

Julieann C Lee

List of Publications by Year in descending order

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Version: 2024-02-01

20
papers

528
citations

687363

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794594

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21
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877
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular characterization of metachronous atypical teratoid rhabdoid tumors occurring in a young man 15 years apart. <i>Pediatric Blood and Cancer</i> , 2023, 70, .	1.5	0
2	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
3	Prospective genomically guided identification of "early/evolving" and "undersampled" IDH-wildtype glioblastoma leads to improved clinical outcomes. <i>Neuro-Oncology</i> , 2022, 24, 1749-1762.	1.2	10
4	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
5	Innumerable Meningiomas Arising in a Patient With Tuberous Sclerosis Complex Decades After Radiation Therapy. <i>Pediatric and Developmental Pathology</i> , 2021, 24, 471-477.	1.0	1
6	Low-grade glioneuronal tumors with FGFR2 fusion resolve into a single epigenetic group corresponding to "Polymorphous low-grade neuroepithelial tumor of the young"™. <i>Acta Neuropathologica</i> , 2021, 142, 595-599.	7.7	16
7	Tumor DNA requirements for accurate epigenetic-based classification of CNS neoplasia. <i>Neuro-Oncology</i> , 2021, 23, 1798-1800.	1.2	2
8	High-grade neuroepithelial tumor with <i>BCOR</i> exon 15 internal tandem duplication "a comprehensive clinical, radiographic, pathologic, and genomic analysis. <i>Brain Pathology</i> , 2020, 30, 46-62.	4.1	69
9	Myxoid glioneuronal tumor, <i>PDGFRA</i> p.K385 mutant: clinical, radiologic, and histopathologic features. <i>Brain Pathology</i> , 2020, 30, 479-494.	4.1	46
10	Clinicopathologic and molecular features of intracranial desmoplastic small round cell tumors. <i>Brain Pathology</i> , 2020, 30, 213-225.	4.1	20
11	Comprehensive analysis of diverse low-grade neuroepithelial tumors with FGFR1 alterations reveals a distinct molecular signature of rosette-forming glioneuronal tumor. <i>Acta Neuropathologica Communications</i> , 2020, 8, 151.	5.2	35
12	Gliomas arising in the setting of Li-Fraumeni syndrome stratify into two molecular subgroups with divergent clinicopathologic features. <i>Acta Neuropathologica</i> , 2020, 139, 953-957.	7.7	18
13	Pediatric bithalamic gliomas have a distinct epigenetic signature and frequent EGFR exon 20 insertions resulting in potential sensitivity to targeted kinase inhibition. <i>Acta Neuropathologica</i> , 2020, 139, 1071-1088.	7.7	50
14	Recurrent non-canonical histone H3 mutations in spinal cord diffuse gliomas. <i>Acta Neuropathologica</i> , 2019, 138, 877-881.	7.7	21
15	Recurrent KBTBD4 small in-frame insertions and absence of DROSHA deletion or DICER1 mutation differentiate pineal parenchymal tumor of intermediate differentiation (PPTID) from pineoblastoma. <i>Acta Neuropathologica</i> , 2019, 137, 851-854.	7.7	45
16	Clinicopathologic features of anaplastic myxopapillary ependymomas. <i>Brain Pathology</i> , 2019, 29, 75-84.	4.1	25
17	Primary intracranial sarcomas with DICER1 mutation often contain prominent eosinophilic cytoplasmic globules and can occur in the setting of neurofibromatosis type 1. <i>Acta Neuropathologica</i> , 2019, 137, 521-525.	7.7	51
18	Oligodendrogliomas, IDH-mutant and 1p/19q-codeleted, arising during teenage years often lack TERT promoter mutation that is typical of their adult counterparts. <i>Acta Neuropathologica Communications</i> , 2018, 6, 95.	5.2	13

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19	The role of histone modifications and telomere alterations in the pathogenesis of diffuse gliomas in adults and children. <i>Journal of Neuro-Oncology</i> , 2017, 132, 1-11.	2.9	35
20	Utility of Pit-1 Immunostaining in Distinguishing Pituitary Adenomas of Primitive Differentiation from Null Cell Adenomas. <i>Endocrine Pathology</i> , 2017, 28, 287-292.	9.0	16