## Kristian W Pajtler

## List of Publications by Citations

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102<br/>papers5,915<br/>citations32<br/>h-index76<br/>g-index115<br/>ext. papers8,319<br/>ext. citations8.8<br/>avg, IF5.1<br/>L-index

#	Paper	IF	Citations
102	DNA methylation-based classification of central nervous system tumours. <i>Nature</i> , <b>2018</b> , 555, 469-474	50.4	992
101	Molecular Classification of Ependymal Tumors across All CNS Compartments, Histopathological Grades, and Age Groups. <i>Cancer Cell</i> , <b>2015</b> , 27, 728-43	24.3	672
100	The landscape of genomic alterations across childhood cancers. <i>Nature</i> , <b>2018</b> , 555, 321-327	50.4	603
99	New Brain Tumor Entities Emerge from Molecular Classification of CNS-PNETs. <i>Cell</i> , <b>2016</b> , 164, 1060-10	) <b>7<del>3</del>6.</b> 2	483
98	Cancer Screening Recommendations for Individuals with Li-Fraumeni Syndrome. <i>Clinical Cancer Research</i> , <b>2017</b> , 23, e38-e45	12.9	245
97	The current consensus on the clinical management of intracranial ependymoma and its distinct molecular variants. <i>Acta Neuropathologica</i> , <b>2017</b> , 133, 5-12	14.3	202
96	Spectrum and prevalence of genetic predisposition in medulloblastoma: a retrospective genetic study and prospective validation in a clinical trial cohort. <i>Lancet Oncology, The</i> , <b>2018</b> , 19, 785-798	21.7	159
95	Lysine-specific demethylase 1 restricts hematopoietic progenitor proliferation and is essential for terminal differentiation. <i>Leukemia</i> , <b>2012</b> , 26, 2039-51	10.7	141
94	Childhood cancer predisposition syndromes-A concise review and recommendations by the Cancer Predisposition Working Group of the Society for Pediatric Oncology and Hematology. <i>American Journal of Medical Genetics, Part A</i> , <b>2017</b> , 173, 1017-1037	2.5	124
93	Therapeutic targeting of ependymoma as informed by oncogenic enhancer profiling. <i>Nature</i> , <b>2018</b> , 553, 101-105	50.4	116
92	Therapeutic Impact of Cytoreductive Surgery and Irradiation of Posterior Fossa Ependymoma in the Molecular Era: A Retrospective Multicohort Analysis. <i>Journal of Clinical Oncology</i> , <b>2016</b> , 34, 2468-77	2.2	113
91	Molecular heterogeneity and CXorf67 alterations in posterior fossa group A (PFA) ependymomas. <i>Acta Neuropathologica</i> , <b>2018</b> , 136, 211-226	14.3	111
90	EPEN-04. CXorf67 MIMICS ONCOGENIC HISTONE H3 K27M MUTATIONS AND FUNCTIONS AS INTRINSIC INHIBITOR OF PRC2 FUNCTION IN AGGRESSIVE POSTERIOR FOSSA EPENDYMOMA. <i>Neuro-Oncology</i> , <b>2019</b> , 21, ii78-ii78	1	78
89	EPEN-36. THE TREATMENT OUTCOME OF PAEDIATRIC SUPRATENTORIAL C11ORF95-RELA FUSED EPENDYMOMA: A COMBINED REPORT FROM E-HIT SERIES AND AUSTRALIAN NEW ZEALAND CHILDRENB HAEMATOLOGY/ONCOLOGY GROUP. <i>Neuro-Oncology</i> , <b>2020</b> , 22, iii315-iii315	1	78
88	EPEN-18. CROSS-SPECIES GENOMICS IDENTIFIES GLI2 AS AN ONCOGENE OF C11orf95 FUSION-POSITIVE SUPRATENTORIAL EPENDYMOMA. <i>Neuro-Oncology</i> , <b>2020</b> , 22, iii311-iii311	1	78
87	EPEN-44. EXTRACELLULAR VESICLES OF SUPRATENTORIAL EPENDYMOMA RELA MEDIATE INTERACTIONS WITH CELLS OF THE TUMOR MICROENVIRONMENT. <i>Neuro-Oncology</i> , <b>2020</b> , 22, iii316-iii	i3 <sup>1</sup> 17	78
86	MBRS-68. SINGLE NUCLEUS RNA-SEQUENCING DECIPHERS INTRATUMORAL HETEROGENEITY IN MEDULLOBLASTOMA WITH EXTENSIVE NODULARITY (MBEN). <i>Neuro-Oncology</i> , <b>2020</b> , 22, iii410-iii410	1	78

85	EPEN-03. ZFTA/C11ORF95 FUSIONS DRIVE SUPRATENTORIAL EPENDYMOMA VIA SHARED ONCOGENIC MECHANISMS. <i>Neuro-Oncology</i> , <b>2021</b> , 23, i13-i14	1	78
84	EPEN-07. OVEREXPRESSION AND MUTATIONS OF CXORF67 IN INFANT-TYPEIPOSTERIOR FOSSA TYPE-A (PFA) EPENDYMOMAS. <i>Neuro-Oncology</i> , <b>2018</b> , 20, i74-i74	1	78
83	Recommendations for Cancer Surveillance in Individuals with RASopathies and Other Rare Genetic Conditions with Increased Cancer Risk. <i>Clinical Cancer Research</i> , <b>2017</b> , 23, e83-e90	12.9	77
82	Sarcoma classification by DNA methylation profiling. <i>Nature Communications</i> , <b>2021</b> , 12, 498	17.4	74
81	EZHIP/CXorf67 mimics K27M mutated oncohistones and functions as an intrinsic inhibitor of PRC2 function in aggressive posterior fossa ependymoma. <i>Neuro-Oncology</i> , <b>2019</b> , 21, 878-889	1	65
8o	Heterogeneity within the PF-EPN-B ependymoma subgroup. <i>Acta Neuropathologica</i> , <b>2018</b> , 136, 227-237	14.3	52
79	cIMPACT-NOW update 7: advancing the molecular classification of ependymal tumors. <i>Brain Pathology</i> , <b>2020</b> , 30, 863-866	6	51
78	MYCN amplification drives an aggressive form of spinal ependymoma. <i>Acta Neuropathologica</i> , <b>2019</b> , 138, 1075-1089	14.3	51
77	Germline Elongator mutations in Sonic Hedgehog medulloblastoma. <i>Nature</i> , <b>2020</b> , 580, 396-401	50.4	47
76	Pharmacological activation of the p53 pathway by nutlin-3 exerts anti-tumoral effects in medulloblastomas. <i>Neuro-Oncology</i> , <b>2012</b> , 14, 859-69	1	44
75	Molecular mechanisms and therapeutic targets in pediatric brain tumors. <i>Science Signaling</i> , <b>2017</b> , 10,	8.8	43
74	Multiple Endocrine Neoplasia and Hyperparathyroid-Jaw Tumor Syndromes: Clinical Features, Genetics, and Surveillance Recommendations in Childhood. <i>Clinical Cancer Research</i> , <b>2017</b> , 23, e123-e13	<sup>12.9</sup>	43
73	Single-Cell RNA-Seq Reveals Cellular Hierarchies and Impaired Developmental Trajectories in Pediatric Ependymoma. <i>Cancer Cell</i> , <b>2020</b> , 38, 44-59.e9	24.3	40
72	YAP1 subgroup supratentorial ependymoma requires TEAD and nuclear factor I-mediated transcriptional programmes for tumorigenesis. <i>Nature Communications</i> , <b>2019</b> , 10, 3914	17.4	39
71	Telomere dysfunction and chromothripsis. <i>International Journal of Cancer</i> , <b>2016</b> , 138, 2905-14	7.5	34
70	DNA methylation-based classification of ependymomas in adulthood: implications for diagnosis and treatment. <i>Neuro-Oncology</i> , <b>2018</b> , 20, 1616-1624	1	32
69	MiR-34a deficiency accelerates medulloblastoma formation in vivo. <i>International Journal of Cancer</i> , <b>2015</b> , 136, 2293-303	7.5	32
68	EANO-EURACAN clinical practice guideline for diagnosis, treatment, and follow-up of post-pubertal and adult patients with medulloblastoma. <i>Lancet Oncology, The</i> , <b>2019</b> , 20, e715-e728	21.7	31

67	Diagnostics of pediatric supratentorial RELA ependymomas: integration of information from histopathology, genetics, DNA methylation and imaging. <i>Brain Pathology</i> , <b>2019</b> , 29, 325-335	6	30
66	FGFR1:TACC1 fusion is a frequent event in molecularly defined extraventricular neurocytoma. <i>Acta Neuropathologica</i> , <b>2018</b> , 136, 293-302	14.3	29
65	Brainstem biopsy in pediatric diffuse intrinsic pontine glioma in the era of precision medicine: the INFORM study experience. <i>European Journal of Cancer</i> , <b>2019</b> , 114, 27-35	7.5	28
64	Evaluation of Storage Tubes for Combined Analysis of Circulating Nucleic Acids in Liquid Biopsies. <i>International Journal of Molecular Sciences</i> , <b>2019</b> , 20,	6.3	27
63	Ependymoma. Seminars in Neurology, 2018, 38, 104-111	3.2	27
62	Papillary Tumor of the Pineal Region: A Distinct Molecular Entity. <i>Brain Pathology</i> , <b>2016</b> , 26, 199-205	6	25
61	Telomerase activation in posterior fossa group A ependymomas is associated with dismal prognosis and chromosome 1q gain. <i>Neuro-Oncology</i> , <b>2017</b> , 19, 1183-1194	1	24
60	The GSK461364 PLK1 inhibitor exhibits strong antitumoral activity in preclinical neuroblastoma models. <i>Oncotarget</i> , <b>2017</b> , 8, 6730-6741	3.3	24
59	YAP1-fusions in pediatric NF2-wildtype meningioma. Acta Neuropathologica, <b>2020</b> , 139, 215-218	14.3	24
58	The KDM1A histone demethylase is a promising new target for the epigenetic therapy of medulloblastoma. <i>Acta Neuropathologica Communications</i> , <b>2013</b> , 1, 19	7.3	23
57	Low-dose Actinomycin-D treatment re-establishes the tumoursuppressive function of P53 in RELA-positive ependymoma. <i>Oncotarget</i> , <b>2016</b> , 7, 61860-61873	3.3	22
56	The Pediatric Precision Oncology INFORM Registry: Clinical Outcome and Benefit for Patients with Very High-Evidence Targets. <i>Cancer Discovery</i> , <b>2021</b> , 11, 2764-2779	24.4	22
55	Comparison of tumor-associated YAP1 fusions identifies a recurrent set of functions critical for oncogenesis. <i>Genes and Development</i> , <b>2020</b> , 34, 1051-1064	12.6	21
54	Molecular characterization of histopathological ependymoma variants. <i>Acta Neuropathologica</i> , <b>2020</b> , 139, 305-318	14.3	20
53	Response to trametinib treatment in progressive pediatric low-grade glioma patients. <i>Journal of Neuro-Oncology</i> , <b>2020</b> , 149, 499-510	4.8	20
52	Cross-Species Genomics Reveals Oncogenic Dependencies in ZFTA/C11orf95 Fusion-Positive Supratentorial Ependymomas. <i>Cancer Discovery</i> , <b>2021</b> , 11, 2230-2247	24.4	20
51	Neuroblastoma in dialog with its stroma: NTRK1 is a regulator of cellular cross-talk with Schwann cells. <i>Oncotarget</i> , <b>2014</b> , 5, 11180-92	3.3	19
50	Epidemiology, molecular classification and WHO grading of ependymoma. <i>Journal of Neurosurgical Sciences</i> , <b>2018</b> , 62, 46-50	1.3	18

49	Expression of NTRK1/TrkA affects immunogenicity of neuroblastoma cells. <i>International Journal of Cancer</i> , <b>2013</b> , 133, 908-19	7.5	16
48	ZFTA-RELA Dictates Oncogenic Transcriptional Programs to Drive Aggressive Supratentorial Ependymoma. <i>Cancer Discovery</i> , <b>2021</b> , 11, 2200-2215	24.4	16
47	Predisposition to cancer in children and adolescents. <i>The Lancet Child and Adolescent Health</i> , <b>2021</b> , 5, 142-154	14.5	15
46	Molecular dissection of ependymomas. <i>Oncoscience</i> , <b>2015</b> , 2, 827-8	0.8	14
45	Ultra high-risk PFA ependymoma is characterized by loss of chromosome 6q. <i>Neuro-Oncology</i> , <b>2021</b> , 23, 1360-1370	1	14
44	Serial assessment of measurable residual disease in medulloblastoma liquid biopsies. <i>Cancer Cell</i> , <b>2021</b> , 39, 1519-1530.e4	24.3	13
43	Translocations Constitute Ependymoma Chromatin Remodeling and Transcription Factors. <i>Cancer Discovery</i> , <b>2021</b> , 11, 2216-2229	24.4	13
42	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> , <b>2020</b> , 20, 523	4.8	11
41	Intraventricular etoposide safety and toxicity profile in children and young adults with refractory or recurrent malignant brain tumors. <i>Journal of Neuro-Oncology</i> , <b>2016</b> , 128, 463-71	4.8	11
40	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> , <b>2021</b> , 23, 1012-1023	1	10
39	The pediatric precision oncology study INFORM: Clinical outcome and benefit for molecular subgroups <i>Journal of Clinical Oncology</i> , <b>2020</b> , 38, LBA10503-LBA10503	2.2	9
38	Genetic confirmation that ependymoma can arise as part of multiple endocrine neoplasia type 1 (MEN1) syndrome. <i>Acta Neuropathologica</i> , <b>2017</b> , 133, 661-663	14.3	8
37	Cerebrospinal Fluid Penetration and Combination Therapy of Entrectinib for Disseminated -Fusion Positive Pediatric High-Grade Glioma. <i>Journal of Personalized Medicine</i> , <b>2020</b> , 10,	3.6	8
36	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> , <b>2021</b> , 23, 848-857	1	7
35	PATZ1 fusions define a novel molecularly distinct neuroepithelial tumor entity with a broad histological spectrum. <i>Acta Neuropathologica</i> , <b>2021</b> , 142, 841-857	14.3	7
34	Limitations of current models for testing the clinical potential of epigenetic inhibitors for treatment of pediatric ependymoma. <i>Oncotarget</i> , <b>2018</b> , 9, 36530-36541	3.3	6
33	Transcriptional profiling of medulloblastoma with extensive nodularity (MBEN) reveals two clinically relevant tumor subsets with VSNL1 as potent prognostic marker. <i>Acta Neuropathologica</i> , <b>2020</b> , 139, 583-596	14.3	6
32	Newly Diagnosed Metastatic Intracranial Ependymoma in Children: Frequency, Molecular Characteristics, Treatment, and Outcome in the Prospective HIT Series. <i>Oncologist</i> , <b>2019</b> , 24, e921-e929	5.7	6

31	Current recommendations for clinical surveillance and genetic testing in rhabdoid tumor predisposition: a report from the SIOPE Host Genome Working Group. <i>Familial Cancer</i> , <b>2021</b> , 20, 305-31	$\vec{6}$	5
30	Recurrent fusions in PLAGL1 define a distinct subset of pediatric-type supratentorial neuroepithelial tumors. <i>Acta Neuropathologica</i> , <b>2021</b> , 142, 827-839	14.3	5
29	Targeting fibroblast growth factor receptors to combat aggressive ependymoma. <i>Acta Neuropathologica</i> , <b>2021</b> , 142, 339-360	14.3	4
28	From Sampling to Sequencing: A Liquid Biopsy Pre-Analytic Workflow to Maximize Multi-Layer Genomic Information from a Single Tube. <i>Cancers</i> , <b>2021</b> , 13,	6.6	3
27	The genetic landscape of choroid plexus tumors in children and adults. <i>Neuro-Oncology</i> , <b>2021</b> , 23, 650-6	60	3
26	How we treat medulloblastoma in adults. <i>ESMO Open</i> , <b>2021</b> , 6, 100173	6	3
25	Systemic chemotherapy of pediatric recurrent ependymomas: results from the German HIT-REZ studies. <i>Journal of Neuro-Oncology</i> , <b>2021</b> , 155, 193-202	4.8	2
24	Development of Randomized Trials in Adults with Medulloblastoma-The Example of EORTC 1634-BTG/NOA-23. <i>Cancers</i> , <b>2021</b> , 13,	6.6	2
23	Controversies and challenges in the management of paediatric non-rhabdomyosarcoma soft tissue sarcomas <i>The Lancet Child and Adolescent Health</i> , <b>2022</b> ,	14.5	2
22	SIOP Ependymoma I: Final results, long term follow-up and molecular analysis of the trial cohort: A BIOMECA Consortium Study <i>Neuro-Oncology</i> , <b>2022</b> ,	1	1
21	Clinical and molecular subgroups of ependymoma in adulthood: An analysis of the German Glioma Network <i>Journal of Clinical Oncology</i> , <b>2017</b> , 35, 2038-2038	2.2	1
20	Cancer predisposition in pediatric neuro-oncology-practical approaches and ethical considerations. <i>Neuro-Oncology Practice</i> , <b>2021</b> , 8, 526-538	2.2	1
19	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> , <b>2021</b> , 1176, 122781	3.2	1
18	Ependymoma <b>2018</b> , 177-192		1
17	The treatment approach to pediatric non-rhabdomyosarcoma soft tissue sarcomas: a critical review from the INternational Soft Tissue SaRcoma ConsorTium <i>European Journal of Cancer</i> , <b>2022</b> , 169, 10-19	7.5	1
16	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	O
15	Clinically aggressive pediatric spinal ependymoma with novel MYC amplification demonstrates molecular and histopathologic similarity to newly described MYCN-amplified spinal ependymomas <i>Acta Neuropathologica Communications</i> , <b>2021</b> , 9, 192	7.3	O
14	Liquid biopsieslals neue Diagnostikplattform in der pfliatrischen Onkologie. <i>Onkologe</i> , <b>2021</b> , 27, 458-46	30.1	

## LIST OF PUBLICATIONS

13	A Mouse Ependymoma Model Provides Molecular Insights into Tumor Formation. <i>Cell Reports</i> , <b>2018</b> , 23, 3699-3700	10.6
12	EPEN-18. Oncogenic 3D genome conformations identify novel therapeutic targets in ependymoma. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i42-i42	1
11	EPEN-19. Impact of molecular classification on prognosis in children and adolescents with spinal ependymoma: Results from the HIT-MED database. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i42-i43	1
10	MODL-04. Drug screening in Disorders with Abnormal DNA Damage Response/Repair (DADDR) andin vivo validation. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i168-i169	1
9	MEDB-60. Medulloblastoma with extensive nodularity mimics cerebellar development and differentiates along the granular precursor lineage. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i120-i120	1
8	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> , <b>2022</b> , 24, i113-i114	1
7	MEDB-14. Clinical outcome of pediatric medulloblastoma patients with Li-Fraumeni syndrome. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i107-i107	1
6	OTHR-32. The Pediatric Targeted Therapy 2.0 registry: robust molecular diagnostics for precision oncology. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i154-i154	1
5	MODL-07. DNA methylation-based biobank of murine models for pediatric tumors. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i169-i170	1
4	EPEN-28. Oncogenic dependency of pediatric ependymomas on extracellular vesicle pathways. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i45-i45	1
3	HGG-61.Landscape of cancer predisposition in pediatric high-grade glioma. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i76-i76	1
2	EPEN-09. Multi-omics characterization of the blood-brain barrier in molecular groups of ependymoma. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i40-i40	1
1	PATH-11. Detection of genetic and epigenetic alterations in Liquid Biopsies from pediatric brain tumor patients. <i>Neuro-Oncology</i> , <b>2022</b> , 24, i160-i161	1