David Capper

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8112466/publications.pdf

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268 papers 30,590 citations

83 h-index 164

279 all docs

279 docs citations

times ranked

279

27814 citing authors

g-index

#	Article	IF	CITATIONS
1	DNA methylation-based classification of central nervous system tumours. Nature, 2018, 555, 469-474.	27.8	1,872
2	A complex secretory program orchestrated by the inflammasome controls paracrine senescence. Nature Cell Biology, 2013, 15, 978-990.	10.3	1,566
3	Type and frequency of IDH1 and IDH2 mutations are related to astrocytic and oligodendroglial differentiation and age: a study of 1,010 diffuse gliomas. Acta Neuropathologica, 2009, 118, 469-474.	7.7	1,020
4	Molecular Classification of Ependymal Tumors across All CNS Compartments, Histopathological Grades, and Age Groups. Cancer Cell, 2015, 27, 728-743.	16.8	933
5	Analysis of BRAF V600E mutation in 1,320 nervous system tumors reveals high mutation frequencies in pleomorphic xanthoastrocytoma, ganglioglioma and extra-cerebellar pilocytic astrocytoma. Acta Neuropathologica, 2011, 121, 397-405.	7.7	914
6	The whole-genome landscape of medulloblastoma subtypes. Nature, 2017, 547, 311-317.	27.8	787
7	Patients with IDH1 wild type anaplastic astrocytomas exhibit worse prognosis than IDH1-mutated glioblastomas, and IDH1 mutation status accounts for the unfavorable prognostic effect of higher age: implications for classification of gliomas. Acta Neuropathologica, 2010, 120, 707-718.	7.7	719
8	New Brain Tumor Entities Emerge from Molecular Classification of CNS-PNETs. Cell, 2016, 164, 1060-1072.	28.9	702
9	Actively personalized vaccination trial for newly diagnosed glioblastoma. Nature, 2019, 565, 240-245.	27.8	637
10	<scp>WHO</scp> 2016 Classification of gliomas. Neuropathology and Applied Neurobiology, 2018, 44, 139-150.	3.2	612
11	DNA methylation-based classification and grading system for meningioma: a multicentre, retrospective analysis. Lancet Oncology, The, 2017, 18, 682-694.	10.7	586
12	Characterization of R132H Mutationâ€specific IDH1 Antibody Binding in Brain Tumors. Brain Pathology, 2010, 20, 245-254.	4.1	463
13	Assessment of BRAF V600E mutation status by immunohistochemistry with a mutation-specific monoclonal antibody. Acta Neuropathologica, 2011, 122, 11-19.	7.7	445
14	Atypical Teratoid/Rhabdoid Tumors Are Comprised of Three Epigenetic Subgroups with Distinct Enhancer Landscapes. Cancer Cell, 2016, 29, 379-393.	16.8	438
15	Temozolomide chemotherapy versus radiotherapy in high-risk low-grade glioma (EORTC 22033-26033): a randomised, open-label, phase 3 intergroup study. Lancet Oncology, The, 2016, 17, 1521-1532.	10.7	396
16	Monoclonal antibody specific for IDH1 R132H mutation. Acta Neuropathologica, 2009, 118, 599-601.	7.7	380
17	ATRX and IDH1-R132H immunohistochemistry with subsequent copy number analysis and IDH sequencing as a basis for an "integrated―diagnostic approach for adult astrocytoma, oligodendroglioma and glioblastoma. Acta Neuropathologica, 2015, 129, 133-146.	7.7	378
18	cIMPACTâ€NOW update 6: new entity and diagnostic principle recommendations of the cIMPACTâ€Utrecht meeting on future CNS tumor classification and grading. Brain Pathology, 2020, 30, 844-856.	4.1	363

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19	Meningeal hemangiopericytoma and solitary fibrous tumors carry the NAB2-STAT6 fusion and can be diagnosed by nuclear expression of STAT6 protein. Acta Neuropathologica, 2013, 125, 651-658.	7.7	324
20	Practical implementation of DNA methylation and copy-number-based CNS tumor diagnostics: the Heidelberg experience. Acta Neuropathologica, 2018, 136, 181-210.	7.7	308
21	ATRX loss refines the classification of anaplastic gliomas and identifies a subgroup of IDH mutant astrocytic tumors with better prognosis. Acta Neuropathologica, 2013, 126, 443-451.	7.7	304
22	Novel, improved grading system(s) for IDH-mutant astrocytic gliomas. Acta Neuropathologica, 2018, 136, 153-166.	7.7	298
23	Immunohistochemistry Is Highly Sensitive and Specific for the Detection of V600E BRAF Mutation in Melanoma. American Journal of Surgical Pathology, 2013, 37, 61-65.	3.7	289
24	TERT Promoter Mutations and Risk of Recurrence in Meningioma. Journal of the National Cancer Institute, 2016, 108, djv377.	6.3	283
25	clMPACT-NOW update 2: diagnostic clarifications for diffuse midline glioma, H3 K27M-mutant and diffuse astrocytoma/anaplastic astrocytoma, IDH-mutant. Acta Neuropathologica, 2018, 135, 639-642.	7.7	281
26	Integrated analysis of pediatric glioblastoma reveals a subset of biologically favorable tumors with associated molecular prognostic markers. Acta Neuropathologica, 2015, 129, 669-678.	7.7	277
27	IDH mutant diffuse and anaplastic astrocytomas have similar age at presentation and little difference in survival: a grading problem for WHO. Acta Neuropathologica, 2015, 129, 867-873.	7.7	272
28	Farewell to oligoastrocytoma: in situ molecular genetics favor classification as either oligodendroglioma or astrocytoma. Acta Neuropathologica, 2014, 128, 551-559.	7.7	268
29	Next-generation personalised medicine for high-risk paediatric cancer patients – The INFORM pilot study. European Journal of Cancer, 2016, 65, 91-101.	2.8	262
30	Combined molecular analysis of BRAF and IDH1 distinguishes pilocytic astrocytoma from diffuse astrocytoma. Acta Neuropathologica, 2009, 118, 401-405.	7.7	255
31	Distribution of TERT promoter mutations in pediatric and adult tumors of the nervous system. Acta Neuropathologica, 2013, 126, 907-915.	7.7	254
32	Adult IDH wild type astrocytomas biologically and clinically resolve into other tumor entities. Acta Neuropathologica, 2015, 130, 407-417.	7.7	237
33	Sarcoma classification by DNA methylation profiling. Nature Communications, 2021, 12, 498.	12.8	237
34	Radiogenomics of Glioblastoma: Machine Learning–based Classification of Molecular Characteristics by Using Multiparametric and Multiregional MR Imaging Features. Radiology, 2016, 281, 907-918.	7.3	236
35	Secretory meningiomas are defined by combined KLF4 K409Q and TRAF7 mutations. Acta Neuropathologica, 2013, 125, 351-358.	7.7	208
36	Immunohistochemical testing of BRAF V600E status in 1,120 tumor tissue samples of patients with brain metastases. Acta Neuropathologica, 2012, 123, 223-233.	7.7	204

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37	Next-generation sequencing in routine brain tumor diagnostics enables an integrated diagnosis and identifies actionable targets. Acta Neuropathologica, 2016, 131, 903-910.	7.7	203
38	Targeting the BRAF V600E Mutation in Multiple Myeloma. Cancer Discovery, 2013, 3, 862-869.	9.4	202
39	BRAFV600E mutant protein is expressed in cells of variable maturation in Langerhans cell histiocytosis. Blood, 2012, 120, e28-e34.	1.4	199
40	Oncolytic H-1 Parvovirus Shows Safety and Signs of Immunogenic Activity in a First Phase I/IIa Glioblastoma Trial. Molecular Therapy, 2017, 25, 2620-2634.	8.2	199
41	A Phase II randomized study of galunisertib monotherapy or galunisertib plus lomustine compared with lomustine monotherapy in patients with recurrent glioblastoma. Neuro-Oncology, 2016, 18, 1146-1156.	1.2	197
42	Distribution of EGFR amplification, combined chromosome 7 gain and chromosome 10 loss, and TERT promoter mutation in brain tumors and their potential for the reclassification of IDHwt astrocytoma to glioblastoma. Acta Neuropathologica, 2018, 136, 793-803.	7.7	195
43	Anaplastic astrocytoma with piloid features, a novel molecular class of IDH wildtype glioma with recurrent MAPK pathway, CDKN2A/B and ATRX alterations. Acta Neuropathologica, 2018, 136, 273-291.	7.7	190
44	Recurrent MET fusion genes represent a drug target in pediatric glioblastoma. Nature Medicine, 2016, 22, 1314-1320.	30.7	183
45	Immunohistochemical Detection of the BRAF V600E-mutated Protein in Papillary Thyroid Carcinoma. American Journal of Surgical Pathology, 2012, 36, 844-850.	3.7	177
46	Mutant BRAF V600E protein in ganglioglioma is predominantly expressed by neuronal tumor cells. Acta Neuropathologica, 2013, 125, 891-900.	7.7	177
47	p53-Dependent Nestin Regulation Links Tumor Suppression to Cellular Plasticity in Liver Cancer. Cell, 2014, 158, 579-592.	28.9	176
48	Integrated DNA methylation and copy-number profiling identify three clinically and biologically relevant groups of anaplastic glioma. Acta Neuropathologica, 2014, 128, 561-571.	7.7	176
49	Polymorphous low-grade neuroepithelial tumor of the young (PLNTY): an epileptogenic neoplasm with oligodendroglioma-like components, aberrant CD34 expression, and genetic alterations involving the MAP kinase pathway. Acta Neuropathologica, 2017, 133, 417-429.	7.7	172
50	Radiomic subtyping improves disease stratification beyond key molecular, clinical, and standard imaging characteristics in patients with glioblastoma. Neuro-Oncology, 2018, 20, 848-857.	1.2	170
51	clMPACTâ€NOW update 7: advancing the molecular classification of ependymal tumors. Brain Pathology, 2020, 30, 863-866.	4.1	168
52	Brain metastases: pathobiology and emerging targeted therapies. Acta Neuropathologica, 2012, 123, 205-222.	7.7	163
53	Histologically distinct neuroepithelial tumors with histone 3 G34 mutation are molecularly similar and comprise a single nosologic entity. Acta Neuropathologica, 2016, 131, 137-146.	7.7	162
54	Infant High-Grade Gliomas Comprise Multiple Subgroups Characterized by Novel Targetable Gene Fusions and Favorable Outcomes. Cancer Discovery, 2020, 10, 942-963.	9.4	157

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55	Methylation-based classification of benign and malignant peripheral nerve sheath tumors. Acta Neuropathologica, 2016, 131, 877-887.	7.7	151
56	Addressing Diffuse Glioma as a Systemic Brain Disease With Single-Cell Analysis. Archives of Neurology, 2012, 69, 523.	4.5	148
57	Pan-mutant IDH1 inhibitor BAY 1436032 for effective treatment of IDH1 mutant astrocytoma in vivo. Acta Neuropathologica, 2017, 133, 629-644.	7.7	146
58	clMPACT-NOW update 1: Not Otherwise Specified (NOS) and Not Elsewhere Classified (NEC). Acta Neuropathologica, 2018, 135, 481-484.	7.7	145
59	Detection of BRAF p.V600E Mutations in Melanomas. Journal of Molecular Diagnostics, 2013, 15, 94-100.	2.8	144
60	H3-/IDH-wild type pediatric glioblastoma is comprised of molecularly and prognostically distinct subtypes with associated oncogenic drivers. Acta Neuropathologica, 2017, 134, 507-516.	7.7	144
61	Molecular, Pathological, Radiological, and Immune Profiling of Non-brainstem Pediatric High-Grade Glioma from the HERBY Phase II Randomized Trial. Cancer Cell, 2018, 33, 829-842.e5.	16.8	140
62	Adamantinomatous and papillary craniopharyngiomas are characterized by distinct epigenomic as well as mutational and transcriptomic profiles. Acta Neuropathologica Communications, 2016, 4, 20.	5.2	136
63	Application of a BRAF V600E Mutation-specific Antibody for the Diagnosis of Hairy Cell Leukemia. American Journal of Surgical Pathology, 2012, 36, 1796-1800.	3.7	135
64	<scp>BRAF V600E /scp> Mutation Is Associated with <scp>mTOR /scp> Signaling Activation in Glioneuronal Tumors. Brain Pathology, 2014, 24, 52-66.</scp></scp>	4.1	129
65	Poorly differentiated chordoma with SMARCB1/INI1 loss: a distinct molecular entity with dismal prognosis. Acta Neuropathologica, 2016, 132, 149-151.	7.7	127
66	BRAFV600E Immunohistochemistry Facilitates Universal Screening of Colorectal Cancers for Lynch Syndrome. American Journal of Surgical Pathology, 2013, 37, 1592-1602.	3.7	125
67	Mutation-specific IDH1 antibody differentiates oligodendrogliomas and oligoastrocytomas from other brain tumors with oligodendroglioma-like morphology. Acta Neuropathologica, 2011, 121, 241-252.	7.7	124
68	Expression pattern of the water channel aquaporin-4 in human gliomas is associated with blood–brain barrier disturbance but not with patient survival. Journal of Neuroscience Research, 2007, 85, 1336-1346.	2.9	120
69	Announcing clMPACT-NOW: the Consortium to Inform Molecular and Practical Approaches to CNS Tumor Taxonomy. Acta Neuropathologica, 2017, 133, 1-3.	7.7	120
70	Molecularly defined diffuse leptomeningeal glioneuronal tumor (DLGNT) comprises two subgroups with distinct clinical and genetic features. Acta Neuropathologica, 2018, 136, 239-253.	7.7	118
71	Low-grade epilepsy-associated neuroepithelial tumours — the 2016 WHO classification. Nature Reviews Neurology, 2016, 12, 732-740.	10.1	113
72	Distinct requirement for an intact dimer interface in wild-type, V600E and kinase-dead B-Raf signalling. EMBO Journal, 2012, 31, 2629-2647.	7.8	110

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73	The Pediatric Precision Oncology INFORM Registry: Clinical Outcome and Benefit for Patients with Very High-Evidence Targets. Cancer Discovery, 2021, 11, 2764-2779.	9.4	110
74	Application of Mutant IDH1 Antibody to Differentiate Diffuse Glioma From Nonneoplastic Central Nervous System Lesions and Therapy-induced Changes. American Journal of Surgical Pathology, 2010, 34, 1199-1204.	3.7	108
75	Diagnostic value of immunohistochemistry for the detection of the BRAF mutation in primary lung adenocarcinoma Caucasian patients. Annals of Oncology, 2013, 24, 742-748.	1.2	103
76	Nuclear relocation of <scp>STAT</scp> 6 reliably predicts <i>NAB2</i> – <i>STAT6</i> fusion for the diagnosis of solitary fibrous tumour. Histopathology, 2014, 65, 613-622.	2.9	101
77	Methylation array profiling of adult brain tumours: diagnostic outcomes in a large, single centre. Acta Neuropathologica Communications, 2019, 7, 24.	5.2	101
78	Machine learning analysis of DNA methylation profiles distinguishes primary lung squamous cell carcinomas from head and neck metastases. Science Translational Medicine, 2019, 11, .	12.4	100
79	BRAF V600E-specific immunohistochemistry reveals low mutation rates in biliary tract cancer and restriction to intrahepatic cholangiocarcinoma. Modern Pathology, 2014, 27, 1028-1034.	5.5	96
80	BRAF V600Eâ€specific immunohistochemistry for the exclusion of Lynch syndrome in MSIâ€H colorectal cancer. International Journal of Cancer, 2013, 133, 1624-1630.	5.1	93
81	Epithelioid glioblastomas stratify into established diagnostic subsets upon integrated molecular analysis. Brain Pathology, 2018, 28, 656-662.	4.1	89
82	Machine learning workflows to estimate class probabilities for precision cancer diagnostics on DNA methylation microarray data. Nature Protocols, 2020, 15, 479-512.	12.0	89
83	Stem-cell-like glioma cells are resistant to TRAIL/Apo2L and exhibit down-regulation of caspase-8 by promoter methylation. Acta Neuropathologica, 2009, 117, 445-456.	7.7	88
84	AKT1E17K mutations cluster with meningothelial and transitional meningiomas and can be detected by SFRP1 immunohistochemistry. Acta Neuropathologica, 2013, 126, 757-762.	7.7	88
85	Isomorphic diffuse glioma is a morphologically and molecularly distinct tumour entity with recurrent gene fusions of MYBL1 or MYB and a benign disease course. Acta Neuropathologica, 2020, 139, 193-209.	7.7	83
86	Malignant astrocytomas of elderly patients lack favorable molecular markers: an analysis of the NOA-08 study collective. Neuro-Oncology, 2013, 15, 1017-1026.	1.2	78
87	Detection of the BRAF V600E mutation in serous ovarian tumors: a comparative analysis of immunohistochemistry with a mutation-specific monoclonal antibody and allele-specific PCR. Human Pathology, 2013, 44, 329-335.	2.0	77
88	Assessing CpG island methylator phenotype, $1p/19q$ codeletion, and MGMT promoter methylation from epigenome-wide data in the biomarker cohort of the NOA-04 trial. Neuro-Oncology, 2014, 16, 1630-1638.	1.2	77
89	2â€Hydroxyglutarate concentration in serum from patients with gliomas does not correlate with IDH1/2 mutation status or tumor size. International Journal of Cancer, 2012, 131, 766-768.	5.1	74
90	Molecular profiling of long-term survivors identifies a subgroup of glioblastoma characterized by chromosome 19/20 co-gain. Acta Neuropathologica, 2015, 130, 419-434.	7.7	74

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91	Gliomatosis cerebri: no evidence for a separate brain tumor entity. Acta Neuropathologica, 2016, 131, 309-319.	7.7	74
92	Evidence of H3 K27M mutations in posterior fossa ependymomas. Acta Neuropathologica, 2016, 132, 635-637.	7.7	73
93	«scp> <i>BRAF</i> â€Mutated Pleomorphic Xanthoastrocytoma is Associated with Temporal Location, Reticulin Fiber Deposition and <scp>CD</scp> 34 Expression. Brain Pathology, 2014, 24, 221-229.	4.1	72
94	Phase 1b/2a study of galunisertib, a small molecule inhibitor of transforming growth factor-beta receptor I, in combination with standard temozolomide-based radiochemotherapy in patients with newly diagnosed malignant glioma. Investigational New Drugs, 2020, 38, 1570-1579.	2.6	70
95	Analysis of IDH mutation, $1p/19q$ deletion, and PTEN loss delineates prognosis in clinical low-grade diffuse gliomas. Neuro-Oncology, 2014, 16, 914-923.	1.2	69
96	Pitfalls in the assessment of MGMT expression and in its correlation with survival in diffuse astrocytomas: proposal of a feasible immunohistochemical approach. Acta Neuropathologica, 2008, 115, 249-259.	7.7	68
97	Tall cell papillary thyroid carcinoma: new diagnostic criteria and mutations in BRAF and TERT. Endocrine-Related Cancer, 2015, 22, 419-429.	3.1	68
98	Integrated molecular characterization of <i><scp>IDH</scp></i> â€mutant glioblastomas. Neuropathology and Applied Neurobiology, 2019, 45, 108-118.	3.2	68
99	Response to trametinib treatment in progressive pediatric low-grade glioma patients. Journal of Neuro-Oncology, 2020, 149, 499-510.	2.9	68
100	Continued Response Off Treatment After BRAF Inhibition in Refractory Hairy Cell Leukemia. Journal of Clinical Oncology, 2013, 31, e300-e303.	1.6	67
101	Methylation profiling of choroid plexus tumors reveals 3 clinically distinct subgroups. Neuro-Oncology, 2016, 18, 790-796.	1.2	67
102	No prognostic value of IDH1 mutations in a series of 100 WHO grade II astrocytomas. Journal of Neuro-Oncology, 2012, 109, 15-22.	2.9	66
103	Melanotic Tumors of the Nervous System are Characterized by Distinct Mutational, Chromosomal and Epigenomic Profiles. Brain Pathology, 2015, 25, 202-208.	4.1	66
104	Utilization of a MAB for BRAFV600E detection in papillary thyroid carcinoma. Endocrine-Related Cancer, 2012, 19, 779-784.	3.1	65
105	Bevacizumab plus hypofractionated radiotherapy versus radiotherapy alone in elderly patients with glioblastoma: the randomized, open-label, phase II ARTE trial. Annals of Oncology, 2018, 29, 1423-1430.	1.2	65
106	DNA methylation-based classification of ependymomas in adulthood: implications for diagnosis and treatment. Neuro-Oncology, 2018, 20, 1616-1624.	1.2	65
107	YAP1 subgroup supratentorial ependymoma requires TEAD and nuclear factor I-mediated transcriptional programmes for tumorigenesis. Nature Communications, 2019, 10, 3914.	12.8	65
108	Genetic Alterations in Gliosarcoma and Giant Cell Glioblastoma. Brain Pathology, 2016, 26, 517-522.	4.1	63

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109	cIMPACTâ€NOW (the consortium to inform molecular and practical approaches to CNS tumor) Tj ETQq1 1 0.78 27, 851-852.	34314 rgBT 4.1	Overlock 10
110	Isocitrate dehydrogenase 1 mutant R132H sensitizes glioma cells to BCNU-induced oxidative stress and cell death. Apoptosis: an International Journal on Programmed Cell Death, 2013, 18, 1416-1425.	4.9	62
111	Acinar cell carcinomas of the pancreas: a molecular analysis in a series of 57 cases. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2014, 465, 661-672.	2.8	61
112	DNA methylation-based reclassification of olfactory neuroblastoma. Acta Neuropathologica, 2018, 136, 255-271.	7.7	59
113	PCR―and Restriction Endonucleaseâ€Based Detection of <i>IDH1</i> Mutations. Brain Pathology, 2010, 20, 298-300.	4.1	58
114	Cribriform neuroepithelial tumor: molecular characterization of a SMARCB1â€deficient nonâ€rhabdoid tumor with favorable longâ€term outcome. Brain Pathology, 2017, 27, 411-418.	4.1	58
115	Extensive Molecular and Clinical Heterogeneity in Patients With Histologically Diagnosed CNS-PNET Treated as a Single Entity: A Report From the Children's Oncology Group Randomized ACNS0332 Trial. Journal of Clinical Oncology, 2018, 36, 3388-3395.	1.6	58
116	Detection of <i>BRAF </i> /i> p.V600E Mutations in Melanoma by Immunohistochemistry Has a Good Interobserver Reproducibility. Archives of Pathology and Laboratory Medicine, 2014, 138, 71-75.	2.5	57
117	A single-arm phase II Austrian/German multicenter trial on continuous daily sunitinib in primary glioblastoma at first recurrence (SURGE 01-07). Neuro-Oncology, 2014, 16, 92-102.	1,2	57
118	Diagnostic Value of Immunohistochemistry for the Detection of the <i>BRAF^{V600E}</i> Mutation in Papillary Thyroid Carcinoma: Comparative Analysis with Three DNA-Based Assays. Thyroid, 2014, 24, 858-866.	4.5	57
119	Rosette-forming glioneuronal tumors share a distinct DNA methylation profile and mutations in FGFR1, with recurrent co-mutation of PIK3CA and NF1. Acta Neuropathologica, 2019, 138, 497-504.	7.7	57
120	Elevated HLA-E levels in human glioblastomas but not in grade I to III astrocytomas correlate with infiltrating CD8+ cells. Journal of Neuroimmunology, 2007, 189, 50-58.	2.3	56
121	BRAFV600E protein expression and outcome from BRAF inhibitor treatment in BRAFV600E metastatic melanoma. British Journal of Cancer, 2013, 108, 924-931.	6.4	55
122	Diagnostics of pediatric supratentorial RELA ependymomas: integration of information from histopathology, genetics, DNA methylation and imaging. Brain Pathology, 2019, 29, 325-335.	4.1	55
123	The Senescence-associated Secretory Phenotype Mediates Oncogene-induced Senescence in Pediatric Pilocytic Astrocytoma. Clinical Cancer Research, 2019, 25, 1851-1866.	7.0	55
124	DNA methylation-based profiling for paediatric CNS tumour diagnosis and treatment: a population-based study. The Lancet Child and Adolescent Health, 2020, 4, 121-130.	5.6	55
125	IDH testing in diagnostic neuropathology: review and practical guideline article invited by the Euro-CNS research committee., 2011, 30, 217-230.		55
126	<i>GRHL1</i> Acts as Tumor Suppressor in Neuroblastoma and Is Negatively Regulated by MYCN and HDAC3. Cancer Research, 2014, 74, 2604-2616.	0.9	54

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127	Recurrent homozygous deletion of DROSHA and microduplication of PDE4DIP in pineoblastoma. Nature Communications, 2018, 9, 2868.	12.8	54
128	Modulation of TGFâ \in β activity by latent TGFâ \in βâ \in binding protein 1 in human malignant glioma cells. International Journal of Cancer, 2009, 125, 530-540.	5.1	53
129	The diagnostic utility of TP53 and CDKN2A to distinguish ovarian high-grade serous carcinoma from low-grade serous ovarian tumors. Modern Pathology, 2013, 26, 1255-1263.	5.5	52
130	Detection of BRAF V600E mutations in skin metastases of malignant melanoma by monoclonal antibody VE1. Journal of the American Academy of Dermatology, 2012, 67, 488-491.	1.2	51
131	BRAF Fusion Analysis in Pilocytic Astrocytomas: KIAA1549-BRAF 15-9 Fusions Are More Frequent in the Midline Than Within the Cerebellum. Journal of Neuropathology and Experimental Neurology, 2015, 74, 867-872.	1.7	51
132	Chemotherapy with BCNU in recurrent glioma: Analysis of clinical outcome and side effects in chemotherapy-na \tilde{A} -ve patients. BMC Cancer, 2016, 16, 81.	2.6	51
133	Histone 3.3 hotspot mutations in conventional osteosarcomas: a comprehensive clinical and molecular characterization of six H3F3A mutated cases. Clinical Sarcoma Research, 2017, 7, 9.	2.3	51
134	Development, Characterization, and Reversal of Acquired Resistance to the MEK1 Inhibitor Selumetinib (AZD6244) in an <i>In Vivo</i> Model of Childhood Astrocytoma. Clinical Cancer Research, 2013, 19, 6716-6729.	7.0	50
135	<scp><i>BRAF V</i></scp> <i>600E</i> analysis for the differentiation of papillary craniopharyngiomas and <scp>R</scp> athke's cleft cysts. Neuropathology and Applied Neurobiology, 2015, 41, 733-742.	3.2	50
136	Detection of 2â€Hydroxyglutarate in Formalinâ€Fixed Paraffinâ€Embedded Glioma Specimens by Gas Chromatography/Mass Spectrometry. Brain Pathology, 2012, 22, 26-31.	4.1	49
137	Improved molecular classification of serrated lesions of the colon by immunohistochemical detection of BRAF V600E. Modern Pathology, 2014, 27, 135-144.	5.5	49
138	Primary glioblastoma cultures: can profiling of stem cell markers predict radiotherapy sensitivity?. Journal of Neurochemistry, 2014, 131, 251-264.	3.9	47
139	Review: Challenges in the histopathological classification of ganglioglioma and DNT: microscopic agreement studies and a preliminary genotypeâ€phenotype analysis. Neuropathology and Applied Neurobiology, 2019, 45, 95-107.	3.2	46
140	Dabrafenib in patients with recurrent, BRAF V600E mutated malignant glioma and leptomeningeal disease. Oncology Reports, 2017, 38, 3291-3296.	2.6	46
141	A pan-tissue DNA methylation atlas enables in silico decomposition of human tissue methylomes at cell-type resolution. Nature Methods, 2022, 19, 296-306.	19.0	46
142	Infratentorial IDH-mutant astrocytoma is a distinct subtype. Acta Neuropathologica, 2020, 140, 569-581.	7.7	45
143	Sellar Region Atypical Teratoid/Rhabdoid Tumors (ATRT) in Adults Display DNA Methylation Profiles of the ATRT-MYC Subgroup. American Journal of Surgical Pathology, 2018, 42, 506-511.	3.7	43
144	Papillary glioneuronal tumor (PGNT) exhibits a characteristic methylation profile and fusions involving PRKCA. Acta Neuropathologica, 2019, 137, 837-846.	7.7	43

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145	Establishment and application of a novel patient-derived KIAA1549:BRAF-driven pediatric pilocytic astrocytoma model for preclinical drug testing. Oncotarget, 2017, 8, 11460-11479.	1.8	43
146	Survival According to BRAF-V600 Tumor Mutations – An Analysis of 437 Patients with Primary Melanoma. PLoS ONE, 2014, 9, e86194.	2.5	42
147	Neurofibromin specific antibody differentiates malignant peripheral nerve sheath tumors (MPNST) from other spindle cell neoplasms. Acta Neuropathologica, 2014, 127, 565-572.	7.7	41
148	Loss of FUBP1 expression in gliomas predicts FUBP1 mutation and is associated with oligodendroglial differentiation, IDH1 mutation and $1p/19q$ loss of heterozygosity. Neuropathology and Applied Neurobiology, 2014, 40, 205-216.	3.2	41
149	Artificial intelligence and pathology: From principles to practice and future applications in histomorphology and molecular profiling. Seminars in Cancer Biology, 2022, 84, 129-143.	9.6	41
150	Increased levels of 2â€hydroxyglutarate in AML patients with IDH1â€R132H and IDH2â€R140Q mutations. European Journal of Haematology, 2010, 85, 457-459.	2.2	39
151	Papillary Tumor of the Pineal Region: A Distinct Molecular Entity. Brain Pathology, 2016, 26, 199-205.	4.1	39
152	Desmoplastic Infantile Ganglioglioma/Astrocytoma (DIG/DIA) Are Distinct Entities with Frequent BRAFV600 Mutations. Molecular Cancer Research, 2018, 16, 1491-1498.	3.4	39
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