

Leonille Schweizer

List of Publications by Year in descending order

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36
papers

4,840
citations

361413
20
h-index

361022
35
g-index

37
all docs

37
docs citations

37
times ranked

7109
citing authors

#	ARTICLE	IF	CITATIONS
1	DNA methylation-based classification of central nervous system tumours. <i>Nature</i> , 2018, 555, 469-474.	27.8	1,872
2	Molecular Classification of Ependymal Tumors across All CNS Compartments, Histopathological Grades, and Age Groups. <i>Cancer Cell</i> , 2015, 27, 728-743.	16.8	933
3	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. <i>Acta Neuropathologica</i> , 2015, 129, 133-146.	7.7	378
4	Meningeal hemangiopericytoma and solitary fibrous tumors carry the NAB2-STAT6 fusion and can be diagnosed by nuclear expression of STAT6 protein. <i>Acta Neuropathologica</i> , 2013, 125, 651-658.	7.7	324
5	Secretory meningiomas are defined by combined KLF4 K409Q and TRAF7 mutations. <i>Acta Neuropathologica</i> , 2013, 125, 351-358.	7.7	208
6	Integrated DNA methylation and copy-number profiling identify three clinically and biologically relevant groups of anaplastic glioma. <i>Acta Neuropathologica</i> , 2014, 128, 561-571.	7.7	176
7	Adamantinomatous and papillary craniopharyngiomas are characterized by distinct epigenomic as well as mutational and transcriptomic profiles. <i>Acta Neuropathologica Communications</i> , 2016, 4, 20.	5.2	136
8	Nuclear relocation of STAT6 reliably predicts NAB2-STAT6 fusion for the diagnosis of solitary fibrous tumour. <i>Histopathology</i> , 2014, 65, 613-622.	2.9	101
9	AKT1E17K mutations cluster with meningotheial and transitional meningiomas and can be detected by SFRP1 immunohistochemistry. <i>Acta Neuropathologica</i> , 2013, 126, 757-762.	7.7	88
10	D-2-Hydroxyglutarate producing neo-enzymatic activity inversely correlates with frequency of the type of isocitrate dehydrogenase 1 mutations found in glioma. <i>Acta Neuropathologica Communications</i> , 2014, 2, 19.	5.2	72
11	Primary mismatch repair deficient IDH-mutant astrocytoma (PMMRDIA) is a distinct type with a poor prognosis. <i>Acta Neuropathologica</i> , 2021, 141, 85-100.	7.7	52
12	BRAF V600E analysis for the differentiation of papillary craniopharyngiomas and Rathke's cleft cysts. <i>Neuropathology and Applied Neurobiology</i> , 2015, 41, 733-742.	3.2	50
13	Infratentorial IDH-mutant astrocytoma is a distinct subtype. <i>Acta Neuropathologica</i> , 2020, 140, 569-581.	7.7	45
14	Papillary glioneuronal tumor (PGNT) exhibits a characteristic methylation profile and fusions involving PRKCA. <i>Acta Neuropathologica</i> , 2019, 137, 837-846.	7.7	43
15	DNA methylation profiling reliably distinguishes pulmonary enteric adenocarcinoma from metastatic colorectal cancer. <i>Modern Pathology</i> , 2019, 32, 855-865.	5.5	36
16	Recurrent fusions in PLAGL1 define a distinct subset of pediatric-type supratentorial neuroepithelial tumors. <i>Acta Neuropathologica</i> , 2021, 142, 827-839.	7.7	33
17	Clear cell meningiomas are defined by a highly distinct DNA methylation profile and mutations in SMARCE1. <i>Acta Neuropathologica</i> , 2021, 141, 281-290.	7.7	31
18	MYCN and HDAC5 transcriptionally repress CD9 to trigger invasion and metastasis in neuroblastoma. <i>Oncotarget</i> , 2016, 7, 66344-66359.	1.8	30

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19	The genetic landscape of choroid plexus tumors in children and adults. <i>Neuro-Oncology</i> , 2021, 23, 650-660.	1.2	26
20	Next generation sequencing of lung adenocarcinoma subtypes with intestinal differentiation reveals distinct molecular signatures associated with histomorphology and therapeutic options. <i>Lung Cancer</i> , 2019, 138, 43-51.	2.0	24
21	Glioblastomas with primitive neuronal component harbor a distinct methylation and copy-number profile with inactivation of TP53, PTEN, and RB1. <i>Acta Neuropathologica</i> , 2021, 142, 179-189.	7.7	24
22	Molecular characterization of CNS paragangliomas identifies cauda equina paragangliomas as a distinct tumor entity. <i>Acta Neuropathologica</i> , 2020, 140, 893-906.	7.7	19
23	Oligosarcomas, IDH-mutant are distinct and aggressive. <i>Acta Neuropathologica</i> , 2022, 143, 263-281.	7.7	18
24	RNA-based analysis of ALK fusions in non-small cell lung cancer cases showing IHC/FISH discordance. <i>BMC Cancer</i> , 2018, 18, 1158.	2.6	17
25	TERT promoter mutation and chromosome 6 loss define a high-risk subtype of ependymoma evolving from posterior fossa subependymoma. <i>Acta Neuropathologica</i> , 2021, 141, 959-970.	7.7	16
26	Histopathological patterns in atypical teratoid/rhabdoid tumors are related to molecular subgroup. <i>Brain Pathology</i> , 2021, 31, e12967.	4.1	16
27	Inflammatory features in sporadic late-onset nemaline myopathy are independent from monoclonal gammopathy. <i>Brain Pathology</i> , 2021, 31, e12962.	4.1	15
28	Neurofibromatosis type 2 predisposes to ependymomas of various localization, histology, and molecular subtype. <i>Acta Neuropathologica</i> , 2021, 141, 971-974.	7.7	12
29	Pleomorphic xanthoastrocytoma is a heterogeneous entity with pTERT mutations prognosticating shorter survival. <i>Acta Neuropathologica Communications</i> , 2022, 10, 5.	5.2	12
30	Comprehensive profiling of myxopapillary ependymomas identifies a distinct molecular subtype with relapsing disease. <i>Neuro-Oncology</i> , 2022, 24, 1689-1699.	1.2	11
31	Cauda equina paragangliomas express HOXB13. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 889-890.	3.2	9
32	Predicting survival in anaplastic astrocytoma patients in a single-center cohort of 108 patients. <i>Radiation Oncology</i> , 2020, 15, 282.	2.7	6
33	Evidence of neural crest cell origin of a DICER1 mutant CNS sarcoma in a child with DICER1 syndrome and NRAS-mutant neurocutaneous melanosis. <i>Neuropathology and Applied Neurobiology</i> , 2022, 48, .	3.2	4
34	Molecular characterisation of sporadic endolymphatic sac tumours and comparison to von Hippel-Lindau disease-related tumours. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 756-767.	3.2	2
35	Fulminant cerebral venous thrombosis associated with the m.3243A>G MELAS mutation: A new guise for an old disease. <i>Brain and Development</i> , 2019, 41, 901-904.	1.1	1
36	Extramedullary relapse after accidental head injury in a patient with multiple myeloma. <i>Annals of Hematology</i> , 2015, 94, 351-353.	1.8	0