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List of Publications by Year in descending order

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66
papers

3,863
citations

147801

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128289

60
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docs citations

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times ranked

4524
citing authors

#	ARTICLE	IF	CITATIONS
1	A Phase 2 Trial of Response-Based Radiation Therapy for Localized Central Nervous System Germ Cell Tumors: Patterns of Failure and Radiation Dosimetry for Nongerminomatous Germ Cell Tumors. <i>International Journal of Radiation Oncology Biology Physics</i> , 2022, 113, 143-151.	0.8	7
2	ADC Histogram Analysis of Pediatric Low-Grade Glioma Treated with Selumetinib: A Report from the Pediatric Brain Tumor Consortium. <i>American Journal of Neuroradiology</i> , 2022, 43, 455-461.	2.4	3
3	Pattern of treatment failures in patients with central nervous system non-germinomatous germ cell tumors (CNS-NGGCT): A pooled analysis of clinical trials. <i>Neuro-Oncology</i> , 2022, 24, 1950-1961.	1.2	12
4	Comprehensive Genomic Profiling of High-Risk Pediatric Cancer Patients Has a Measurable Impact on Clinical Care. <i>JCO Precision Oncology</i> , 2022, 6, e2100451.	3.0	3
5	Imaging response assessment for CNS germ cell tumours: consensus recommendations from the European Society for Paediatric Oncology Brain Tumour Group and North American Children's Oncology Group. <i>Lancet Oncology</i> , The, 2022, 23, e218-e228.	10.7	4
6	NFB-04. Evaluating focal areas of signal intensity (FASI) in children with neurofibromatosis type-1 (NF1) treated with selumetinib on PBTC-029B. <i>Neuro-Oncology</i> , 2022, 24, i128-i129.	1.2	0
7	MEK inhibitors for neurofibromatosis type 1 manifestations: Clinical evidence and consensus. <i>Neuro-Oncology</i> , 2022, 24, 1845-1856.	1.2	30
8	Phase 1 study of pomalidomide in children with recurrent, refractory, and progressive central nervous system tumors: A Pediatric Brain Tumor Consortium trial. <i>Pediatric Blood and Cancer</i> , 2021, 68, e28756.	1.5	9
9	A phase II trial of selumetinib in children with recurrent optic pathway and hypothalamic low-grade glioma without NF1: a Pediatric Brain Tumor Consortium study. <i>Neuro-Oncology</i> , 2021, 23, 1777-1788.	1.2	68
10	Neonatal Central Nervous System Tumors. <i>Clinics in Perinatology</i> , 2021, 48, 35-51.	2.1	3
11	Phase 2 Study of Pomalidomide (CC-4047) Monotherapy for Children and Young Adults With Recurrent or Progressive Primary Brain Tumors. <i>Frontiers in Oncology</i> , 2021, 11, 660892.	2.8	7
12	Clinical phenotypes and prognostic features of embryonal tumours with multi-layered rosettes: a Rare Brain Tumor Registry study. <i>The Lancet Child and Adolescent Health</i> , 2021, 5, 800-813.	5.6	12
13	Multicenter Analysis of Genomically Targeted Single Patient Use Requests for Pediatric Neoplasms. <i>Journal of Clinical Oncology</i> , 2021, 39, 3822-3828.	1.6	4
14	Clinical Pharmacokinetics and Pharmacodynamics of Selumetinib. <i>Clinical Pharmacokinetics</i> , 2021, 60, 283-303.	3.5	16
15	Advances in the classification and treatment of pediatric brain tumors. <i>Current Opinion in Pediatrics</i> , 2021, 33, 26-32.	2.0	11
16	Pineoblastoma segregates into molecular sub-groups with distinct clinico-pathologic features: a Rare Brain Tumor Consortium registry study. <i>Acta Neuropathologica</i> , 2020, 139, 223-241.	7.7	65
17	Response assessment in paediatric low-grade glioma: recommendations from the Response Assessment in Pediatric Neuro-Oncology (RAPNO) working group. <i>Lancet Oncology</i> , The, 2020, 21, e305-e316.	10.7	115
18	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.	1.2	44

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19	The “Risk” in Pediatric Low-Grade Glioma. <i>Cancer Cell</i> , 2020, 37, 424-425.	16.8	5
20	Advanced ADC Histogram, Perfusion, and Permeability Metrics Show an Association with Survival and Pseudoprogression in Newly Diagnosed Diffuse Intrinsic Pontine Glioma: A Report from the Pediatric Brain Tumor Consortium. <i>American Journal of Neuroradiology</i> , 2020, 41, 718-724.	2.4	14
21	Phase II Trial of Response-Based Radiation Therapy for Patients With Localized CNS Nongerminomatous Germ Cell Tumors: A Children's Oncology Group Study. <i>Journal of Clinical Oncology</i> , 2019, 37, 3283-3290.	1.6	78
22	Transcriptional repressor REST drives lineage stage-specific chromatin compaction at <i>Ptch1</i> and increases AKT activation in a mouse model of medulloblastoma. <i>Science Signaling</i> , 2019, 12, .	3.6	19
23	Selumetinib in paediatric patients with BRAF-aberrant or neurofibromatosis type 1-associated recurrent, refractory, or progressive low-grade glioma: a multicentre, phase 2 trial. <i>Lancet Oncology</i> , 2019, 20, 1011-1022.	10.7	315
24	Improved neuropsychological outcomes following proton therapy relative to X-ray therapy for pediatric brain tumor patients. <i>Neuro-Oncology</i> , 2019, 21, 934-943.	1.2	51
25	Management of pediatric low-grade glioma. <i>Current Opinion in Pediatrics</i> , 2019, 31, 21-27.	2.0	87
26	Prospective feasibility and safety assessment of surgical biopsy for patients with newly diagnosed diffuse intrinsic pontine glioma. <i>Neuro-Oncology</i> , 2018, 20, 1547-1555.	1.2	82
27	Pediatric low-grade gliomas: next biologically driven steps. <i>Neuro-Oncology</i> , 2018, 20, 160-173.	1.2	116
28	REST upregulates gremlin to modulate diffuse intrinsic pontine glioma vasculature. <i>Oncotarget</i> , 2018, 9, 5233-5250.	1.8	12
29	Evaluation of age-dependent treatment strategies for children and young adults with pineoblastoma: analysis of pooled European Society for Paediatric Oncology (SIOP-E) and US Head Start data. <i>Neuro-Oncology</i> , 2017, 19, now234.	1.2	33
30	Pediatric low-grade gliomas: implications of the biologic era. <i>Neuro-Oncology</i> , 2017, 19, now209.	1.2	73
31	Phase II trial of pegylated interferon alfa-2b in young patients with neurofibromatosis type 1 and unresectable plexiform neurofibromas. <i>Neuro-Oncology</i> , 2017, 19, now158.	1.2	41
32	A phase I trial of the MEK inhibitor selumetinib (AZD6244) in pediatric patients with recurrent or refractory low-grade glioma: a Pediatric Brain Tumor Consortium (PBTC) study. <i>Neuro-Oncology</i> , 2017, 19, 1135-1144.	1.2	236
33	Rethinking childhood ependymoma: a retrospective, multi-center analysis reveals poor long-term overall survival. <i>Journal of Neuro-Oncology</i> , 2017, 135, 201-211.	2.9	72
34	Severe Radiation Necrosis Successfully Treated With Bevacizumab in an Infant with Low-Grade Glioma and Tumor-Associated Intractable Trigeminal Neuralgia. <i>Pediatric Blood and Cancer</i> , 2016, 63, 1671-1673.	1.5	13
35	Improving vaccine efficacy against malignant glioma. <i>Oncolmmunology</i> , 2016, 5, e1196311.	4.6	16
36	Involvement of the neural stem cell compartment by pediatric and adult gliomas: a retrospective review of 377 cases. <i>Journal of Neuro-Oncology</i> , 2015, 122, 105-110.	2.9	2

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37	Consensus on the management of intracranial germ-cell tumours. <i>Lancet Oncology, The</i> , 2015, 16, e470-e477.	10.7	173
38	Successful treatment of metastatic ^{125}I HCG-secreting germ cell tumor occurring 3 years after total resection of a pineal mature teratoma. <i>European Journal of Pediatrics</i> , 2014, 173, 1011-5.	2.7	5
39	Efficacy of bevacizumab plus irinotecan in children with recurrent low-grade gliomas—a Pediatric Brain Tumor Consortium study. <i>Neuro-Oncology</i> , 2014, 16, 310-317.	1.2	132
40	CNS-PNETs with C19MC amplification and/or LIN28 expression comprise a distinct histogenetic diagnostic and therapeutic entity. <i>Acta Neuropathologica</i> , 2014, 128, 291-303.	7.7	141
41	Unclear standard of care for pediatric high grade glioma patients. <i>Journal of Neuro-Oncology</i> , 2013, 113, 341-342.	2.9	22
42	Response to bevacizumab, irinotecan, and temozolomide in children with relapsed medulloblastoma: a multi-institutional experience. <i>Child's Nervous System</i> , 2013, 29, 589-596.	1.1	66
43	Bevacizumab-Associated Osteonecrosis of the Wrist and Knee in Three Pediatric Patients With Recurrent CNS Tumors. <i>Journal of Clinical Oncology</i> , 2013, 31, e24-e27.	1.6	18
44	Bevacizumab (BVZ)-associated toxicities in children with recurrent central nervous system tumors treated with BVZ and irinotecan (CPT-11). <i>Cancer</i> , 2013, 119, 4180-4187.	4.1	33
45	Lack of efficacy of bevacizumab + irinotecan in cases of pediatric recurrent ependymoma—a Pediatric Brain Tumor Consortium study. <i>Neuro-Oncology</i> , 2012, 14, 1404-1412.	1.2	50
46	REST Is a Novel Prognostic Factor and Therapeutic Target for Medulloblastoma. <i>Molecular Cancer Therapeutics</i> , 2012, 11, 1713-1723.	4.1	47
47	Pediatric High Grade Glioma: a Review and Update on Tumor Clinical Characteristics and Biology. <i>Frontiers in Oncology</i> , 2012, 2, 105.	2.8	137
48	Markers of survival and metastatic potential in childhood CNS primitive neuro-ectodermal brain tumours: an integrative genomic analysis. <i>Lancet Oncology, The</i> , 2012, 13, 838-848.	10.7	148
49	Variable response to propranolol treatment of kaposiform hemangioendothelioma, tufted angioma, and Kasabach-Merritt phenomenon. <i>Pediatric Blood and Cancer</i> , 2012, 59, 934-938.	1.5	107
50	Evaluating the incidence and utility of microscopic metastatic dissemination as diagnosed by lumbar cerebro-spinal fluid (CSF) samples in children with newly diagnosed intracranial ependymoma. <i>Journal of Neuro-Oncology</i> , 2011, 103, 693-698.	2.9	24
51	Recurrent pure CNS germinoma with markedly elevated serum and cerebrospinal fluid human chorionic gonadotropin- β (HCG β). <i>Pediatric Blood and Cancer</i> , 2011, 56, 863-864.	1.5	3
52	Bevacizumab and irinotecan in the treatment of children with recurrent/refractory medulloblastoma. <i>Pediatric Blood and Cancer</i> , 2011, 56, 491-494.	1.5	27
53	Long-term survival in a pediatric patient with supratentorial primitive neuroectodermal tumor and extraneural metastasis at diagnosis. <i>Pediatric Blood and Cancer</i> , 2011, 57, 341-344.	1.5	2
54	Phase I Trial of Lenalidomide in Pediatric Patients With Recurrent, Refractory, or Progressive Primary CNS Tumors: Pediatric Brain Tumor Consortium Study PBTC-018. <i>Journal of Clinical Oncology</i> , 2011, 29, 324-329.	1.6	83

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55	Non-cerebellar primitive neuroectodermal tumors (PNET): Summary of the Milan consensus and state of the art workshop on marrow ablative chemotherapy with hematopoietic cell rescue for malignant brain tumors of childhood and adolescents. <i>Pediatric Blood and Cancer</i> , 2010, 54, 638-640.	1.5	14
56	Incidental Resolution of a Radiation-Induced Cavernous Hemangioma of the Brain following the Use of Bevacizumab in a Child with Recurrent Medulloblastoma. <i>Pediatric Neurosurgery</i> , 2010, 46, 303-307.	0.7	19
57	Aggressive variant of a papillary glioneuronal tumor. <i>Journal of Neurosurgery: Pediatrics</i> , 2009, 3, 46-52.	1.3	43
58	Introduction to a Special Issue on Pediatric Neuro-Oncology. <i>Journal of Child Neurology</i> , 2009, 24, 1341-1342.	1.4	5
59	Pediatric High-Grade Gliomas and Diffuse Intrinsic Pontine Gliomas. <i>Journal of Child Neurology</i> , 2009, 24, 1409-1417.	1.4	84
60	Glioblastoma Multiforme in a Patient With Chronic Granulomatous Disease Treated With Subtotal Resection, Radiation, and Thalidomide. <i>Journal of Pediatric Hematology/Oncology</i> , 2009, 31, 965-969.	0.6	5
61	Acute transient encephalopathy following paclitaxel treatment in an adolescent with a recurrent suprasellar germinoma. <i>Pediatric Blood and Cancer</i> , 2008, 50, 699-700.	1.5	9
62	Intensive chemotherapy followed by consolidative myeloablative chemotherapy with autologous hematopoietic cell rescue (AuHCR) in young children with newly diagnosed supratentorial primitive neuroectodermal tumors (sPNETs): Report of the Head Start I and II experience. <i>Pediatric Blood and Cancer</i> , 2008, 50, 312-318.	1.5	125
63	Radiation therapy approaches do not currently improve overall survival in young children with sPNET as compared to the Head Start I and II experience. <i>Pediatric Blood and Cancer</i> , 2008, 51, 149-150.	1.5	1
64	Pediatric Central Nervous System Germ Cell Tumors: A Review. <i>Oncologist</i> , 2008, 13, 690-699.	3.7	370
65	Survivin splice variants regulate the balance between proliferation and cell death. <i>Oncogene</i> , 2005, 24, 1994-2007.	5.9	176
66	Essential Role for Survivin in Early Brain Development. <i>Journal of Neuroscience</i> , 2005, 25, 6962-6970.	3.6	116