitive neuroectodermal tumors. <i>Human Pathology</i> , 2001 , 32, 156-62	3.7	90
331	Poorly differentiated chordoma with SMARCB1/INI1 loss: a distinct molecular entity with dismal prognosis. <i>Acta Neuropathologica</i> , 2016 , 132, 149-51	14.3	89

330	Array-based comparative genomic hybridization identifies CDK4 and FOXM1 alterations as independent predictors of survival in malignant peripheral nerve sheath tumor. <i>Clinical Cancer Research</i> , 2011 , 17, 1924-34	12.9	88
329	Widespread CXCR4 activation in astrocytomas revealed by phospho-CXCR4-specific antibodies. <i>Cancer Research</i> , 2005 , 65, 11392-9	10.1	88
328	LIN28A immunoreactivity is a potent diagnostic marker of embryonal tumor with multilayered rosettes (ETMR). <i>Acta Neuropathologica</i> , 2012 , 124, 875-81	14.3	87
327	Clinical utility of fluorescence in situ hybridization (FISH) in morphologically ambiguous gliomas with hybrid oligodendroglial/astrocytic features. <i>Journal of Neuropathology and Experimental Neurology</i> , 2003 , 62, 1118-28	3.1	86
326	Metastatic adenocarcinoma to the brain: an immunohistochemical approach. <i>Human Pathology</i> , 1997 , 28, 938-43	3.7	85
325	Meningiomas in pregnancy: a clinicopathologic study of 17 cases. <i>Neurosurgery</i> , 2012 , 71, 951-61	3.2	83
324	High-resolution, dual-platform aCGH analysis reveals frequent HIPK2 amplification and increased expression in pilocytic astrocytomas. <i>Oncogene</i> , 2008 , 27, 4745-51	9.2	82
323	The alternative lengthening of telomere phenotype is significantly associated with loss of ATRX expression in high-grade pediatric and adult astrocytomas: a multi-institutional study of 214 astrocytomas. <i>Modern Pathology</i> , 2013 , 26, 1425-32	9.8	81
322	DNA sequence of the translocation breakpoints in undifferentiated embryonal sarcoma arising in mesenchymal hamartoma of the liver harboring the t(11;19)(q11;q13.4) translocation. <i>Genes Chromosomes and Cancer</i> , 2007 , 46, 508-13	5	81
321	Pediatric oligodendrogliomas: a study of molecular alterations on 1p and 19q using fluorescence in situ hybridization. <i>Journal of Neuropathology and Experimental Neurology</i> , 2003 , 62, 530-7	3.1	81
320	Spindle cell oncocytoma of the adenohypophysis: report of two recurrent cases. <i>American Journal of Surgical Pathology</i> , 2005 , 29, 247-53	6.7	81
319	Epithelioid Glioblastomas and Anaplastic Epithelioid Pleomorphic Xanthoastrocytomas--Same Entity or First Cousins?. <i>Brain Pathology</i> , 2016 , 26, 215-23	6	80
318	Glioma formation in neurofibromatosis 1 reflects preferential activation of K-RAS in astrocytes. <i>Cancer Research</i> , 2005 , 65, 236-45	10.1	80
317	DNA methylation profiling to predict recurrence risk in meningioma: development and validation of a nomogram to optimize clinical management. <i>Neuro-Oncology</i> , 2019 , 21, 901-910	1	79
316	A New Senior Editor. <i>Brain Pathology</i> , 2012 , 22, i-i	6	78
315	Announcing free color: another great reason to publish in Brain Pathology. <i>Brain Pathology</i> , 2013 , 23, 1-1	6	78
314	Clonal expansion and epigenetic reprogramming following deletion or amplification of mutant. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017 , 114, 10743-10748	11.5	78
313	The Juggler with Half the Balls on the Ground. <i>Brain Pathology</i> , 2010 , 20, i-i	6	78

312	Neuropathology, the Next Generation. <i>Brain Pathology</i> , 2010 , 20, iii-iii	6	78
311	New Blood In the Editorial Board. <i>Brain Pathology</i> , 2009 , 19, i-ii	6	78
310	Job Hunting at the Neuropathology Blog?. <i>Brain Pathology</i> , 2009 , 20, i-i	6	78
309	Brain Pathology Impact Factor on the Rise. <i>Brain Pathology</i> , 2007 , 17, v-vi	6	78
308	EPEN-07. PATTERNS OF EXTRANEURAL METASTASES IN PEDIATRIC SUPRATENTORIAL EPENDYMOMA: CASE SERIES AND REVIEW OF THE LITERATURE. <i>Neuro-Oncology</i> , 2020 , 22, iii309-iii309	1	78
307	SURG-02. A NOVEL RISK MODEL TO DEFINE THE RELATIVE BENEFIT OF MAXIMAL EXTENT OF RESECTION WITHIN PROGNOSTIC GROUPS IN NEWLY DIAGNOSED GLIOBLASTOMA. <i>Neuro-Oncology</i> , 2018 , 20, vi250-vi250	1	78
306	PDTM-38. PEDIATRIC MENINGIOMAS ARE CHARACTERIZED BY DISTINCT METHYLATION PROFILES DIFFERENT FROM ADULT MENINGIOMAS. <i>Neuro-Oncology</i> , 2018 , 20, vi212-vi212	1	78
305	PATH-05. IMPLEMENTATION OF A TARGETED NEXT-GENERATION SEQUENCING PANEL FOR THE DIAGNOSIS AND PRECISION MEDICINE TREATMENT OF ADULT PATIENTS WITH WHO GRADE IV DIFFUSE GLIOMAS. <i>Neuro-Oncology</i> , 2018 , 20, vi158-vi159	1	78
304	MNGI-30. RADIOLOGIC FEATURES ARE PROGNOSTIC FOR CLINICAL OUTCOMES OF CHORDOID MENINGIOMA. <i>Neuro-Oncology</i> , 2018 , 20, vi155-vi155	1	78
303	Large cell/anaplastic medulloblastomas and medulloblastomas: clinicopathological and genetic features. <i>Journal of Neurosurgery</i> , 2001 , 95, 82-8	3.2	77
302	Gone FISHing: clinical lessons learned in brain tumor molecular diagnostics over the last decade. <i>Brain Pathology</i> , 2011 , 21, 57-73	6	76
301	Anaplastic meningioma versus meningeal hemangiopericytoma: immunohistochemical and genetic markers. <i>Human Pathology</i> , 2004 , 35, 1413-8	3.7	76
300	The genetic landscape of ganglioglioma. <i>Acta Neuropathologica Communications</i> , 2018 , 6, 47	7.3	75
299	High rate of concurrent BRAF-KIAA1549 gene fusion and 1p deletion in disseminated oligodendroglioma-like leptomenigeal neoplasms (DOLN). <i>Acta Neuropathologica</i> , 2015 , 129, 609-610	14.3	73
298	Frequent promoter hypermethylation and transcriptional downregulation of the NDRG2 gene at 14q11.2 in primary glioblastoma. <i>International Journal of Cancer</i> , 2008 , 123, 2080-6	7.5	73
297	Interobserver and intraobserver reproducibility in focal cortical dysplasia (malformations of cortical development). <i>Epilepsia</i> , 2009 , 50, 2593-8	6.4	72
296	PDGFRA amplification is common in pediatric and adult high-grade astrocytomas and identifies a poor prognostic group in IDH1 mutant glioblastoma. <i>Brain Pathology</i> , 2013 , 23, 565-73	6	71
295	Clinicopathologic features of recurrent dysembryoplastic neuroepithelial tumor and rare malignant transformation: a report of 5 cases and review of the literature. <i>Journal of Neuro-Oncology</i> , 2009 , 94, 283-92	4.8	70

294	Primary intracerebral angiomatoid fibrous histiocytoma: report of a case with a t(12;22)(q13;q12) causing type 1 fusion of the EWS and ATF-1 genes. <i>American Journal of Surgical Pathology</i> , 2008 , 32, 478-84	6.7	70
293	Mapping of the chromosome 19 q-arm glioma tumor suppressor gene using fluorescence in situ hybridization and novel microsatellite markers. <i>Genes Chromosomes and Cancer</i> , 2000 , 29, 16-25	5	70
292	Somatostatin receptor 2a is a more sensitive diagnostic marker of meningioma than epithelial membrane antigen. <i>Acta Neuropathologica</i> , 2015 , 130, 441-3	14.3	69
291	Identification of a third Protein 4.1 tumor suppressor, Protein 4.1R, in meningioma pathogenesis. <i>Neurobiology of Disease</i> , 2003 , 13, 191-202	7.5	69
290	Meningioma metastatic to the lung. <i>Mayo Clinic Proceedings</i> , 1999 , 74, 1129-33	6.4	68
289	EWS-ATF1 fusion transcripts in gastrointestinal tumors previously diagnosed as malignant melanoma. <i>Human Pathology</i> , 2005 , 36, 74-81	3.7	67
288	The in vivo antitumoral effects of lipopolysaccharide against glioblastoma multiforme are mediated in part by Toll-like receptor 4. <i>Neurosurgery</i> , 2007 , 60, 372-80; discussion 381	3.2	66
287	Epithelioid glioblastomas stratify into established diagnostic subsets upon integrated molecular analysis. <i>Brain Pathology</i> , 2018 , 28, 656-662	6	65
286	Meningeal tumors of childhood and infancy. An update and literature review. <i>Brain Pathology</i> , 2003 , 13, 386-408	6	65
285	Postoperative radiation therapy for grade II and III intracranial ependymoma. <i>International Journal of Radiation Oncology Biology Physics</i> , 2005 , 61, 387-91	4	65
284	Cooperative interactions of BRAFV600E kinase and CDKN2A locus deficiency in pediatric malignant astrocytoma as a basis for rational therapy. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012 , 109, 8710-5	11.5	64
283	Gene expression profiling reveals unique molecular subtypes of Neurofibromatosis Type I-associated and sporadic malignant peripheral nerve sheath tumors. <i>Brain Pathology</i> , 2004 , 14, 297-303	6	63
282	Atypical teratoid/rhabdoid tumor arising in the setting of a pleomorphic xanthoastrocytoma. <i>Journal of Neuro-Oncology</i> , 2007 , 84, 217-22	4.8	62
281	Expression and prognostic impact of immune modulatory molecule PD-L1 in meningioma. <i>Journal of Neuro-Oncology</i> , 2016 , 130, 543-552	4.8	61
280	Pathology concordance levels for meningioma classification and grading in NRG Oncology RTOG Trial 0539. <i>Neuro-Oncology</i> , 2016 , 18, 565-74	1	61
279	Germline and somatic BAP1 mutations in high-grade rhabdoid meningiomas. <i>Neuro-Oncology</i> , 2017 , 19, 535-545	1	60
278	Ependymomas with neuronal differentiation: a morphologic and immunohistochemical spectrum. <i>Acta Neuropathologica</i> , 2007 , 113, 313-24	14.3	60
277	Expression of epidermal growth factor receptor in squamous cell carcinomas of the anal canal is independent of gene amplification. <i>Modern Pathology</i> , 2006 , 19, 942-9	9.8	60

276	Aggressive enterogenous cyst with extensive craniospinal spread: case report. <i>Neurosurgery</i> , 1999 , 44, 401-4; discussion 404-5	3.2	60
275	Comparative gene expression profile analysis of neurofibromatosis 1-associated and sporadic pilocytic astrocytomas. <i>Cancer Research</i> , 2002 , 62, 2085-91	10.1	60
274	Chemotherapy for adult low-grade gliomas: clinical outcomes by molecular subtype in a phase II study of adjuvant temozolomide. <i>Neuro-Oncology</i> , 2017 , 19, 242-251	1	59
273	Spectrum of pilomyxoid astrocytomas: intermediate pilomyxoid tumors. <i>American Journal of Surgical Pathology</i> , 2010 , 34, 1783-91	6.7	58
272	Natural history of neurofibromatosis 1-associated optic nerve glioma in mice. <i>Annals of Neurology</i> , 2005 , 57, 119-27	9.4	58
271	Characterization of gliomas: from morphology to molecules. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2017 , 471, 257-269	5.1	57
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