Stefan M Pfister

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

574	57,205	111	227
papers	citations	h-index	g-index
625 ext. papers	73,938 ext. citations	11.3 avg, IF	7.78 L-index

#	Paper	IF	Citations
574	Important Requirements for the Selection of Internal Standards during the Development of Desorption/Ionization Assays for Drug Quantification in Biological Matrices-A Practical Example <i>Molecules</i> , 2022 , 27,	4.8	1
573	Pleomorphic xanthoastrocytoma is a heterogeneous entity with pTERT mutations prognosticating shorter survival <i>Acta Neuropathologica Communications</i> , 2022 , 10, 5	7.3	2
572	Integrative gene network and functional analyses identify a prognostically relevant key regulator of metastasis in Ewing sarcoma <i>Molecular Cancer</i> , 2022 , 21, 1	42.1	3
571	Clinical and molecular characterization of isolated M1 disease in pediatric medulloblastoma: experience from the German HIT-MED studies <i>Journal of Neuro-Oncology</i> , 2022 , 157, 37	4.8	
570	Predictive modeling of resistance to SMO inhibition in a patient-derived orthotopic xenograft model of SHH medulloblastoma <i>Neuro-Oncology Advances</i> , 2022 , 4, vdac026	0.9	O
569	Rapid-CNS: rapid comprehensive adaptive nanopore-sequencing of CNS tumors, a proof-of-concept study <i>Acta Neuropathologica</i> , 2022 , 1	14.3	O
568	The genomic landscape of pediatric renal cell carcinomas <i>IScience</i> , 2022 , 25, 104167	6.1	O
567	MRI Radiogenomics of Pediatric Medulloblastoma: A Multicenter Study Radiology, 2022, 212137	20.5	2
566	MODL-02. A novelCre-conditionalcMYC-driven MB Group 3 transgenic mouse model shows traceable leptomeningeal dissemination <i>Neuro-Oncology</i> , 2022 , 24, i168-i168	1	
565	RARE-12. Pineoblastoma of children and young adults in a national population: An analysis of the HIT-MED study cohort. <i>Neuro-Oncology</i> , 2022 , 24, i11-i12	1	
564	HGG-50. Specific sensitivity of pediatric high-grade glioma with ATRX inactivation to PARP inhibitor combinations. <i>Neuro-Oncology</i> , 2022 , 24, i73-i73	1	
563	PATH-08. DNA methylation profiling improves routine diagnostics of paediatric CNS tumours: a prospective population-based study. <i>Neuro-Oncology</i> , 2022 , 24, i159-i160	1	
562	HGG-27. Understanding the role of PLAG family transcription factors in cortex development and tumorigenesis. <i>Neuro-Oncology</i> , 2022 , 24, i66-i66	1	
561	MODL-04. Drug screening in Disorders with Abnormal DNA Damage Response/Repair (DADDR) andin vivo validation. <i>Neuro-Oncology</i> , 2022 , 24, i168-i169	1	
560	MEDB-60. Medulloblastoma with extensive nodularity mimics cerebellar development and differentiates along the granular precursor lineage. <i>Neuro-Oncology</i> , 2022 , 24, i120-i120	1	
559	MEDB-38. Significance of CSF cytology and neurologic deterioration in relapsed medulloblastomas in the German HIT-REZ-97/-2005 Studies and the HIT-REZ-Register. <i>Neuro-Oncology</i> , 2022 , 24, i113-i114	1	
558	MEDB-14. Clinical outcome of pediatric medulloblastoma patients with Li-Fraumeni syndrome. <i>Neuro-Oncology</i> , 2022 , 24, i107-i107	1	

557	OTHR-32. The Pediatric Targeted Therapy 2.0 registry: robust molecular diagnostics for precision oncology. <i>Neuro-Oncology</i> , 2022 , 24, i154-i154	1
556	PATH-13. Methylation analysis in the diagnosis of pediatric CNS tumors; a single center experience. <i>Neuro-Oncology</i> , 2022 , 24, i161-i161	1
555	Analytical Performance Evaluation of New DESI Enhancements for Targeted Drug Quantification in Tissue Sections. <i>Pharmaceuticals</i> , 2022 , 15, 694	5.2 1
554	MEDB-41. Identifying a subgroup of patients with early childhood sonic hedgehog-activated medulloblastoma with unfavorable prognosis after treatment with radiation-sparing regimens including intraventricular methotrexate. <i>Neuro-Oncology</i> , 2022 , 24, i114-i115	1
553	PATH-03. Clinically Tractable Outcome Prediction of Group 3/4 Medulloblastoma Based on TPD52 Immunohistochemistry: a Multicohort Study. <i>Neuro-Oncology</i> , 2022 , 24, i158-i158	1
552	HGG-11. Clinical characteristics and clinical evolution of a large cohort of pediatric patients with primary central nervous system (CNS) tumors and tropomyosin receptor kinase (TRK) fusion <i>Neuro-Oncology</i> , 2022 , 24, i61-i62	1
551	MEDB-36. Clinical and molecular heterogeneity withinMYC andMYCN amplified medulloblastoma. <i>Neuro-Oncology</i> , 2022 , 24, i113-i113	1
550	LGG-14. LOGGIC (Low Grade Glioma in Children) Core BioClinical Data Bank: Establishment and added clinical value of an international molecular diagnostic registry for pediatric low-grade glioma patients. <i>Neuro-Oncology</i> , 2022 , 24, i90-i90	1
549	MEDB-15. Dynamic chromatin alteration induces oncogenic hijacking by essential transcriptional factors during SHH medulloblastoma tumorigenesis. <i>Neuro-Oncology</i> , 2022 , 24, i107-i108	1
548	MODL-07. DNA methylation-based biobank of murine models for pediatric tumors. <i>Neuro-Oncology</i> , 2022 , 24, i169-i170	1
547	EPEN-28. Oncogenic dependency of pediatric ependymomas on extracellular vesicle pathways. <i>Neuro-Oncology</i> , 2022 , 24, i45-i45	1
546	IMMU-04. Transcriptional analysis reveals distinct microenvironmental subgroups across pediatric nervous system tumors. <i>Neuro-Oncology</i> , 2022 , 24, i81-i81	1
545	HGG-61.Landscape of cancer predisposition in pediatric high-grade glioma. <i>Neuro-Oncology</i> , 2022 , 24, i76-i76	1
544	EPEN-09. Multi-omics characterization of the blood-brain barrier in molecular groups of ependymoma. <i>Neuro-Oncology</i> , 2022 , 24, i40-i40	1
543	DIPG-19. FOXR2 is an oncogenic driver across pediatric and adult cancers. <i>Neuro-Oncology</i> , 2022 , 24, i21-i22	1
542	PATH-11. Detection of genetic and epigenetic alterations in Liquid Biopsies from pediatric brain tumor patients. <i>Neuro-Oncology</i> , 2022 , 24, i160-i161	1
541	MEDB-04. Young children with metastatic medulloblastoma: frequent requirement for radiotherapy in children with non-WNT/non-SHH medulloblastoma despite highly intensified chemotherapy [Results of the MET-HIT2000-BIS4 trial. <i>Neuro-Oncology</i> , 2022 , 24, i104-i104	1
540	RARE-15. Astroblastoma, MN1 altered comprises two molecularly and clinically distinct subgroups defined by the fusion partners BEND2 and CXXC5. <i>Neuro-Oncology</i> , 2022 , 24, i12-i13	1

539	THER-01. Precision brain tumor therapy by AAV-mediated oncogene editing. <i>Neuro-Oncology</i> , 2022 , 24, i185-i186	1	
538	MODL-01. Targeting replication stress in pediatric brain tumors. <i>Neuro-Oncology</i> , 2022 , 24, i168-i168	1	
537	Target actionability review to evaluate CDK4/6 as a therapeutic target in paediatric solid and brain tumours. <i>European Journal of Cancer</i> , 2022 , 170, 196-208	7.5	1
536	EZHIP: a new piece of the puzzle towards understanding pediatric posterior fossa ependymoma. <i>Acta Neuropathologica</i> , 2021 , 143, 1	14.3	3
535	Target Actionability Review: a systematic evaluation of replication stress as a therapeutic target for paediatric solid malignancies <i>European Journal of Cancer</i> , 2021 , 162, 107-117	7.5	О
534	Systemic chemotherapy of pediatric recurrent ependymomas: results from the German HIT-REZ studies. <i>Journal of Neuro-Oncology</i> , 2021 , 155, 193-202	4.8	2
533	Genetic and epigenetic characterization of posterior pituitary tumors. <i>Acta Neuropathologica</i> , 2021 , 142, 1025-1043	14.3	1
532	Primary central nervous system sarcoma with DICER1 mutation-treatment results of a novel molecular entity in pediatric Peruvian patients. <i>Cancer</i> , 2021 ,	6.4	2
531	Integrated Molecular-Morphologic Meningioma Classification: A Multicenter Retrospective Analysis, Retrospectively and Prospectively Validated. <i>Journal of Clinical Oncology</i> , 2021 , 39, 3839-3852	2.2	8
530	A systematic analysis of genetic interactions and their underlying biology in childhood cancer. <i>Communications Biology</i> , 2021 , 4, 1139	6.7	O
529	Targeting integrated epigenetic and metabolic pathways in lethal childhood PFA ependymomas. <i>Science Translational Medicine</i> , 2021 , 13, eabc0497	17.5	3
528	Pr⊠isionsonkologie und Phase-I/II-Netzwerke in der Kinderkrebsmedizin 2021 , 36, 485	0.2	
527	Spatial Dissection of Invasive Front from Tumor Mass Enables Discovery of Novel microRNA Drivers of Glioblastoma Invasion. <i>Advanced Science</i> , 2021 , 8, e2101923	13.6	2
526	Clear cell meningiomas are defined by a highly distinct DNA methylation profile and mutations in SMARCE1. <i>Acta Neuropathologica</i> , 2021 , 141, 281-290	14.3	9
525	Local and systemic therapy of recurrent ependymoma in children and adolescents: short- and long-term results of the E-HIT-REZ 2005 study. <i>Neuro-Oncology</i> , 2021 , 23, 1012-1023	1	10
524	The age of adult pilocytic astrocytoma cells. <i>Oncogene</i> , 2021 , 40, 2830-2841	9.2	2
523	Notch Signaling between Cerebellar Granule Cell Progenitors. <i>ENeuro</i> , 2021 , 8,	3.9	3
522	ZFTA-RELA Dictates Oncogenic Transcriptional Programs to Drive Aggressive Supratentorial Ependymoma. <i>Cancer Discovery</i> , 2021 , 11, 2200-2215	24.4	16

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521	Translocations Constitute Ependymoma Chromatin Remodeling and Transcription Factors. <i>Cancer Discovery</i> , 2021 , 11, 2216-2229	24.4	13
520	Outcomes by Clinical and Molecular Features in Children With Medulloblastoma Treated With Risk-Adapted Therapy: Results of an International Phase III Trial (SJMB03). <i>Journal of Clinical Oncology</i> , 2021 , 39, 822-835	2.2	25
519	H3.3-K27M drives neural stem cell-specific gliomagenesis in a human iPSC-derived model. <i>Cancer Cell</i> , 2021 , 39, 407-422.e13	24.3	13
518	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	14.3	5
517	Cross-Species Genomics Reveals Oncogenic Dependencies in ZFTA/C11orf95 Fusion-Positive Supratentorial Ependymomas. <i>Cancer Discovery</i> , 2021 , 11, 2230-2247	24.4	20
516	Clinicopathologic and molecular analysis of embryonal rhabdomyosarcoma of the genitourinary tract: evidence for a distinct DICER1-associated subgroup. <i>Modern Pathology</i> , 2021 , 34, 1558-1569	9.8	5
515	Cancer predisposition in pediatric neuro-oncology-practical approaches and ethical considerations. <i>Neuro-Oncology Practice</i> , 2021 , 8, 526-538	2.2	1
514	Carbon ion radiotherapy eradicates medulloblastomas with chromothripsis in an orthotopic Li-Fraumeni patient-derived mouse model. <i>Neuro-Oncology</i> , 2021 , 23, 2028-2041	1	1
513	Single cell derived mRNA signals across human kidney tumors. <i>Nature Communications</i> , 2021 , 12, 3896	17.4	4
512	FOXR2 Stabilizes MYCN Protein and Identifies NonAmplified Neuroblastoma Patients With Unfavorable Outcome. <i>Journal of Clinical Oncology</i> , 2021 , 39, 3217-3228	2.2	4
511	EMBR-01. CLASS I HDAC INHIBITORS AND PLK1 INHIBITORS SYNERGIZE IN MYC-AMPLIFIED MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2021 , 23, i5-i5	1	78
510	The 2021 WHO Classification of Tumors of the Central Nervous System: a summary. <i>Neuro-Oncology</i> , 2021 , 23, 1231-1251	1	708
509	EMBR-21. CLINICALLY TRACTABLE OUTCOME PREDICTION OF GROUP 3/4 MEDULLOBLASTOMA BASED ON TPD52 IMMUNOHISTOCHEMISTRY: A MULTICOHORT STUDY. <i>Neuro-Oncology</i> , 2021 , 23, i10-	i 1 0	78
508	IMMU-14. COMPUTATIONAL DECONVOLUTION OF TUMOR-INFILTRATING IMMUNE COMPONENTS IN PEDIATRIC NERVOUS SYSTEM TUMORS. <i>Neuro-Oncology</i> , 2021 , 23, i30-i30	1	78
507	International Consensus on Minimum Preclinical Testing Requirements for the Development of Innovative Therapies For Children and Adolescents with Cancer. <i>Molecular Cancer Therapeutics</i> , 2021 , 20, 1462-1468	6.1	3
506	From Sampling to Sequencing: A Liquid Biopsy Pre-Analytic Workflow to Maximize Multi-Layer Genomic Information from a Single Tube. <i>Cancers</i> , 2021 , 13,	6.6	3
505	High-Resolution Cartography of the Transcriptome and Methylome Landscapes of Diffuse Gliomas. <i>Cancers</i> , 2021 , 13,	6.6	4
504	LGG-04. MULTIOMIC ANALYSIS OF MAPK PATHWAY ACTIVITY IN PEDIATRIC PILOCYTIC ASTROCYTOMA. <i>Neuro-Oncology</i> , 2021 , 23, i31-i32	1	

503	Therapeutic implications of improved molecular diagnostics for rare CNS embryonal tumor entities: results of an international, retrospective study. <i>Neuro-Oncology</i> , 2021 , 23, 1597-1611	1	3
502	Bioanalysis of selinexor in mouse plasma micro-samples utilizing UPLC-MS/MS. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2021 , 1176, 122781	3.2	1
501	TMOD-03. A NOVEL MB GR3 TRANSGENIC MOUSE MODEL IS GENERATED BY MYCN AND P53 DEFECTS IN VENTRICULAR ZONE PROGENITORS <i>Neuro-Oncology</i> , 2021 , 23, i36-i36	1	
500	LGG-06. COMPREHENSIVE GENOMIC CHARACTERIZATION AND INTEGRATED CLINICAL ANALYSIS OF LOW-GRADE GLIOMAS IN CHILDREN WITH NEUROFIBROMATOSIS TYPE 1. <i>Neuro-Oncology</i> , 2021 , 23, i32-i32	1	78
499	EPCT-06. PRECISION ONCOLOGY IN THE PEDIATRIC TARGETED THERAPY 2.0 PROGRAM. Neuro-Oncology, 2021 , 23, i47-i48	1	78
498	EPEN-03. ZFTA/C11ORF95 FUSIONS DRIVE SUPRATENTORIAL EPENDYMOMA VIA SHARED ONCOGENIC MECHANISMS. <i>Neuro-Oncology</i> , 2021 , 23, i13-i14	1	78
497	LGG-11. BH3-MIMETICS TARGETING BCL-XL SELECTIVELY IMPACT THE SENESCENT COMPARTMENT OF PILOCYTIC ASTROCYTOMA. <i>Neuro-Oncology</i> , 2021 , 23, i33-i34	1	78
496	Development of Randomized Trials in Adults with Medulloblastoma-The Example of EORTC 1634-BTG/NOA-23. <i>Cancers</i> , 2021 , 13,	6.6	2
495	Emergence and maintenance of actionable genetic drivers at medulloblastoma relapse. <i>Neuro-Oncology</i> , 2021 ,	1	3
494	Intimal sarcomas and undifferentiated cardiac sarcomas carry mutually exclusive MDM2, MDM4, and CDK6 amplifications and share a common DNA methylation signature. <i>Modern Pathology</i> , 2021 , 34, 2122-2129	9.8	4
493	Second series by the Italian Association of Pediatric Hematology and Oncology of children and adolescents with intracranial ependymoma: an integrated molecular and clinical characterization with a long-term follow-up. <i>Neuro-Oncology</i> , 2021 , 23, 848-857	1	7
492	Downregulation of miR-326 and its host gene [larrestin1 induces pro-survival activity of E2F1 and promotes medulloblastoma growth. <i>Molecular Oncology</i> , 2021 , 15, 523-542	7.9	5
491	Primary mismatch repair deficient IDH-mutant astrocytoma (PMMRDIA) is a distinct type with a poor prognosis. <i>Acta Neuropathologica</i> , 2021 , 141, 85-100	14.3	14
490	An extracellular vesicle-related gene expression signature identifies high-risk patients in medulloblastoma. <i>Neuro-Oncology</i> , 2021 , 23, 586-598	1	2
489	Super enhancers define regulatory subtypes and cell identity in neuroblastoma <i>Nature Cancer</i> , 2021 , 2, 114-128	15.4	15
488	Accurate calling of KIAA1549-BRAF fusions from DNA of human brain tumours using methylation array-based copy number and gene panel sequencing data. <i>Neuropathology and Applied Neurobiology</i> , 2021 , 47, 406-414	5.2	2
487	DNA methylation based glioblastoma subclassification is related to tumoral T-cell infiltration and patient survival. <i>Neuro-Oncology</i> , 2021 , 23, 240-250	1	9
486	A subset of pediatric-type thalamic gliomas share a distinct DNA methylation profile, H3K27me3 loss and frequent alteration of EGFR. <i>Neuro-Oncology</i> , 2021 , 23, 34-43	1	22

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485	Reduced chromatin binding of MYC is a key effect of HDAC inhibition in MYC amplified medulloblastoma. <i>Neuro-Oncology</i> , 2021 , 23, 226-239	1	6
484	Thrombospondin-1 mimetics are promising novel therapeutics for MYC-associated medulloblastoma. <i>Neuro-Oncology Advances</i> , 2021 , 3, vdab002	0.9	O
483	ABCB1 inhibition provides a novel therapeutic target to block TWIST1-induced migration in medulloblastoma. <i>Neuro-Oncology Advances</i> , 2021 , 3, vdab030	0.9	
482	Small-molecule screen reveals synergy of cell cycle checkpoint kinase inhibitors with DNA-damaging chemotherapies in medulloblastoma. <i>Science Translational Medicine</i> , 2021 , 13,	17.5	7
481	Rapid MALDI-MS Assays for Drug Quantification in Biological Matrices: Lessons Learned, New Developments, and Future Perspectives. <i>Molecules</i> , 2021 , 26,	4.8	5
480	Clinical and molecular heterogeneity of pineal parenchymal tumors: a consensus study. <i>Acta Neuropathologica</i> , 2021 , 141, 771-785	14.3	9
479	Alternative lengthening of telomeres in childhood neuroblastoma from genome to proteome. <i>Nature Communications</i> , 2021 , 12, 1269	17.4	12
478	Molecular analysis of pediatric CNS-PNET revealed nosologic heterogeneity and potent diagnostic markers for CNS neuroblastoma with FOXR2-activation. <i>Acta Neuropathologica Communications</i> , 2021 , 9, 20	7.3	2
477	Integrated molecular analysis of adult sonic hedgehog (SHH)-activated medulloblastomas reveals two clinically relevant tumor subsets with VEGFA as potent prognostic indicator. <i>Neuro-Oncology</i> , 2021 , 23, 1576-1585	1	3
476	Alterations in Pediatric High-Risk Malignancies Identified Through European Clinical Sequencing Programs Constitute Promising Drug Targets <i>JCO Precision Oncology</i> , 2021 , 5, 450-454	3.6	
475	Integrated molecular and clinical analysis of low-grade gliomas in children with neurofibromatosis type 1 (NF1). <i>Acta Neuropathologica</i> , 2021 , 141, 605-617	14.3	9
474	Maturation Block in Childhood Cancer. <i>Cancer Discovery</i> , 2021 , 11, 542-544	24.4	4
473	Ultra high-risk PFA ependymoma is characterized by loss of chromosome 6q. <i>Neuro-Oncology</i> , 2021 , 23, 1360-1370	1	14
472	Clinical Outcomes and Patient-Matched Molecular Composition of Relapsed Medulloblastoma. <i>Journal of Clinical Oncology</i> , 2021 , 39, 807-821	2.2	7
471	PATZ1 fusions define a novel molecularly distinct neuroepithelial tumor entity with a broad histological spectrum. <i>Acta Neuropathologica</i> , 2021 , 142, 841-857	14.3	7
470	Recurrent fusions in PLAGL1 define a distinct subset of pediatric-type supratentorial neuroepithelial tumors. <i>Acta Neuropathologica</i> , 2021 , 142, 827-839	14.3	5
469	The Pediatric Precision Oncology INFORM Registry: Clinical Outcome and Benefit for Patients with Very High-Evidence Targets. <i>Cancer Discovery</i> , 2021 , 11, 2764-2779	24.4	22
468	Subgroup and subtype-specific outcomes in adult medulloblastoma. <i>Acta Neuropathologica</i> , 2021 , 142, 859-871	14.3	2

467	Developmental and evolutionary dynamics of cis-regulatory elements in mouse cerebellar cells. <i>Science</i> , 2021 , 373,	33.3	8
466	Radiation-induced gliomas represent H3-/IDH-wild type pediatric gliomas with recurrent PDGFRA amplification and loss of CDKN2A/B. <i>Nature Communications</i> , 2021 , 12, 5530	17.4	3
465	GOPC:ROS1 and other ROS1 fusions represent a rare but recurrent drug target in a variety of glioma types. <i>Acta Neuropathologica</i> , 2021 , 142, 1065-1069	14.3	1
464	Molecular profiling of pediatric meningiomas shows tumor characteristics distinct from adult meningiomas. <i>Acta Neuropathologica</i> , 2021 , 142, 873-886	14.3	1
463	Sarcoma classification by DNA methylation profiling. <i>Nature Communications</i> , 2021 , 12, 498	17.4	74
462	A Summary of the Inaugural WHO Classification of Pediatric Tumors: Transitioning from the Optical into the Molecular Era <i>Cancer Discovery</i> , 2021 ,	24.4	5
461	Molecular correlates of cerebellar mutism syndrome in medulloblastoma. <i>Neuro-Oncology</i> , 2020 , 22, 290-297	1	8
460	Evaluation of Prognostic Factors and Role of Participation in a Randomized Trial or a Prospective Registry in Pediatric and Adolescent Nonmetastatic Medulloblastoma - A Report From the HIT 2000 Trial. <i>Advances in Radiation Oncology</i> , 2020 , 5, 1158-1169	3.3	6
459	Systematic target actionability reviews of preclinical proof-of-concept papers to match targeted drugs to paediatric cancers. <i>European Journal of Cancer</i> , 2020 , 130, 168-181	7.5	4
458	An optimized workflow to improve reliability of detection of KIAA1549:BRAF fusions from RNA sequencing data. <i>Acta Neuropathologica</i> , 2020 , 140, 237-239	14.3	3
457	INFORM2 NivEnt: The first trial of the INFORM2 biomarker driven phase I/II trial series: the combination of nivolumab and entinostat in children and adolescents with refractory high-risk malignancies. <i>BMC Cancer</i> , 2020 , 20, 523	4.8	11
456	Histone H3 wild-type DIPG/DMG overexpressing EZHIP extend the spectrum diffuse midline gliomas with PRC2 inhibition beyond H3-K27M mutation. <i>Acta Neuropathologica</i> , 2020 , 139, 1109-1113	14.3	33
455	Germline Elongator mutations in Sonic Hedgehog medulloblastoma. <i>Nature</i> , 2020 , 580, 396-401	50.4	47
454	ETMR: a tumor entity in its infancy. <i>Acta Neuropathologica</i> , 2020 , 140, 249-266	14.3	20
453	CDKN2A/B homozygous deletion is associated with early recurrence in meningiomas. <i>Acta Neuropathologica</i> , 2020 , 140, 409-413	14.3	26
452	Implications of new understandings of gliomas in children and adults with NF1: report of a consensus conference. <i>Neuro-Oncology</i> , 2020 , 22, 773-784	1	21
451	Larotrectinib in patients with TRK fusion-positive solid tumours: a pooled analysis of three phase 1/2 clinical trials. <i>Lancet Oncology, The</i> , 2020 , 21, 531-540	21.7	279
450	Pilocytic astrocytoma demethylation and transcriptional landscapes link bZIP transcription factors to immune response. <i>Neuro-Oncology</i> , 2020 , 22, 1327-1338	1	4

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449	Rapid and Sensitive Drug Quantification in Tissue Sections Using Matrix Assisted Laser Desorption Ionization-Ion Mobility-Mass Spectrometry Profiling. <i>Journal of the American Society for Mass Spectrometry</i> , 2020 , 31, 742-751	3.5	11
448	Pan-cancer analysis of whole genomes. <i>Nature</i> , 2020 , 578, 82-93	50.4	840
447	Nonmetastatic Medulloblastoma of Early Childhood: Results From the Prospective Clinical Trial HIT-2000 and An Extended Validation Cohort. <i>Journal of Clinical Oncology</i> , 2020 , 38, 2028-2040	2.2	21
446	Locoregionally administered B7-H3-targeted CAR T cells for treatment of atypical teratoid/rhabdoid tumors. <i>Nature Medicine</i> , 2020 , 26, 712-719	50.5	74
445	A Cell-Based MAPK Reporter Assay Reveals Synergistic MAPK Pathway Activity Suppression by MAPK Inhibitor Combination in -Driven Pediatric Low-Grade Glioma Cells. <i>Molecular Cancer Therapeutics</i> , 2020 , 19, 1736-1750	6.1	5
444	The pediatric precision oncology study INFORM: Clinical outcome and benefit for molecular subgroups <i>Journal of Clinical Oncology</i> , 2020 , 38, LBA10503-LBA10503	2.2	9
443	EPEN-09. IMPACT OF MOLECULAR SUBGROUP ON OUTCOME FOR INFANTS . <i>Neuro-Oncology</i> , 2020 , 22, iii309-iii309	1	78
442	QOL-13. NEUROCOGNITIVE OUTCOMES ACCORDING TO RISK-ADAPTED TREATMENT REGIMENS FOR CHILDREN OLDER THAN 4 WITH MEDULLOBLASTOMA AND POSTERIOR FOSSA EPENDYMOMA IRESULTS OF THE HIT2000 TRIAL. <i>Neuro-Oncology</i> , 2020 , 22, iii433-iii433	1	78
441	EPEN-36. THE TREATMENT OUTCOME OF PAEDIATRIC SUPRATENTORIAL C11ORF95-RELA FUSED EPENDYMOMA: A COMBINED REPORT FROM E-HIT SERIES AND AUSTRALIAN NEW ZEALAND CHILDRENS HAEMATOLOGY/ONCOLOGY GROUP. <i>Neuro-Oncology</i> , 2020 , 22, iii315-iii315	1	78
440	MBCL-11. TIME TO RADIOTHERAPY IMPACTS SURVIVAL IN PEDIATRIC AND ADOLESCENT NON-METASTATIC MEDULLOBLASTOMA TREATED BY UPFRONT RADIOTHERAPY A REPORT FROM THE HIT 2000 TRIAL. <i>Neuro-Oncology</i> , 2020 , 22, iii389-iii390	1	78
439	HGG-56. EXTENSIVE MOLECULAR HETEROGENEITY WITHIN H3-/IDH-WILDTYPE PEDIATRIC GLIOBLASTOMA. <i>Neuro-Oncology</i> , 2020 , 22, iii354-iii354	1	78
438	ETMR-21. META-ANALYSIS OF PINEAL REGION TUMOURS DEMONSTRATES MOLECULAR SUBGROUPS WITH DISTINCT CLINICO-PATHOLOGICAL FEATURES: A CONSENSUS STUDY. <i>Neuro-Oncology</i> , 2020 , 22, iii327-iii327	1	78
437	EPEN-18. CROSS-SPECIES GENOMICS IDENTIFIES GLI2 AS AN ONCOGENE OF C11orf95 FUSION-POSITIVE SUPRATENTORIAL EPENDYMOMA. <i>Neuro-Oncology</i> , 2020 , 22, iii311-iii311	1	78
436	EPEN-44. EXTRACELLULAR VESICLES OF SUPRATENTORIAL EPENDYMOMA RELA MEDIATE INTERACTIONS WITH CELLS OF THE TUMOR MICROENVIRONMENT. <i>Neuro-Oncology</i> , 2020 , 22, iii316-iii	3 ¹ 17	78
435	EPEN-39. CLINICAL STRATIFIED TREATMENT OF LOCALIZED PEDIATRIC INTRACRANIAL EPENDYMOMA WITH COMBINED LOCAL IRRADIATION AND CHEMOTHERAPY WITHIN THE PROSPECTIVE, MULTICENTER E-HIT TRIAL ITHE MOLECULAR SUBGROUP MATTERS.	1	0
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411	Germline Mutations Predispose to Pediatric Medulloblastoma. <i>Journal of Clinical Oncology</i> , 2020 , 38, 43-50	2.2	28
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354	Recurrent noncoding U1IsnRNA mutations drive cryptic splicing in SHH medulloblastoma. <i>Nature</i> , 2019 , 574, 707-711	50.4	78
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	32, 886-96 Whole exome sequencing reveals that the majority of schwannomatosis cases remain unexplained		
151	Whole exome sequencing reveals that the majority of schwannomatosis cases remain unexplained after excluding SMARCB1 and LZTR1 germline variants. <i>Acta Neuropathologica</i> , 2014 , 128, 449-52 Phase II study of sorafenib in children with recurrent or progressive low-grade astrocytomas.	14.3	28
151 150	32, 886-96 Whole exome sequencing reveals that the majority of schwannomatosis cases remain unexplained after excluding SMARCB1 and LZTR1 germline variants. <i>Acta Neuropathologica</i> , 2014 , 128, 449-52 Phase II study of sorafenib in children with recurrent or progressive low-grade astrocytomas. <i>Neuro-Oncology</i> , 2014 , 16, 1408-16 Integrated DNA methylation and copy-number profiling identify three clinically and biologically	14.3	28
151 150 149	Whole exome sequencing reveals that the majority of schwannomatosis cases remain unexplained after excluding SMARCB1 and LZTR1 germline variants. <i>Acta Neuropathologica</i> , 2014 , 128, 449-52 Phase II study of sorafenib in children with recurrent or progressive low-grade astrocytomas. <i>Neuro-Oncology</i> , 2014 , 16, 1408-16 Integrated DNA methylation and copy-number profiling identify three clinically and biologically relevant groups of anaplastic glioma. <i>Acta Neuropathologica</i> , 2014 , 128, 561-71 Farewell to oligoastrocytoma: in situ molecular genetics favor classification as either	14.3 1 14.3	28 140 148
151 150 149 148	Whole exome sequencing reveals that the majority of schwannomatosis cases remain unexplained after excluding SMARCB1 and LZTR1 germline variants. <i>Acta Neuropathologica</i> , 2014 , 128, 449-52 Phase II study of sorafenib in children with recurrent or progressive low-grade astrocytomas. <i>Neuro-Oncology</i> , 2014 , 16, 1408-16 Integrated DNA methylation and copy-number profiling identify three clinically and biologically relevant groups of anaplastic glioma. <i>Acta Neuropathologica</i> , 2014 , 128, 561-71 Farewell to oligoastrocytoma: in situ molecular genetics favor classification as either oligodendroglioma or astrocytoma. <i>Acta Neuropathologica</i> , 2014 , 128, 551-9	14.3 1 14.3	28 140 148 200
151 150 149 148	Whole exome sequencing reveals that the majority of schwannomatosis cases remain unexplained after excluding SMARCB1 and LZTR1 germline variants. <i>Acta Neuropathologica</i> , 2014 , 128, 449-52 Phase II study of sorafenib in children with recurrent or progressive low-grade astrocytomas. <i>Neuro-Oncology</i> , 2014 , 16, 1408-16 Integrated DNA methylation and copy-number profiling identify three clinically and biologically relevant groups of anaplastic glioma. <i>Acta Neuropathologica</i> , 2014 , 128, 561-71 Farewell to oligoastrocytoma: in situ molecular genetics favor classification as either oligodendroglioma or astrocytoma. <i>Acta Neuropathologica</i> , 2014 , 128, 551-9 Enhancer hijacking activates GFI1 family oncogenes in medulloblastoma. <i>Nature</i> , 2014 , 511, 428-34 Genomic and transcriptomic analyses match medulloblastoma mouse models to their human	14.3 14.3 14.3	28 140 148 200

143	BI-30 * CHARACTERIZATION OF L1CAM AS A CLINICAL MARKER FOR THE C11orf95-RELA FUSION IN SUPRATENTORIAL EPENDYMOMAS. <i>Neuro-Oncology</i> , 2014 , 16, v30-v30	1	4
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118	Emerging insights into the ependymoma epigenome. <i>Brain Pathology</i> , 2013 , 23, 206-9	6	18
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88	Genome sequencing of pediatric medulloblastoma links catastrophic DNA rearrangements with TP53 mutations. <i>Cell</i> , 2012 , 148, 59-71	56.2	600
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43	DNA methylation pattern changes upon long-term culture and aging of human mesenchymal stromal cells. <i>Aging Cell</i> , 2010 , 9, 54-63	9.9	325
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35	Focal genomic amplification at 19q13.42 comprises a powerful diagnostic marker for embryonal tumors with ependymoblastic rosettes. <i>Acta Neuropathologica</i> , 2010 , 120, 253-60	14.3	115
34	Molecular diagnostics of CNS embryonal tumors. <i>Acta Neuropathologica</i> , 2010 , 120, 553-66	14.3	72
33	The genetics of pediatric brain tumors. Current Neurology and Neuroscience Reports, 2010, 10, 215-23	6.6	58
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31	Abstract 4347: Medulloblastoma comprises four distinct diseases 2010 ,		5
30	Outcome prediction in pediatric medulloblastoma based on DNA copy-number aberrations of chromosomes 6q and 17q and the MYC and MYCN loci. <i>Journal of Clinical Oncology</i> , 2009 , 27, 1627-36	2.2	238
29	From glioblastoma to gangliocytoma: an unforeseen but welcome shift in biological behavior. <i>Journal of Neurosurgery: Pediatrics</i> , 2009 , 4, 475-8	2.1	7
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16	High-Resolution Genomic Profiling (array-CGH) of Childhood T-ALL Identifies Deletions at 6q15-16.1 as a Predictive Marker for Early Treatment Response <i>Blood</i> , 2008 , 112, 1484-1484	2.2	
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7	CD28 signaling via VAV/SLP-76 adaptors: regulation of cytokine transcription independent of TCR ligation. <i>Immunity</i> , 2001 , 15, 921-33	32.3	88
6	TelomereHunter: telomere content estimation and characterization from whole genome sequencing data		11
5	Dissecting telomere maintenance mechanisms in pediatric glioblastoma		4
4	Single cell derived mRNA signals across human kidney tumors		4
3	Establishment of a simplified preparation method for single-nucleus RNA-sequencing and its application to long-term frozen tumor tissues		2
2	Integrated phospho-proteogenomic and single-cell transcriptomic analysis of meningiomas establishes robust subtyping and reveals subtype-specific immune invasion		2
1	Systematic multi-omics cell line profiling uncovers principles of Ewing sarcoma fusion oncogene-mediated gene regulation		1