

Jacques Grill

List of Publications by Year in descending order

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191
papers

12,556
citations

28274

55
h-index

28297

105
g-index

195
all docs

195
docs citations

195
times ranked

11785
citing authors

#	ARTICLE	IF	CITATIONS
1	Repurposing Vandetanib plus Everolimus for the Treatment of <i>ACVR1</i> -Mutant Diffuse Intrinsic Pontine Glioma. <i>Cancer Discovery</i> , 2022, 12, 416-431.	9.4	25
2	DIPG Harbors Alterations Targetable by MEK Inhibitors, with Acquired Resistance Mechanisms Overcome by Combinatorial Inhibition. <i>Cancer Discovery</i> , 2022, 12, 712-729.	9.4	15
3	Deciphering the genetic and epigenetic landscape of pediatric bithalamic tumors. <i>Brain Pathology</i> , 2022, 32, e13039.	4.1	5
4	High Prevalence of Early Endocrine Disorders After Childhood Brain Tumors in a Large Cohort. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e2156-e2166.	3.6	6
5	MRI and Molecular Characterization of Pediatric High-Grade Midline Thalamic Gliomas: The HERBY Phase II Trial. <i>Radiology</i> , 2022, 304, 174-182.	7.3	12
6	HGG-41. Glioma oncogenesis in the constitutional mismatch repair deficiency (CMMRD) syndrome. <i>Neuro-Oncology</i> , 2022, 24, i70-i70.	1.2	0
7	EPEN-24. Biological markers of ependymoma in children and adolescents (BIOMECA): Systematic comparison of methods for the precise evaluation of biomarkers for ependymoma diagnosis and prognostication. <i>Neuro-Oncology</i> , 2022, 24, i44-i44.	1.2	0
8	HGG-49. Gliomatosis cerebri in children: A collaborative report from the European Society for Pediatric Oncology (SIOPE). <i>Neuro-Oncology</i> , 2022, 24, i72-i73.	1.2	0
9	HGG-40. NF1 mosaicism in a CMMRD-patient with a glioblastoma. <i>Neuro-Oncology</i> , 2022, 24, i69-i70.	1.2	0
10	NF1 optic pathway glioma: analyzing risk factors for visual outcome and indications to treat. <i>Neuro-Oncology</i> , 2021, 23, 100-111.	1.2	27
11	A subset of pediatric-type thalamic gliomas share a distinct DNA methylation profile, H3K27me3 loss and frequent alteration of <i>EGFR</i> . <i>Neuro-Oncology</i> , 2021, 23, 34-43.	1.2	75
12	Clinical and molecular analysis of smoothened inhibitors in Sonic Hedgehog medulloblastoma. <i>Neuro-Oncology Advances</i> , 2021, 3, vdab097.	0.7	5
13	Droplet digital PCR-based detection of circulating tumor DNA from pediatric high grade and diffuse midline glioma patients. <i>Neuro-Oncology Advances</i> , 2021, 3, vdab013.	0.7	27
14	Radiogenomics of diffuse intrinsic pontine gliomas (DIPGs): correlation of histological and biological characteristics with multimodal MRI features. <i>European Radiology</i> , 2021, 31, 8913-8924.	4.5	11
15	A novel case of cribriform neuroepithelial tumor: A potential diagnostic pitfall in the ventricular system. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29037.	1.5	3
16	A CBF decrease in the left supplementary motor areas: New insight into postoperative pediatric cerebellar mutism syndrome using arterial spin labeling perfusion MRI. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2021, 41, 3339-3349.	4.3	10
17	Supratentorial non-RELA, ZFTA-fused ependymomas: a comprehensive phenotype genotype correlation highlighting the number of zinc fingers in ZFTA-NCOA1/2 fusions. <i>Acta Neuropathologica Communications</i> , 2021, 9, 135.	5.2	21
18	Pediatric brain tumors as a developmental disease. <i>Current Opinion in Oncology</i> , 2021, 33, 608-614.	2.4	2

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19	CNS tumors with YWHAE:NUTM2 and KDM2B-fusions present molecular similarities to extra-CNS tumors having BCOR internal tandem duplication or alternative fusions. <i>Acta Neuropathologica Communications</i> , 2021, 9, 176.	5.2	1
20	Prognostic Clinical and Biologic Features for Overall Survival after Relapse in Childhood Medulloblastoma. <i>Cancers</i> , 2021, 13, 53.	3.7	10
21	Object Detection Improves Tumour Segmentation in MR Images of Rare Brain Tumours. <i>Cancers</i> , 2021, 13, 6113.	3.7	9
22	High Prevalence of Developmental Venous Anomaly in Diffuse Intrinsic Pontine Gliomas: A Pediatric Control Study. <i>Neurosurgery</i> , 2020, 86, 517-523.	1.1	13
23	WHO grade has no prognostic value in the pediatric high-grade glioma included in the HERBY trial. <i>Neuro-Oncology</i> , 2020, 22, 116-127.	1.2	26
24	The histomolecular criteria established for adult anaplastic pilocytic astrocytoma are not applicable to the pediatric population. <i>Acta Neuropathologica</i> , 2020, 139, 287-303.	7.7	19
25	Regarding "Neuro-Oncology Practice Clinical Debate: targeted therapy vs conventional chemotherapy in pediatric low-grade glioma". <i>Neuro-Oncology Practice</i> , 2020, 7, 572-573.	1.6	2
26	Modeling the Interaction between the Microenvironment and Tumor Cells in Brain Tumors. <i>Neuron</i> , 2020, 108, 1025-1044.	8.1	31
27	Phase I study of vinblastine in combination with nilotinib in children, adolescents, and young adults with refractory or recurrent low-grade glioma. <i>Neuro-Oncology Advances</i> , 2020, 2, vdaa075.	0.7	2
28	The EP300:BCOR fusion extends the genetic alteration spectrum defining the new tumoral entity of "CNS tumors with BCOR internal tandem duplication". <i>Acta Neuropathologica Communications</i> , 2020, 8, 178.	5.2	17
29	Focal Areas of High Signal Intensity in Children with Neurofibromatosis Type 1: Expected Evolution on MRI. <i>American Journal of Neuroradiology</i> , 2020, 41, 1733-1739.	2.4	8
30	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.	7.7	104
31	High-grade gliomas in adolescents and young adults highlight histomolecular differences from their adult and pediatric counterparts. <i>Neuro-Oncology</i> , 2020, 22, 1190-1202.	1.2	50
32	Ultrasound-induced blood-brain barrier disruption for the treatment of gliomas and other primary CNS tumors. <i>Cancer Letters</i> , 2020, 479, 13-22.	7.2	38
33	Radiological Evaluation of Newly Diagnosed Non-Brainstem Pediatric High-Grade Glioma in the HERBY Phase II Trial. <i>Clinical Cancer Research</i> , 2020, 26, 1856-1865.	7.0	10
34	Role of neoadjuvant chemotherapy in metastatic medulloblastoma: a comparative study in 92 children. <i>Neuro-Oncology</i> , 2020, 22, 1686-1695.	1.2	14
35	Blood-brain barrier disruption with low-intensity pulsed ultrasound for the treatment of pediatric brain tumors: a review and perspectives. <i>Neurosurgical Focus</i> , 2020, 48, E10.	2.3	31
36	A kinome-wide shRNA screen uncovers vaccinia-related kinase 3 (VRK3) as an essential gene for diffuse intrinsic pontine glioma survival. <i>Oncogene</i> , 2019, 38, 6479-6490.	5.9	13

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37	The dark matter of diffuse intrinsic pontine gliomas: an update. Expert Opinion on Orphan Drugs, 2019, 7, 11-20.	0.8	1
38	Diagnostics and treatment of diffuse intrinsic pontine glioma: where do we stand?. Journal of Neuro-Oncology, 2019, 145, 177-184.	2.9	36
39	Patients with High-Grade Gliomas and Café-au-Lait Macules: Is Neurofibromatosis Type 1 the Only Diagnosis?. American Journal of Neuroradiology, 2019, 40, E30-E31.	2.4	4
40	Evaluation of the Implementation of the Response Assessment in Neuro-Oncology Criteria in the HERBY Trial of Pediatric Patients with Newly Diagnosed High-Grade Gliomas. American Journal of Neuroradiology, 2019, 40, 568-575.	2.4	4
41	Anatomo-functional study of the cerebellum in working memory in children treated for medulloblastoma. Journal of Neuroradiology, 2019, 46, 207-213.	1.1	9
42	Constitutional mismatch repair deficiency-associated brain tumors: report from the European C4CMMRD consortium. Neuro-Oncology Advances, 2019, 1, vdz033.	0.7	23
43	TP53 Pathway Alterations Drive Radioresistance in Diffuse Intrinsic Pontine Gliomas (DIPG). Clinical Cancer Research, 2019, 25, 6788-6800.	7.0	66
44	International experience in the development of patient-derived xenograft models of diffuse intrinsic pontine glioma. Journal of Neuro-Oncology, 2019, 141, 253-263.	2.9	30
45	Maternal stress and pediatric brain cancer: A French study. Journal of Psychosocial Oncology, 2019, 37, 96-109.	1.2	7
46	Integrated analysis of long-term growth and bone development in pediatric and adolescent patients receiving bevacizumab. Pediatric Blood and Cancer, 2019, 66, e27487.	1.5	5
47	Diagnostics of pediatric supratentorial RELA ependymomas: integration of information from histopathology, genetics, DNA methylation and imaging. Brain Pathology, 2019, 29, 325-335.	4.1	55
48	Childhood supratentorial ependymomas with <i>YAP1</i> – <i>MAML2</i> fusion: an entity with characteristic clinical, radiological, cytogenetic and histopathological features. Brain Pathology, 2019, 29, 205-216.	4.1	75
49	CT and Multimodal MR Imaging Features of Embryonal Tumors with Multilayered Rosettes in Children. American Journal of Neuroradiology, 2019, 40, 732-736.	2.4	9
50	Germline <i>SUFU</i> mutation carriers and medulloblastoma: clinical characteristics, cancer risk, and prognosis. Neuro-Oncology, 2018, 20, 1122-1132.	1.2	52
51	Cerebral blood flow changes after radiation therapy identifies pseudoprogression in diffuse intrinsic pontine gliomas. Neuro-Oncology, 2018, 20, 994-1002.	1.2	21
52	Historadiological correlations in high-grade glioma with the histone 3.3 G34R mutation. Journal of Neuroradiology, 2018, 45, 316-322.	1.1	26
53	Co-occurrence of histone H3 K27M and BRAF V600E mutations in paediatric midline grade I ganglioglioma. Brain Pathology, 2018, 28, 103-111.	4.1	80
54	Clinical, Radiologic, Pathologic, and Molecular Characteristics of Long-Term Survivors of Diffuse Intrinsic Pontine Glioma (DIPG): A Collaborative Report From the International and European Society for Pediatric Oncology DIPG Registries. Journal of Clinical Oncology, 2018, 36, 1963-1972.	1.6	250

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55	Phase II, Open-Label, Randomized, Multicenter Trial (HERBY) of Bevacizumab in Pediatric Patients With Newly Diagnosed High-Grade Glioma. <i>Journal of Clinical Oncology</i> , 2018, 36, 951-958.	1.6	95
56	Transcriptomic and epigenetic profiling of diffuse midline gliomas, H3 K27M-mutant™ discriminate two subgroups based on the type of histone H3 mutated and not supratentorial or infratentorial location. <i>Acta Neuropathologica Communications</i> , 2018, 6, 117.	5.2	83
57	DIPG-20. PRE-RANDOMISATION CENTRAL REVIEW AND REAL-TIME BIOMARKERS SCREENING IN THE MULTICENTRE BIOLOGICAL MEDICINE FOR DIPG ERADICATION (BIOMEDE) TRIAL: LESSONS LEARNT FROM THE FIRST 120 BIOPSIES. <i>Neuro-Oncology</i> , 2018, 20, i52-i53.	1.2	2
58	Parental stress and paediatric acquired brain injury. <i>Brain Injury</i> , 2018, 32, 1780-1786.	1.2	10
59	Tumor dissemination through surgical tracts in diffuse intrinsic pontine glioma. <i>Journal of Neurosurgery: Pediatrics</i> , 2018, 22, 678-683.	1.3	9
60	New stratification for early childhood medulloblastoma. <i>Pediatric Medicine</i> , 2018, 1, 10-10.	2.7	2
61	Cognitive Profile of Children With Intracranial Germ Cell Tumor According to Tumor Location. <i>Journal of Pediatric Hematology/Oncology</i> , 2018, 40, e424-e428.	0.6	4
62	Molecular, Pathological, Radiological, and Immune Profiling of Non-brainstem Pediatric High-Grade Glioma from the HERBY Phase II Randomized Trial. <i>Cancer Cell</i> , 2018, 33, 829-842.e5.	16.8	140
63	Development of the SIOPE DIPG network, registry and imaging repository: a collaborative effort to optimize research into a rare and lethal disease. <i>Journal of Neuro-Oncology</i> , 2017, 132, 255-266.	2.9	42
64	The international diffuse intrinsic pontine glioma registry: an infrastructure to accelerate collaborative research for an orphan disease. <i>Journal of Neuro-Oncology</i> , 2017, 132, 323-331.	2.9	27
65	Diffuse intrinsic pontine gliomas—current management and new biologic insights. Is there a glimmer of hope?. <i>Neuro-Oncology</i> , 2017, 19, 1025-1034.	1.2	91
66	Integrated Molecular Meta-Analysis of 1,000 Pediatric High-Grade and Diffuse Intrinsic Pontine Glioma. <i>Cancer Cell</i> , 2017, 32, 520-537.e5.	16.8	716
67	Molecular Screening for Cancer Treatment Optimization (MOSCATO-01) in Pediatric Patients: A Single-Institutional Prospective Molecular Stratification Trial. <i>Clinical Cancer Research</i> , 2017, 23, 6101-6112.	7.0	102
68	Epileptic seizures in anaplastic gangliogliomas. <i>British Journal of Neurosurgery</i> , 2017, 31, 227-233.	0.8	6
69	Relationships between Regional Radiation Doses and Cognitive Decline in Children Treated with Cranio-Spinal Irradiation for Posterior Fossa Tumors. <i>Frontiers in Oncology</i> , 2017, 7, 166.	2.8	20
70	Integrating Tenascin-C protein expression and 1q25 copy number status in pediatric intracranial ependymoma prognostication: A new model for risk stratification. <i>PLoS ONE</i> , 2017, 12, e0178351.	2.5	15
71	Quality of survival and cognitive performance in children treated for medulloblastoma in the PNET 4 randomized controlled trial. <i>Neuro-Oncology Practice</i> , 2017, 4, 161-170.	1.6	9
72	New <i>in vivo</i> avatars of diffuse intrinsic pontine gliomas (DIPG) from stereotactic biopsies performed at diagnosis. <i>Oncotarget</i> , 2017, 8, 52543-52559.	1.8	41

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73	HG-46 RECURRENT DIFFUSE INTRINSIC PONTINE GLIOMAS: CLINICAL, BIOLOGICAL, RADIOLOGICAL AND THERAPEUTIC FACTORS CORRELATING WITH THE SURVIVAL. <i>Neuro-Oncology</i> , 2016, 18, iii57.4-iii58.	1.2	0
74	Primary Leptomeningeal Gliomatosis in Children and Adults. <i>Neurosurgery</i> , 2016, 78, 343-352.	1.1	10
75	Chordoma in children: Case-report and review of literature. <i>Reports of Practical Oncology and Radiotherapy</i> , 2016, 21, 1-7.	0.6	11
76	Response Assessment in Pediatric Neuro-Oncology: Implementation and Expansion of the RANO Criteria in a Randomized Phase II Trial of Pediatric Patients with Newly Diagnosed High-Grade Gliomas. <i>American Journal of Neuroradiology</i> , 2016, 37, 1581-1587.	2.4	31
77	Histone H3 genotyping refines clinico-radiological diagnostic and prognostic criteria in DIPG. <i>Acta Neuropathologica</i> , 2016, 131, 795-796.	7.7	11
78	Clinical, Imaging, Histopathological and Molecular Characterization of Anaplastic Ganglioglioma. <i>Journal of Neuropathology and Experimental Neurology</i> , 2016, 75, 971-980.	1.7	54
79	Re-irradiation of recurrent pediatric ependymoma: modalities and outcomes: a twenty-year survey. <i>SpringerPlus</i> , 2016, 5, 879.	1.2	35
80	Bevacizumab dosing strategy in paediatric cancer patients based on population pharmacokinetic analysis with external validation. <i>British Journal of Clinical Pharmacology</i> , 2016, 81, 148-160.	2.4	38
81	Arterial Spin Labeling to Predict Brain Tumor Grading in Children: Correlations between Histopathologic Vascular Density and Perfusion MR Imaging. <i>Radiology</i> , 2016, 281, 553-566.	7.3	82
82	Reprint of "Chordoma in children: Case-report and review of literature". <i>Reports of Practical Oncology and Radiotherapy</i> , 2016, 21, 412-417.	0.6	4
83	MYB-QKI rearrangements in angiocentric glioma drive tumorigenicity through a tripartite mechanism. <i>Nature Genetics</i> , 2016, 48, 273-282.	21.4	214
84	New Brain Tumor Entities Emerge from Molecular Classification of CNS-PNETs. <i>Cell</i> , 2016, 164, 1060-1072.	28.9	702
85	Palliative and end-of-life care for children with diffuse intrinsic pontine glioma: results from a London cohort study and international survey. <i>Neuro-Oncology</i> , 2016, 18, 582-588.	1.2	25
86	Long survival in a child with a mutated K27M-H3.3 pilocytic astrocytoma. <i>Annals of Clinical and Translational Neurology</i> , 2015, 2, 439-443.	3.7	51
87	Patients in Pediatric Phase I and Early Phase II Clinical Oncology Trials at Gustave Roussy. <i>Journal of Pediatric Hematology/Oncology</i> , 2015, 37, e102-e110.	0.6	31
88	Mortality in Children with Optic Pathway Glioma Treated with Up-Front BB-SFOP Chemotherapy. <i>PLoS ONE</i> , 2015, 10, e0127676.	2.5	38
89	Expression profiles of 151 pediatric low-grade gliomas reveal molecular differences associated with location and histological subtype. <i>Neuro-Oncology</i> , 2015, 17, 1486-1496.	1.2	39
90	A common polymorphism in the 5' UTR of ERCC5 creates an upstream ORF that confers resistance to platinum-based chemotherapy. <i>Genes and Development</i> , 2015, 29, 1891-1896.	5.9	32

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91	Preclinical evaluation of dasatinib alone and in combination with cabozantinib for the treatment of diffuse intrinsic pontine glioma. <i>Neuro-Oncology</i> , 2015, 17, 953-964.	1.2	56
92	Functionally defined therapeutic targets in diffuse intrinsic pontine glioma. <i>Nature Medicine</i> , 2015, 21, 555-559.	30.7	473
93	Inhibition of the NOTCH pathway using \hat{I}^3 -secretase inhibitor RO4929097 has limited antitumor activity in established glial tumors. <i>Anti-Cancer Drugs</i> , 2015, 26, 272-283.	1.4	26
94	Histone H3F3A and HIST1H3B K27M mutations define two subgroups of diffuse intrinsic pontine gliomas with different prognosis and phenotypes. <i>Acta Neuropathologica</i> , 2015, 130, 815-827.	7.7	482
95	Translating preclinical hopes into clinical reality for children with ependymoma. <i>Neuro-Oncology</i> , 2015, 17, 1545-1546.	1.2	1
96	Biopsy in a series of 130 pediatric diffuse intrinsic Pontine gliomas. <i>Child's Nervous System</i> , 2015, 31, 1773-1780.	1.1	145
97	Conceptions of time in children treated for malignant cerebellar tumours. <i>Brain Injury</i> , 2014, 28, 1334-1341.	1.2	3
98	Variable selection for generalized canonical correlation analysis. <i>Biostatistics</i> , 2014, 15, 569-583.	1.5	168
99	Water and Electrolyte Disorders at Long-Term Post-Treatment Follow-Up in Paediatric Patients with Suprasellar Tumours Include Unexpected Persistent Cerebral Salt-Wasting Syndrome. <i>Hormone Research in Paediatrics</i> , 2014, 82, 364-371.	1.8	20
100	Vemurafenib in pediatric patients with <i>BRAFV600E</i> mutated high-grade gliomas. <i>Pediatric Blood and Cancer</i> , 2014, 61, 1101-1103.	1.5	125
101	Central nervous system germ cell tumors. <i>Current Opinion in Oncology</i> , 2014, 26, 622-626.	2.4	26
102	Tandem high-dose chemotherapy and autologous stem cell rescue in children with newly diagnosed high-risk medulloblastoma or supratentorial primitive neuroectodermic tumors. <i>Pediatric Blood and Cancer</i> , 2014, 61, 1398-1402.	1.5	46
103	High-dose busulfan-thiotepa with autologous stem cell transplantation followed by posterior fossa irradiation in young children with classical or incompletely resected medulloblastoma. <i>Pediatric Blood and Cancer</i> , 2014, 61, 907-912.	1.5	18
104	Pediatric low-grade gliomas: How modern biology reshapes the clinical field. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2014, 1845, 294-307.	7.4	45
105	Stability of medulloblastoma subgroups at tumour recurrence. <i>Nature Reviews Neurology</i> , 2014, 10, 5-6.	10.1	4
106	Paediatric and adult glioblastoma: multiform (epi)genomic culprits emerge. <i>Nature Reviews Cancer</i> , 2014, 14, 92-107.	28.4	469
107	Rubinstein-Taybi syndrome predisposing to non-WNT, non-SHH, group 3 medulloblastoma. <i>Pediatric Blood and Cancer</i> , 2014, 61, 383-386.	1.5	33
108	Recurrent activating ACVR1 mutations in diffuse intrinsic pontine glioma. <i>Nature Genetics</i> , 2014, 46, 457-461.	21.4	423

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109	Stability of Etoposide Solutions in Disposable Infusion Devices for Day Hospital Cancer Practices. <i>Drugs in R and D</i> , 2014, 14, 13-23.	2.2	4
110	Disrupted sensorimotor synchronization, but intact rhythm discrimination, in children treated for a cerebellar medulloblastoma. <i>Research in Developmental Disabilities</i> , 2014, 35, 2053-2068.	2.2	14
111	A phase II single-arm study of irinotecan in combination with temozolomide (TEMIRI) in children with newly diagnosed high grade glioma: a joint ITCC and SIOPE-brain tumour study. <i>Journal of Neuro-Oncology</i> , 2013, 113, 127-134.	2.9	9
112	Primary gliomatosis cerebri involving gray matter in pediatrics: a distinct entity? A multicenter study of 14 cases. <i>Child's Nervous System</i> , 2013, 29, 565-571.	1.1	13
113	Phase II study of irinotecan in combination with temozolomide (TEMIRI) in children with recurrent or refractory medulloblastoma: a joint ITCC and SIOPE brain tumor study. <i>Neuro-Oncology</i> , 2013, 15, 1236-1243.	1.2	41
114	Reduced H3K27me3 and DNA Hypomethylation Are Major Drivers of Gene Expression in K27M Mutant Pediatric High-Grade Gliomas. <i>Cancer Cell</i> , 2013, 24, 660-672.	16.8	633
115	Time perception in children treated for a cerebellar medulloblastoma. <i>Research in Developmental Disabilities</i> , 2013, 34, 480-494.	2.2	12
116	Impact of extensive surgery in multidisciplinary approach of pterygopalatine/infratemporal fossa soft tissue sarcoma. <i>Pediatric Blood and Cancer</i> , 2013, 60, 928-934.	1.5	9
117	Current and evolving knowledge of prognostic factors for pediatric ependymomas. <i>Future Oncology</i> , 2013, 9, 183-191.	2.4	13
118	Copy Number Gain of 1q25 Predicts Poor Progression-Free Survival for Pediatric Intracranial Ependymomas and Enables Patient Risk Stratification: A Prospective European Clinical Trial Cohort Analysis on Behalf of the Children's Cancer Leukaemia Group (CCLG), Soci�t� Fran�saise d'Oncologie P�diatrique (SFOP), and International Society for Pediatric Oncology (SIOP). <i>Clinical Cancer Research</i> , 2012, 18, 2001-2011.	7.0	111
119	Delays in diagnosis of paediatric cancers: a systematic review and comparison with expert testimony in lawsuits. <i>Lancet Oncology</i> , The, 2012, 13, e445-e459.	10.7	134
120	Teachers' report of learning and behavioural difficulties in children treated for cerebellar tumours. <i>Brain Injury</i> , 2012, 26, 1014-1020.	1.2	21
121	Pseudoprogression after high-dose busulfan-thiotepa with autologous stem cell transplantation and radiation therapy in children with brain tumors: Impact on survival. <i>Neuro-Oncology</i> , 2012, 14, 1413-1421.	1.2	14
122	Mesenchymal Transition and PDGFRA Amplification/Mutation Are Key Distinct Oncogenic Events in Pediatric Diffuse Intrinsic Pontine Gliomas. <i>PLoS ONE</i> , 2012, 7, e30313.	2.5	200
123	Long Time to Diagnosis of Medulloblastoma in Children Is Not Associated with Decreased Survival or with Worse Neurological Outcome. <i>PLoS ONE</i> , 2012, 7, e33415.	2.5	34
124	Critical oncogenic mutations in newly diagnosed pediatric diffuse intrinsic pontine glioma. <i>Pediatric Blood and Cancer</i> , 2012, 58, 489-491.	1.5	111
125	Clinicopathologic prognostic factors in childhood atypical teratoid and rhabdoid tumor of the central nervous system. <i>Cancer</i> , 2012, 118, 3812-3821.	4.1	101
126	Radiotherapy with concurrent and adjuvant temozolomide in children with newly diagnosed diffuse intrinsic pontine glioma. <i>Journal of Neuro-Oncology</i> , 2012, 106, 399-407.	2.9	100

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127	Interval between onset of symptoms and diagnosis of medulloblastoma in children: distribution and determinants in a population-based study. <i>European Journal of Pediatrics</i> , 2012, 171, 25-32.	2.7	39
128	Neuropathological and Neuroradiological Spectrum of Pediatric Malignant Gliomas: Correlation With Outcome. <i>Neurosurgery</i> , 2011, 69, 215-224.	1.1	11
129	Pediatric ependymomas. <i>Current Opinion in Oncology</i> , 2011, 23, 638-642.	2.4	9
130	Hypofractionated radiotherapy in the treatment of diffuse intrinsic pontine glioma in children: a single institution's experience. <i>Journal of Neuro-Oncology</i> , 2011, 104, 773-777.	2.9	48
131	Cerebellar mutism: definitions, classification and grading of symptoms. <i>Child's Nervous System</i> , 2011, 27, 1361-1363.	1.1	35
132	Appraisal of the current staging system for residual medulloblastoma by volumetric analysis. <i>Child's Nervous System</i> , 2011, 27, 2101-2106.	1.1	6
133	Atypical teratoid rhabdoid tumor mimicking beta-catenin-positive nodular medulloblastoma. <i>Acta Neuropathologica</i> , 2011, 121, 429-430.	7.7	3
134	Relationship between the brain radiation dose for the treatment of childhood cancer and the risk of long-term cerebrovascular mortality. <i>Brain</i> , 2011, 134, 1362-1372.	7.6	60
135	Clinical Relevance of Tumor Cells with Stem-Like Properties in Pediatric Brain Tumors. <i>PLoS ONE</i> , 2011, 6, e16375.	2.5	57
136	Utility of Cerebrospinal Fluid Cytology in Newly Diagnosed Childhood Ependymoma. <i>Journal of Pediatric Hematology/Oncology</i> , 2010, 32, 515-518.	0.6	10
137	Medulloblastoma in young children. <i>Pediatric Blood and Cancer</i> , 2010, 54, 635-637.	1.5	52
138	Portrait of Ependymoma Recurrence in Children: Biomarkers of Tumor Progression Identified by Dual-Color Microarray-Based Gene Expression Analysis. <i>PLoS ONE</i> , 2010, 5, e12932.	2.5	35
139	Survival and Prognostic Factors of Early Childhood Medulloblastoma: An International Meta-Analysis. <i>Journal of Clinical Oncology</i> , 2010, 28, 4961-4968.	1.6	273
140	Low Bone Mineral Density and High Incidences of Fractures and Vitamin D Deficiency in 52 Pediatric Cancer Survivors. <i>Hormone Research in Paediatrics</i> , 2010, 74, 319-327.	1.8	33
141	Neuronal differentiation distinguishes supratentorial and infratentorial childhood ependymomas. <i>Neuro-Oncology</i> , 2010, 12, 1126-1134.	1.2	54
142	Incomplete penetrance of the predisposition to medulloblastoma associated with germ-line SUFU mutations. <i>Journal of Medical Genetics</i> , 2010, 47, 142-144.	3.2	51
143	Forniceal glioma in children. <i>Journal of Neurosurgery: Pediatrics</i> , 2009, 4, 249-253.	1.3	5
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