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List of Publications by Year in descending order

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Version: 2024-02-01

687363 794594 20 528 13 19 citations h-index g-index papers 21 21 21 877 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Highâ€grade neuroepithelial tumor with <i>BCOR</i> exon 15 internal tandem duplication—a comprehensive clinical, radiographic, pathologic, and genomic analysis. Brain Pathology, 2020, 30, 46-62.	4.1	69
2	Primary intracranial sarcomas with DICER1 mutation often contain prominent eosinophilic cytoplasmic globules and can occur in the setting of neurofibromatosis type 1. Acta Neuropathologica, 2019, 137, 521-525.	7.7	51
3	Pediatric bithalamic gliomas have a distinct epigenetic signature and frequent EGFR exon 20 insertions resulting in potential sensitivity to targeted kinase inhibition. Acta Neuropathologica, 2020, 139, 1071-1088.	7.7	50
4	Myxoid glioneuronal tumor, <i>PDGFRA</i> p.K385â€mutant: clinical, radiologic, and histopathologic features. Brain Pathology, 2020, 30, 479-494.	4.1	46
5	Recurrent KBTBD4 small in-frame insertions and absence of DROSHA deletion or DICER1 mutation differentiate pineal parenchymal tumor of intermediate differentiation (PPTID) from pineoblastoma. Acta Neuropathologica, 2019, 137, 851-854.	7.7	45
6	Intracranial mesenchymal tumor with FETâ€CREB fusionâ€"A unifying diagnosis for the spectrum of intracranial myxoid mesenchymal tumors and angiomatoid fibrous histiocytomaâ€like neoplasms. Brain Pathology, 2021, 31, e12918.	4.1	44
7	The role of histone modifications and telomere alterations in the pathogenesis of diffuse gliomas in adults and children. Journal of Neuro-Oncology, 2017, 132, 1-11.	2.9	35
8	Comprehensive analysis of diverse low-grade neuroepithelial tumors with FGFR1 alterations reveals a distinct molecular signature of rosette-forming glioneuronal tumor. Acta Neuropathologica Communications, 2020, 8, 151.	5.2	35
9	Clinicopathologic features of anaplastic myxopapillary ependymomas. Brain Pathology, 2019, 29, 75-84.	4.1	25
10	Recurrent non-canonical histone H3 mutations in spinal cord diffuse gliomas. Acta Neuropathologica, 2019, 138, 877-881.	7.7	21
11	Clinicopathologic and molecular features of intracranial desmoplastic small round cell tumors. Brain Pathology, 2020, 30, 213-225.	4.1	20
12	Gliomas arising in the setting of Li-Fraumeni syndrome stratify into two molecular subgroups with divergent clinicopathologic features. Acta Neuropathologica, 2020, 139, 953-957.	7.7	18
13	Utility of Pit-1 Immunostaining in Distinguishing Pituitary Adenomas of Primitive Differentiation from Null Cell Adenomas. Endocrine Pathology, 2017, 28, 287-292.	9.0	16
14	Low-grade glioneuronal tumors with FGFR2 fusion resolve into a single epigenetic group corresponding to †Polymorphous low-grade neuroepithelial tumor of the young'. Acta Neuropathologica, 2021, 142, 595-599.	7.7	16
15	Oligodendrogliomas, IDH-mutant and $1p/19q$ -codeleted, arising during teenage years often lack TERT promoter mutation that is typical of their adult counterparts. Acta Neuropathologica Communications, 2018, 6, 95.	5.2	13
16	Intracranial mesenchymal tumors with FETâ€CREB fusion are composed of at least two epigenetic subgroups distinct from meningioma and extracranial sarcomas. Brain Pathology, 2022, 32, e13037.	4.1	11
17	Prospective genomically guided identification of "early/evolving―and "undersampled―IDH-wildtype glioblastoma leads to improved clinical outcomes. Neuro-Oncology, 2022, 24, 1749-1762.	1.2	10
18	Tumor DNA requirements for accurate epigenetic-based classification of CNS neoplasia. Neuro-Oncology, 2021, 23, 1798-1800.	1.2	2

#	Article	IF	CITATIONS
19	Innumerable Meningiomas Arising in a Patient With Tuberous Sclerosis Complex Decades After Radiation Therapy. Pediatric and Developmental Pathology, 2021, 24, 471-477.	1.0	1
20	Molecular characterization of metachronous atypical teratoid rhabdoid tumors occurring in a young man 15Âyears apart. Pediatric Blood and Cancer, 2023, 70, .	1.5	0