Cynthia Hawkins

List of Publications by Year in Descending Order

Source: https://exaly.com/author-pdf/8434785/cynthia-hawkins-publications-by-year.pdf

Version: 2024-04-25

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

23,218 78 151 239 h-index g-index citations papers 266 6.2 6.64 29,335 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
239	Clinical and economic impact of molecular testing for BRAF fusion in pediatric low-grade Glioma <i>BMC Pediatrics</i> , 2022 , 22, 13	2.6	
238	Splicing is an alternate oncogenic pathway activation mechanism in glioma <i>Nature Communications</i> , 2022 , 13, 588	17.4	0
237	Genomic predictors of response to PD-1 inhibition in children with germline DNA replication repair deficiency <i>Nature Medicine</i> , 2022 ,	50.5	2
236	Comprehensive analysis of the ErbB receptor family in pediatric nervous system tumors and rhabdomyosarcoma. <i>Pediatric Blood and Cancer</i> , 2022 , 69, e29316	3	0
235	Ependymal Tumors Pediatric and Developmental Pathology, 2022, 25, 59-67	2.2	O
234	Medulloblastoma: WHO 2021 and Beyond Pediatric and Developmental Pathology, 2022, 25, 23-33	2.2	1
233	The diverse landscape of histone-mutant pediatric high-grade gliomas: A narrative review. <i>Glioma</i> (Mumbai, India), 2022 , 5, 5	0.3	
232	Immune Checkpoint Inhibition as Single Therapy for Synchronous Cancers Exhibiting Hypermutation: An IRRDC Study <i>JCO Precision Oncology</i> , 2022 , 6, e2100286	3.6	1
231	Building the ecosystem for pediatric neuro-oncology care in Pakistan: Results of a 7-year long twinning program between Canada and Pakistan <i>Pediatric Blood and Cancer</i> , 2022 , e29726	3	1
230	IMMU-13. Dual CTLA4/PD-1 blockade improves survival for replication-repair deficient high-grade gliomas failing single agent PD-1 inhibition: An IRRDC study. <i>Neuro-Oncology</i> , 2022 , 24, i84-i84	1	0
229	IMMU-17. Comprehensive immunological gene expression profiling of pediatric brain tumors. <i>Neuro-Oncology</i> , 2022 , 24, i85-i85	1	1
228	LGG-41. The clinical and molecular landscape of gliomas in adolescents and young adults. <i>Neuro-Oncology</i> , 2022 , 24, i97-i97	1	
227	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	
226	Giant choroid plexus cysts with calvarial erosion: a case report and literature review. <i>Childls Nervous System</i> , 2021 , 37, 2381-2385	1.7	
225	Germline predisposition to glial neoplasms in children and young adults: A narrative review. <i>Glioma</i> (Mumbai, India), 2021 , 4, 68	0.3	
224	Investigating Urinary Circular RNA Biomarkers for Improved Detection of Renal Cell Carcinoma <i>Frontiers in Oncology</i> , 2021 , 11, 814228	5.3	1
223	EXTH-30. HARNESSING CELLULAR STRESS FOR IMMUNE TARGETING OF DIPGS. <i>Neuro-Oncology</i> , 2021 , 23, vi169-vi170	1	

222	CTNI-06. TRAM-01: A PHASE 2 STUDY OF TRAMETINIB FOR PATIENTS WITH PEDIATRIC GLIOMA WITH ACTIVATION OF THE MAPK/ERK PATHWAY. <i>Neuro-Oncology</i> , 2021 , 23, vi59-vi60	1	
221	INNV-43. MORE THAN WHAT MEETS THE EYE: ETMR AN UNDER RECOGNISED ATYPICAL BRAINSTEM PRIMARY. A RARE BRAIN TUMOR CONSORTIUM (RBTC) STUDY. <i>Neuro-Oncology</i> , 2021 , 23, vi114-vi115	1	
220	Pediatric Glial Tumors. Pediatric and Developmental Pathology, 2021, 10935266211009101	2.2	0
219	A Practical Approach to the Evaluation and Diagnosis of Pediatric CNS Tumors. <i>Pediatric and Developmental Pathology</i> , 2021 , 10935266211007022	2.2	
218	MetaFusion: A high-confidence metacaller for filtering and prioritizing RNA-seq gene fusion candidates. <i>Bioinformatics</i> , 2021 ,	7.2	2
217	Characteristics of Patients © 0 Years of Age with Diffuse Intrinsic Pontine Glioma: A Report from the International DIPG Registry. <i>Neuro-Oncology</i> , 2021 ,	1	1
216	The 2021 WHO Classification of Tumors of the Central Nervous System: a summary. <i>Neuro-Oncology</i> , 2021 , 23, 1231-1251	1	708
215	HGG-39. ALTERNATIVE SPLICING OF NEUROFIBROMIN 1 IS ASSOCIATED WITH ELEVATED MAPK ACTIVITY AND POOR PROGNOSIS IN HIGH-GRADE GLIOMA. <i>Neuro-Oncology</i> , 2021 , 23, i25-i25	1	78
214	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
213	OMRT-8. Precision targeting of cellular pathways with complementary diagnostics. <i>Neuro-Oncology Advances</i> , 2021 , 3, ii8-ii8	0.9	78
212	Salvage chemotherapy after failure of targeted therapy in a child with BRAF V600E low-grade glioma. <i>Pediatric Blood and Cancer</i> , 2021 , 68, e28561	3	1
211	Re-irradiation with concurrent BRAF and MEK inhibitor therapy. <i>Pediatric Blood and Cancer</i> , 2021 , 68, e28838	3	O
210	Clinical and molecular heterogeneity of pineal parenchymal tumors: a consensus study. <i>Acta Neuropathologica</i> , 2021 , 141, 771-785	14.3	9
209	Longitudinal Assessment of Enhancing Foci of Abnormal Signal Intensity in Neurofibromatosis Type 1. <i>American Journal of Neuroradiology</i> , 2021 , 42, 766-773	4.4	
208	Radiomics of Pediatric Low-Grade Gliomas: Toward a Pretherapeutic Differentiation of Mutated and -Fused Tumors. <i>American Journal of Neuroradiology</i> , 2021 , 42, 759-765	4.4	5
207	Clinical Outcomes and Patient-Matched Molecular Composition of Relapsed Medulloblastoma. <i>Journal of Clinical Oncology</i> , 2021 , 39, 807-821	2.2	7
206	Local FK506 drug delivery enhances nerve regeneration through fresh, unprocessed peripheral nerve allografts. <i>Experimental Neurology</i> , 2021 , 341, 113680	5.7	7
205	Upfront Adjuvant Immunotherapy of Replication Repair-Deficient Pediatric Glioblastoma With Chemoradiation-Sparing Approach <i>JCO Precision Oncology</i> , 2021 , 5, 1426-1431	3.6	O

204	Survival Benefit for Individuals With Constitutional Mismatch Repair Deficiency Undergoing Surveillance. <i>Journal of Clinical Oncology</i> , 2021 , 39, 2779-2790	2.2	10
203	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	14.5	1
202	SYST-04. TRAM-01: A PHASE 2 STUDY OF TRAMETINIB FOR PATIENTS WITH PEDIATRIC GLIOMA WITH ACTIVATION OF THE MAPK/ERK PATHWAY. <i>Neuro-Oncology Advances</i> , 2021 , 3, iv9-iv9	0.9	O
201	EPCO-16. ONCOHISTONE INTERACTOME PROFILING UNCOVERS MECHANISMS OF CHROMATIN DISRUPTION AND IDENTIFIES POTENTIAL THERAPEUTIC TARGETS IN PEDIATRIC HIGH-GRADE GLIOMA. <i>Neuro-Oncology</i> , 2021 , 23, vi5-vi5	1	
200	A novel central nervous system embryonal tumor successfully treated with multi-modal therapy highlights limitation of methylation-based tumor classification <i>Pediatric Blood and Cancer</i> , 2021 , e295	52 ð	
199	Outcomes of BRAF V600E Pediatric Gliomas Treated With Targeted BRAF Inhibition. <i>JCO Precision Oncology</i> , 2020 , 4,	3.6	23
198	MR imaging features of diffuse intrinsic pontine glioma and relationship to overall survival: report from the International DIPG Registry. <i>Neuro-Oncology</i> , 2020 , 22, 1647-1657	1	12
197	cIMPACT-NOW update 7: advancing the molecular classification of ependymal tumors. <i>Brain Pathology</i> , 2020 , 30, 863-866	6	51
196	Pediatric low-grade glioma in the era of molecular diagnostics. <i>Acta Neuropathologica Communications</i> , 2020 , 8, 30	7.3	52
195	Mutant ACVR1 Arrests Glial Cell Differentiation to Drive Tumorigenesis in Pediatric Gliomas. <i>Cancer Cell</i> , 2020 , 37, 308-323.e12	24.3	21
194	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
193	Immunohistochemical and nanoString-Based Subgrouping of Clinical Medulloblastoma Samples. Journal of Neuropathology and Experimental Neurology, 2020 , 79, 437-447	3.1	8
192	An update on the CNS manifestations of brain tumor polyposis syndromes. <i>Acta Neuropathologica</i> , 2020 , 139, 703-715	14.3	19
191	Integrated Molecular and Clinical Analysis of 1,000 Pediatric Low-Grade Gliomas. <i>Cancer Cell</i> , 2020 , 37, 569-583.e5	24.3	92
190	Clinical impact of combined epigenetic and molecular analysis of pediatric low-grade gliomas. <i>Neuro-Oncology</i> , 2020 , 22, 1474-1483	1	14
189	Locoregional delivery of CAR T cells to the cerebrospinal fluid for treatment of metastatic medulloblastoma and ependymoma. <i>Nature Medicine</i> , 2020 , 26, 720-731	50.5	60
188	TAMI-29. MULTIFACTORIAL UPREGULATION OF ID1 DRIVES DIPG INVASIVENESS AND IS THERAPEUTICALLY TARGETABLE. <i>Neuro-Oncology</i> , 2020 , 22, ii219-ii219	1	
187	NIMG-31. NON-DIPG PATIENTS ENROLLED IN THE INTERNATIONAL DIPG REGISTRY: HISTOPATHOLOGIC EVALUATION OF CENTRAL NEURO-IMAGING REVIEW. <i>Neuro-Oncology</i> , 2020 ,	1	

186	CTNI-24. A PHASE 2 STUDY OF TRAMETINIB FOR PATIENTS WITH PEDIATRIC GLIOMA WITH ACTIVATION OF THE MAPK/ERK PATHWAY. TRAM-01. <i>Neuro-Oncology</i> , 2020 , 22, ii47-ii47	1	
185	DDRE-06. CELLULAR STRESS RESPONSE IN DIPG THERAPY. <i>Neuro-Oncology</i> , 2020 , 22, ii62-ii62	1	
184	ATRT-33. ENABLING RAPID CLASSIFICATION OF ATRT WITH NANOSTRING NCOUNTER PLATFORM. <i>Neuro-Oncology</i> , 2020 , 22, iii282-iii282	1	78
183	MBCL-25. PILOT STUDY OF A SURGERY AND CHEMOTHERAPY-ONLY APPROACH IN THE UPFRONT THERAPY OF CHILDREN WITH WNT-POSITIVE STANDARD RISK MEDULLOBLASTOMA: UPDATED OUTCOMES. <i>Neuro-Oncology</i> , 2020 , 22, iii393-iii394	1	1
182	MODL-25. REPLICATION REPAIR DEFICIENT MOUSE MODELS PROVIDE INSIGHT ON HYPERMUTANT BRAIN TUMOURS, MECHANISMS OF IMMUNE EVASION, AND COMBINATORIAL IMMUNOTHERAPY. <i>Neuro-Oncology</i> , 2020 , 22, iii416-iii416	1	78
181	DIPG-46. NON-DIPG PATIENTS ENROLLED IN THE INTERNATIONAL DIPG REGISTRY: HISTOPATHOLOGIC EVALUATION OF CENTRAL NEURO-IMAGING REVIEW. <i>Neuro-Oncology</i> , 2020 , 22, iii295-iii296	1	78
180	LGG-13. THE CLINICAL AND MOLECULAR LANDSCAPE OF GLIOMAS IN ADOLESCENTS AND YOUNG ADULTS. <i>Neuro-Oncology</i> , 2020 , 22, iii368-iii368	1	78
179	LGG-19. SPINAL LOW-GRADE GLIOMAS IN CANADIAN CHILDREN: A MULTI-CENTRE RETROSPECTIVE REVIEW. <i>Neuro-Oncology</i> , 2020 , 22, iii369-iii370	1	78
178	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
177	LGG-34. CLINICAL AND MOLECULAR CHARACTERIZATION OF A MULTI-INSTITUTIONAL COHORT OF PEDIATRIC SPINAL CORD LOW-GRADE GLIOMAS. <i>Neuro-Oncology</i> , 2020 , 22, iii373-iii373	1	78
176	HGG-20. DIAGNOSTIC AND BIOLOGICAL ROLE OF METHYLATION PATTERNS IN REPLICATION REPAIR DEFICIENT HIGH GRADE GLIOMAS. <i>Neuro-Oncology</i> , 2020 , 22, iii347-iii348	1	78
175	LGG-50. INTEGRATED MOLECULAR AND CLINICAL ANALYSIS OF 1,000 PEDIATRIC LOW-GRADE GLIOMAS UNCOVERS NOVEL SUBGROUPS FOR CLINICAL RISK STRATIFICATION. <i>Neuro-Oncology</i> , 2020 , 22, iii375-iii376	1	78
174	DIPG-59. UPREGULATION OF PRENATAL PONTINE ID1 SIGNALING IN DIPG. <i>Neuro-Oncology</i> , 2020 , 22, iii298-iii299	1	78
173	PATH-14. GENETIC SUSCEPTIBILITY AND OUTCOMES OF PEDIATRIC, ADOLESCENT AND YOUNG ADULT IDH-MUTANT ASTROCYTOMAS. <i>Neuro-Oncology</i> , 2020 , 22, iii427-iii427	1	78
172	LGG-25. A PHASE 2 STUDY OF TRAMETINIB FOR PATIENTS WITH PEDIATRIC GLIOMA WITH ACTIVATION OF THE MAPK/ERK PATHWAY. TRAM-01. <i>Neuro-Oncology</i> , 2020 , 22, iii371-iii371	1	1
171	LGG-55. OUTCOME OF BRAF V600E PEDIATRIC GLIOMAS TREATED WITH TARGETED BRAF INHIBITION. <i>Neuro-Oncology</i> , 2020 , 22, iii377-iii377	1	78
170	Medulloblastoma Arises from the Persistence of a Rare and Transient Sox2 Granule Neuron Precursor. <i>Cell Reports</i> , 2020 , 31, 107511	10.6	10
169	IMMU-18. FAVORABLE OUTCOME IN REPLICATION REPAIR DEFICIENT HYPERMUTANT BRAIN TUMORS TO IMMUNE CHECKPOINT INHIBITION: AN INTERNATIONAL RRD CONSORTIUM REGISTRY STUDY. <i>Neuro-Oncology</i> , 2020 , 22, iii363-iii363	1	1

168	MBRS-54. POOR SURVIVAL IN REPLICATION REPAIR DEFICIENT HYPERMUTANT MEDULLOBLASTOMA AND CNS EMBRYONAL TUMORS: A REPORT FROM THE INTERNATIONAL RRD CONSORTIUM. <i>Neuro-Oncology</i> , 2020 , 22, iii407-iii407	1	1
167	Multiplexed Digital Detection of B-Cell Acute Lymphoblastic Leukemia Fusion Transcripts Using the NanoString nCounter System. <i>Journal of Molecular Diagnostics</i> , 2020 , 22, 72-80	5.1	6
166	Modeling DIPG in the mouse brainstem. <i>Neuro-Oncology</i> , 2020 , 22, 307-308	1	1
165	B7-H3 as a Prognostic Biomarker and Therapeutic Target in Pediatric central nervous system Tumors. <i>Translational Oncology</i> , 2020 , 13, 365-371	4.9	15
164	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	14.3	28
163	Phase II Study of Nonmetastatic Desmoplastic Medulloblastoma in Children Younger Than 4 Years of Age: A Report of the Children's Oncology Group (ACNS1221). <i>Journal of Clinical Oncology</i> , 2020 , 38, 223-231	2.2	19
162	Indolent course of brainstem tumors with K27M-H3.3 mutation. <i>Pediatric Blood and Cancer</i> , 2020 , 67, e28102	3	1
161	Clinical and molecular characterization of a multi-institutional cohort of pediatric spinal cord low-grade gliomas. <i>Neuro-Oncology Advances</i> , 2020 , 2, vdaa103	0.9	3
160	Senescence Induced by BMI1 Inhibition Is a Therapeutic Vulnerability in H3K27M-Mutant DIPG. <i>Cell Reports</i> , 2020 , 33, 108286	10.6	10
159	An OTX2-PAX3 signaling axis regulates Group 3 medulloblastoma cell fate. <i>Nature Communications</i> , 2020 , 11, 3627	17.4	8
158	Pontine gliomas a 10-year population-based study: a report from The Canadian Paediatric Brain Tumour Consortium (CPBTC). <i>Journal of Neuro-Oncology</i> , 2020 , 149, 45-54	4.8	4
157	Epigenetic activation of a RAS/MYC axis in H3.3K27M-driven cancer. <i>Nature Communications</i> , 2020 , 11, 6216	17.4	8
156	Germline-driven replication repair-deficient high-grade gliomas exhibit unique hypomethylation patterns. <i>Acta Neuropathologica</i> , 2020 , 140, 765-776	14.3	8
155	Cancer proteome and metabolite changes linked to SHMT2. <i>PLoS ONE</i> , 2020 , 15, e0237981	3.7	7
154	Diffuse midline glioma: review of epigenetics. <i>Journal of Neuro-Oncology</i> , 2020 , 150, 27-34	4.8	11
153	ETMR-22. TITLE: DEFINING THE CLINICAL AND PROGNOSTIC LANDSCAPE OF EMBRYONAL TUMORS WITH MULTI-LAYERED ROSETTES (ETMRs), A RARE BRAIN TUMOR REGISTRY (RBTC) STUDY. <i>Neuro-Oncology</i> , 2020 , 22, iii327-iii328	1	78
152	BRAF V600E mutant oligodendroglioma-like tumors with chromosomal instability in adolescents and young adults. <i>Brain Pathology</i> , 2020 , 30, 515-523	6	5
151	Targeting reduced mitochondrial DNA quantity as a therapeutic approach in pediatric high-grade gliomas. <i>Neuro-Oncology</i> , 2020 , 22, 139-151	1	21

(2019-2020)

150	cIMPACT-NOW update 6: new entity and diagnostic principle recommendations of the cIMPACT-Utrecht meeting on future CNS tumor classification and grading. <i>Brain Pathology</i> , 2020 , 30, 844-856	6	196
149	Pearls & Oy-sters: Fatal brain edema is a rare complication of severe CACNA1A-related disorder. <i>Neurology</i> , 2020 , 94, 631-634	6.5	3
148	Re-irradiation for children with recurrent medulloblastoma in Toronto, Canada: a 20-year experience. <i>Journal of Neuro-Oncology</i> , 2019 , 145, 107-114	4.8	5
147	Alterations in ALK/ROS1/NTRK/MET drive a group of infantile hemispheric gliomas. <i>Nature Communications</i> , 2019 , 10, 4343	17.4	95
146	Transcriptional repressor REST drives lineage stage-specific chromatin compaction at and increases AKT activation in a mouse model of medulloblastoma. <i>Science Signaling</i> , 2019 , 12,	8.8	11
145	Diffuse intrinsic pontine glioma ventricular peritoneal shunt metastasis: a case report and literature review. <i>Childls Nervous System</i> , 2019 , 35, 861-864	1.7	3
144	Repeat irradiation for children with supratentorial high-grade glioma. <i>Pediatric Blood and Cancer</i> , 2019 , 66, e27881	3	5
143	Ongoing issues with the management of children with Constitutional Mismatch Repair Deficiency syndrome. <i>European Journal of Medical Genetics</i> , 2019 , 62, 103706	2.6	3
142	DIPG-22. GENETIC MODELING IMPLICATES RAS AND MYC AS KEY EPIGENETICALLY ACTIVATED TRANSCRIPTIONAL TARGETS OF H3K27M-DRIVEN CANCER. <i>Neuro-Oncology</i> , 2019 , 21, ii73-ii73	1	78
141	DIPG-35. OPEN DIPG INITIATIVE: A PLATFORM FOR ACCELERATING DISCOVERY THROUGH DATA ACCESS, CONSOLIDATION AND HARMONIZATION. <i>Neuro-Oncology</i> , 2019 , 21, ii76-ii76	1	78
140	HGG-22. CHARACTERIZING THE ROLE H3.3G34R MUTATION IN PEDIATRIC HIGH GRADE ASTROCYTOMA. <i>Neuro-Oncology</i> , 2019 , 21, ii91-ii91	1	78
139	LGG-07. CLINICAL FEATURES OF NON-CANONICAL MOLECULAR DRIVERS IN PLGG; AN UPDATE FORM THE INTERNATIONAL PLGG TASKFORCE. <i>Neuro-Oncology</i> , 2019 , 21, ii100-ii100	1	78
138	DIPG-36. CLINICAL, RADIOLOGICAL, AND HISTO-MOLECULAR CHARACTERISTICS OF DIFFUSE INTRINSIC PONTINE GLIOMA IN PATIENTS WHO SURVIVE LESS THAN 3 MONTHS FROM DIAGNOSIS: A REPORT FROM THE INTERNATIONAL DIPG REGISTRY. <i>Neuro-Oncology</i> , 2019 , 21, ii76-ii77	, 1	78
137	IMMU-20. IMMUNE AND TUMOR BIOMARKERS OF OUTCOME IN REPLICATION REPAIR DEFICIENT BRAIN TUMORS TREATED WITH IMMUNE CHECKPOINT INHIBITORS: UPDATES FROM THE INTERNATIONAL REPLICATION REPAIR DEFICIENCY CONSORTIUM. <i>Neuro-Oncology</i> , 2019 , 21, ii96-ii97	1	78
136	LGG-01. BRAF V600E MUTANT OLIGODENDROGLIOMA-LIKE TUMORS WITH CHROMOSOMAL INSTABILITY IN ADOLESCENT AND YOUNG ADULT. <i>Neuro-Oncology</i> , 2019 , 21, ii98-ii98	1	78
135	HGG-18. ALTERNATIVE SPLICING OF NEUROFIBROMIN 1 IS ASSOCIATED WITH ELEVATED MAPK ACTIVITY AND POOR PROGNOSIS IN HIGH-GRADE GLIOMA. <i>Neuro-Oncology</i> , 2019 , 21, ii90-ii90	1	78
134	HGG-19. MOLECULAR ANALYSIS UNCOVERS 3 DISTINCT SUBGROUPS AND MULTIPLE TARGETABLE GENE FUSIONS IN INFANT GLIOMAS. <i>Neuro-Oncology</i> , 2019 , 21, ii90-ii91	1	78
133	cIMPACT-NOW update 4: diffuse gliomas characterized by MYB, MYBL1, or FGFR1 alterations or BRAF mutation. <i>Acta Neuropathologica</i> , 2019 , 137, 683-687	14.3	92

132	Recessive mutations in muscle-specific isoforms of FXR1 cause congenital multi-minicore myopathy. <i>Nature Communications</i> , 2019 , 10, 797	17.4	10
131	Survival and functional outcomes of molecularly defined childhood posterior fossa ependymoma: Cure at a cost. <i>Cancer</i> , 2019 , 125, 1867-1876	6.4	26
130	Sarcoma Subgrouping by Detection of Fusion Transcripts Using NanoString nCounter Technology. <i>Pediatric and Developmental Pathology</i> , 2019 , 22, 205-213	2.2	7
129	DNA methylation signature is prognostic of choroid plexus tumor aggressiveness. <i>Clinical Epigenetics</i> , 2019 , 11, 117	7.7	12
128	MEDU-04. AN OTX2-PAX GENE NETWORK REGULATES GROUP 3 MEDULLOBLASTOMA DIFFERENTIATION AND TUMOR GROWTH. <i>Neuro-Oncology</i> , 2019 , 21, ii103-ii104	1	78
127	Apparent Lack of Derived HLA Class I Presented Neoantigens Hampers Neoplastic Cell Targeting by CD8 T Cells in Langerhans Cell Histiocytosis. <i>Frontiers in Immunology</i> , 2019 , 10, 3045	8.4	1
126	LGG-16. PREDICTORS OF OUTCOME IN BRAF-V600E PEDIATRIC GLIOMAS TREATED WITH BRAF INHIBITORS: A REPORT FROM THE PLGG TASKFORCE. <i>Neuro-Oncology</i> , 2019 , 21, ii102-ii102	1	78
125	MEDU-34. PILOT STUDY OF A SURGERY AND CHEMOTHERAPY-ONLY APPROACH IN THE UPFRONT THERAPY OF CHILDREN WITH WNT-POSITIVE STANDARD RISK MEDULLOBLASTOMA. Neuro-Oncology, 2019, 21, ii110-ii110	1	7
124	Acute MR-Guided High-Intensity Focused Ultrasound Lesion Assessment Using Diffusion-Weighted Imaging and Histological Analysis. <i>Frontiers in Neurology</i> , 2019 , 10, 1069	4.1	6
123	TMOD-10. REPLICATION REPAIR DEFICIENT MOUSE MODELS PROVIDE INSIGHT ON HYPERMUTANT BRAIN TUMOURS AND COMBINATIONAL IMMUNOTHERAPY. <i>Neuro-Oncology</i> , 2019 , 21, ii123-ii123	1	78
122	International experience in the development of patient-derived xenograft models of diffuse intrinsic pontine glioma. <i>Journal of Neuro-Oncology</i> , 2019 , 141, 253-263	4.8	19
121	Craniospinal irradiation as part of re-irradiation for children with recurrent intracranial ependymoma. <i>Neuro-Oncology</i> , 2019 , 21, 547-557	1	16
120	Detecting Stem Cell Marker Expression Using the NanoString nCounter System. <i>Methods in Molecular Biology</i> , 2019 , 1869, 57-67	1.4	2
119	Reirradiation in patients with diffuse intrinsic pontine gliomas: The Canadian experience. <i>Pediatric Blood and Cancer</i> , 2018 , 65, e26988	3	33
118	Mitochondrial POLG related disorder presenting prenatally with fetal cerebellar growth arrest. <i>Metabolic Brain Disease</i> , 2018 , 33, 1369-1373	3.9	6
117	CD271 Cells Are Diagnostic and Prognostic and Exhibit Elevated MAPK Activity in SHH Medulloblastoma. <i>Cancer Research</i> , 2018 , 78, 4745-4759	10.1	16
116	EMBR-15. DIAGNOSTIC RE-EVALUATION AND POOLED CLINICAL DATA ANALYSIS OF PATIENTS WITH PREVIOUS DIAGNOSIS OF CNS-PNET. <i>Neuro-Oncology</i> , 2018 , 20, i72-i72	1	2
115	DIPG-70. CLINICAL, RADIOLOGICAL, PATHOLOGICAL AND MOLECULAR CHARACTERISTICS OF CHILDREN . <i>Neuro-Oncology</i> , 2018 , 20, i63-i63	1	78

114	Sustained Response to Targeted Therapy in a Patient With Disseminated Anaplastic Pleomorphic Xanthoastrocytoma. <i>Journal of Pediatric Hematology/Oncology</i> , 2018 , 40, 478-482	1.2	14
113	Pediatric low-grade gliomas: next biologically driven steps. <i>Neuro-Oncology</i> , 2018 , 20, 160-173	1	76
112	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. <i>Journal of Clinical Oncology</i> , 2018 , 36, 1963-1972	2.2	125
111	LGG-10. EPIGENETIC/GENETIC/MORPHOLOGIC ANALYSES REVEAL CLINICAL/PROGNOSTIC INSIGHT OF PEDIATRIC LOW GRADE GLIOMAS. <i>Neuro-Oncology</i> , 2018 , 20, i106-i106	1	78
110	ATRT-40. IMPACT OF MOLECULAR SUBTYPES ON TREATMENT OUTCOMES IN RHABDOID TUMORS - A REPORT FROM THE RARE TUMOR CONSORTIUM. <i>Neuro-Oncology</i> , 2018 , 20, i36-i36	1	78
109	DIPG-55. TARGETING SENESCENT CELLS WITH ABT-263 ENHANCES CELL DEATH INDUCED BY BMI1 INHIBITION AND IONIZING RADIATION IN DIPG. <i>Neuro-Oncology</i> , 2018 , 20, i60-i60	1	78
108	EPEN-31. SUBGROUP SPECIFIC LONG-TERM SURVIVAL AND NEUROCOGNITIVE OUTCOMES IN POSTERIOR FOSSA EPENDYMOMA (PFE). <i>Neuro-Oncology</i> , 2018 , 20, i79-i79	1	78
107	HGG-17. TUMOR MUTATIONAL BURDEN ANALYSIS OF PEDIATRIC TUMORS PROVIDES A DIAGNOSTIC TOOL FOR GERMLINE REPLICATION REPAIR DEFICIENCY AND PREDICT RESPONSE TO IMMUNE CHECKPOINT INHIBITION. <i>Neuro-Oncology</i> , 2018 , 20, i92-i92	1	78
106	MBCL-08. MOLECULAR CHARACTERIZATION OF NODULAR DESMOPLASTIC MEDULLOBLASTOMAS IN YOUNG CHILDREN TREATED ON ACNS1221. A REPORT FROM THE CHILDREN ONCOLOGY GROUP. <i>Neuro-Oncology</i> , 2018 , 20, i118-i119	1	2
105	MBRS-62. REPRESSIVE CHROMATIN REMODELERS IN SHH-DRIVEN MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2018 , 20, i141-i141	1	78
104	IMMU-25. PROGRAMMED CELL DEATH-LIGAND 1 (PD-L1) IS NOT EXPRESSED IN DIFFUSE INTRINSIC PONTINE GLIOMA (DIPG) TUMOR CELLS. <i>Neuro-Oncology</i> , 2018 , 20, vi126-vi126	1	1
103	EAPH-06. HYPERMUTANT PEDIATRIC HIGH GRADE GLIOMAS ARE DRIVEN BY RAS/MAPK MUTATIONS AND RESPOND TO MEK INHIBITION. <i>Neuro-Oncology</i> , 2018 , 20, i66-i66	1	78
102	LGG-60. THE GENETIC LANDSCAPE OF PEDIATRIC LOW-GRADE GLIOMAS: INCIDENCE, PROGNOSIS AND RESPONSE TO THERAPY. <i>Neuro-Oncology</i> , 2018 , 20, i117-i117	1	1
101	TBIO-30. MOLECULAR LANDSCAPE AND CLINICAL CORRELATIONS OF CNS SARCOMAS. Neuro-Oncology, 2018 , 20, i186-i186	1	78
100	LGG-49. MOLECULAR ALTERATIONS IN PREGNANT ADOLESCENT AND YOUNG ADULT WOMEN WITH GLIOMA. <i>Neuro-Oncology</i> , 2018 , 20, i115-i115	1	78
99	DIPG-69. CHARACTERISTICS OF PATIENTS 110 YEARS OF AGE WITH DIFFUSE INTRINSIC PONTINE GLIOMA: A REPORT FROM THE INTERNATIONAL DIPG REGISTRY. <i>Neuro-Oncology</i> , 2018 , 20, i63-i63	1	1
98	EMBR-12. IMPROVED DIAGNOSTIC ALGORITHM FOR DIFFERENTIAL DIAGNOSTICS OF CNS EMBRYONAL TUMORS (FORMER CNS-PNET) BY NEUROPATHOLOGICAL RE-EVALUATION OF 256 CASES AND CROSSVALIDATION BY METHYLATION CLASSIFICATION. <i>Neuro-Oncology</i> , 2018 , 20, i71-i71	1	78
97	LGG-59. REMARKABLE OBJECTIVE RESPONSE AND FAVORABLE SURVIVAL FOR BRAF-V600E CHILDHOOD LOW-GRADE GLIOMAS TO BRAF INHIBITORS COMPARED CONVENTIONAL CHEMOTHERAPY. <i>Neuro-Oncology</i> , 2018 , 20, i117-i117	1	78

96	Two different STAT1 gain-of-function mutations lead to diverse IFN-Emediated gene expression. <i>Npj Genomic Medicine</i> , 2018 , 3, 23	6.2	7
95	DIPG-38. ID1 EXPRESSION CORRELATES WITH H3F3A K27M MUTATION AND EXTRA-PONTINE INVASION IN DIPG. <i>Neuro-Oncology</i> , 2018 , 20, i56-i56	1	78
94	EMBR-17. PINEOBLASTOMA SEGREGATES INTO MOLECULAR SUBTYPES WITH DISTINCT CLINICOPATHOLOGIC FEATURES: REPORT FROM THE RARE BRAIN TUMOR CONSORTIUM. <i>Neuro-Oncology</i> , 2018 , 20, i72-i73	1	78
93	Atypical teratoid rhabdoid tumor in the first year of life: the Canadian ATRT registry experience and review of the literature. <i>Journal of Neuro-Oncology</i> , 2017 , 132, 155-162	4.8	29
92	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	4.8	19
91	Prognostic relevance of miR-124-3p and its target TP53INP1 in pediatric ependymoma. <i>Genes Chromosomes and Cancer</i> , 2017 , 56, 639-650	5	14
90	H3 K27M mutations are extremely rare in posterior fossa group A ependymoma. <i>Childls Nervous System</i> , 2017 , 33, 1047-1051	1.7	29
89	A comprehensive review of paediatric low-grade diffuse glioma: pathology, molecular genetics and treatment. <i>Brain Tumor Pathology</i> , 2017 , 34, 51-61	3.2	33
88	Noncompaction cardiomyopathy in an infant with Walker-Warburg syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2017 , 173, 3082-3086	2.5	5
87	Integrated Molecular Meta-Analysis of 1,000 Pediatric High-Grade and Diffuse Intrinsic Pontine Glioma. <i>Cancer Cell</i> , 2017 , 32, 520-537.e5	24.3	423
86	Comprehensive Analysis of Hypermutation in Human Cancer. <i>Cell</i> , 2017 , 171, 1042-1056.e10	56.2	417
85	Therapeutic and Prognostic Implications of BRAF V600E in Pediatric Low-Grade Gliomas. <i>Journal of Clinical Oncology</i> , 2017 , 35, 2934-2941	2.2	153
84	Pediatric high-grade glioma: biologically and clinically in need of new thinking. <i>Neuro-Oncology</i> , 2017 , 19, 153-161	1	125
83	Immunohistochemical analysis of H3K27me3 demonstrates global reduction in group-A childhood posterior fossa ependymoma and is a powerful predictor of outcome. <i>Acta Neuropathologica</i> , 2017 , 134, 705-714	14.3	114
82	Identification of complex genomic rearrangements in cancers using CouGaR. <i>Genome Research</i> , 2017 , 27, 107-117	9.7	21
81	cIMPACT-NOW (the consortium to inform molecular and practical approaches to CNS tumor taxonomy): a new initiative in advancing nervous system tumor classification. <i>Brain Pathology</i> , 2017 , 27, 851-852	6	36
80	Cribriform neuroepithelial tumor: molecular characterization of a SMARCB1-deficient non-rhabdoid tumor with favorable long-term outcome. <i>Brain Pathology</i> , 2017 , 27, 411-418	6	34
79	Embryonal tumor with multilayered rosettes, C19MC-altered: Report of an extremely rare malignant pediatric central nervous system neoplasm. SAGE Open Medical Case Reports, 2017, 5, 2050:	313 X 17	745208

78	Germline and somatic mutations in with diverse neurodevelopmental phenotypes. <i>Neurology: Genetics</i> , 2017 , 3, e199	3.8	28
77	ACNS1221: A phase II study for the treatment of non metastatic desmoplastic medulloblastoma in children less than 4 years of ageA report from the Children Oncology Group <i>Journal of Clinical Oncology</i> , 2017 , 35, 10505-10505	2.2	7
76	Molecular alterations to predict survival and response to chemotherapy of pediatric low-grade glioma <i>Journal of Clinical Oncology</i> , 2017 , 35, 10503-10503	2.2	
75	Pediatric low-grade gliomas: implications of the biologic era. <i>Neuro-Oncology</i> , 2017 , 19, 750-761	1	47
74	Synchronous glioblastoma and medulloblastoma in a child with mismatch repair mutation. <i>Childls Nervous System</i> , 2016 , 32, 553-7	1.7	11
73	Lowered H3K27me3 and DNA hypomethylation define poorly prognostic pediatric posterior fossa ependymomas. <i>Science Translational Medicine</i> , 2016 , 8, 366ra161	17.5	109
72	Profound clinical and radiological response to BRAF inhibition in a 2-month-old diencephalic child with hypothalamic/chiasmatic glioma. <i>Pediatric Blood and Cancer</i> , 2016 , 63, 2038-41	3	43
71	LG-19IMMUNOHISTOCHEMISTRY IS HIGHLY SENSITIVE AND SPECIFIC FOR THE DETECTION OF BRAF V600E STATUS IN PEDIATRIC LOW-GRADE GLIOMA. <i>Neuro-Oncology</i> , 2016 , 18, iii82.3-iii82	1	0
70	PINK1 Is a Negative Regulator of Growth and the Warburg Effect in Glioblastoma. <i>Cancer Research</i> , 2016 , 76, 4708-19	10.1	80
69	Clinical and treatment factors determining long-term outcomes for adult survivors of childhood low-grade glioma: A population-based study. <i>Cancer</i> , 2016 , 122, 1261-9	6.4	77
68	Medulloblastoma subgroup-specific outcomes in irradiated children: who are the true high-risk patients?. <i>Neuro-Oncology</i> , 2016 , 18, 291-7	1	86
67	Rasmussen's encephalitis: advances in management and patient outcomes. <i>Childls Nervous System</i> , 2016 , 32, 629-40	1.7	21
66	Spatial genomic heterogeneity in diffuse intrinsic pontine and midline high-grade glioma: implications for diagnostic biopsy and targeted therapeutics. <i>Acta Neuropathologica Communications</i> , 2016 , 4, 1	7.3	93
65	Pediatric thalamic tumors in the MRI era: a Canadian perspective. <i>Childls Nervous System</i> , 2016 , 32, 269-6	8 0 7	29
64	JPO2/CDCA7L and LEDGF/p75 Are Novel Mediators of PI3K/AKT Signaling and Aggressive Phenotypes in Medulloblastoma. <i>Cancer Research</i> , 2016 , 76, 2802-12	10.1	11
63	Lethal Disorder of Mitochondrial Fission Caused by Mutations in DNM1L. <i>Journal of Pediatrics</i> , 2016 , 171, 313-6.e1-2	3.6	52
62	MYB-QKI rearrangements in angiocentric glioma drive tumorigenicity through a tripartite mechanism. <i>Nature Genetics</i> , 2016 , 48, 273-82	36.3	154
61	Hemorrhagic presentations of cerebellar pilocytic astrocytomas in children resulting in death: report of 2 cases. <i>Journal of Neurosurgery: Pediatrics</i> , 2016 , 17, 446-52	2.1	11

60	Relationship of BRAF V600E and associated secondary mutations on survival rate and response to conventional therapies in childhood low-grade glioma <i>Journal of Clinical Oncology</i> , 2016 , 34, 10509-10	5 09	3
59	Re-irradiation for relapsed paediatric ependymoma <i>Journal of Clinical Oncology</i> , 2016 , 34, 10565-1056	52.2	
58	Imaging of metastatic medulloblastoma in the molecular era Journal of Clinical Oncology, 2016, 34, e2	2 00 3-e	22003
57	LG-66CLINICAL AND TREATMENT FACTORS DETERMINING LONG-TERM OUTCOMES FOR ADULT SURVIVORS OF CHILDHOOD LOW-GRADE GLIOMA: A POPULATION-BASED STUDY. <i>Neuro-Oncology</i> , 2016 , 18, iii94.1-iii94	1	78
56	PNR-32UPDATE OF DIAGNOSTICS OF PRIMITIVE NEUROECTODERMAL TUMOURS OF THE CNS - NEUROPATHOLOGICAL RE-EVALUATION OF 99 CASES. <i>Neuro-Oncology</i> , 2016 , 18, iii13.1-iii13	1	78
55	High frequency of mismatch repair deficiency among pediatric high grade gliomas in Jordan. <i>International Journal of Cancer</i> , 2016 , 138, 380-5	7.5	48
54	An integrative molecular and genomic analysis of pediatric hemispheric low-grade gliomas: an update. <i>Childls Nervous System</i> , 2016 , 32, 1789-97	1.7	22
53	Management and outcome of chordomas in the pediatric population: The Hospital for Sick Children experience and review of the literature. <i>Journal of Clinical Neuroscience</i> , 2016 , 34, 169-176	2.2	19
52	Targeted detection of genetic alterations reveal the prognostic impact of H3K27M and MAPK pathway aberrations in paediatric thalamic glioma. <i>Acta Neuropathologica Communications</i> , 2016 , 4, 93	7.3	77
51	Intellectual Outcome in Molecular Subgroups of Medulloblastoma. <i>Journal of Clinical Oncology</i> , 2016 , 34, 4161-4170	2.2	56
50	Molecular characterization of choroid plexus tumors reveals novel clinically relevant subgroups. <i>Clinical Cancer Research</i> , 2015 , 21, 184-92	12.9	63
49	Phenotypic and genotypic characterisation of biallelic mismatch repair deficiency (BMMR-D) syndrome. <i>European Journal of Cancer</i> , 2015 , 51, 977-83	7.5	77
48	A preclinical study demonstrating the efficacy of nilotinib in inhibiting the growth of pediatric high-grade glioma. <i>Journal of Neuro-Oncology</i> , 2015 , 122, 471-80	4.8	4
47	Functionally defined therapeutic targets in diffuse intrinsic pontine glioma. <i>Nature Medicine</i> , 2015 , 21, 555-9	50.5	319
46	Spinal Myxopapillary Ependymomas Demonstrate a Warburg Phenotype. <i>Clinical Cancer Research</i> , 2015 , 21, 3750-8	12.9	35
45	Poly-ADP-Ribose Polymerase as a Therapeutic Target in Pediatric Diffuse Intrinsic Pontine Glioma and Pediatric High-Grade Astrocytoma. <i>Molecular Cancer Therapeutics</i> , 2015 , 14, 2560-8	6.1	37
44	Pathological Findings of a Subependymal Giant Cell Astrocytoma Following Treatment With Rapamycin. <i>Pediatric Neurology</i> , 2015 , 53, 238-242.e1	2.9	4
43	Pathology, Molecular Genetics, and Epigenetics of Diffuse Intrinsic Pontine Glioma. <i>Frontiers in Oncology</i> , 2015 , 5, 147	5.3	67

(2013-2015)

42	Massive CAG repeat expansion and somatic instability in maternally transmitted infantile spinocerebellar ataxia type 7. <i>JAMA Neurology</i> , 2015 , 72, 219-23	17.2	19
41	Study of the biodistribution of fluorescein in glioma-infiltrated mouse brain and histopathological correlation of intraoperative findings in high-grade gliomas resected under fluorescein fluorescence guidance. <i>Journal of Neurosurgery</i> , 2015 , 122, 1360-9	3.2	119
40	BT-02 * FUNCTIONALLY-DEFINED THERAPEUTIC TARGETS IN DIFFUSE INTRINSIC PONTINE GLIOMA. <i>Neuro-Oncology</i> , 2015 , 17, iii3-iii3	1	1
39	Combined hereditary and somatic mutations of replication error repair genes result in rapid onset of ultra-hypermutated cancers. <i>Nature Genetics</i> , 2015 , 47, 257-62	36.3	253
38	Outcome of neurofibromatosis type 1 patients treated with first line vinblastine for optic pathway gliomas: A Canadian multicenter study <i>Journal of Clinical Oncology</i> , 2015 , 33, 2019-2019	2.2	1
37	A microRNA-1280/JAG2 network comprises a novel biological target in high-risk medulloblastoma. <i>Oncotarget</i> , 2015 , 6, 2709-24	3.3	18
36	Genetic alterations in paediatric high grade astrocytomas. <i>Diagnostic Histopathology</i> , 2014 , 20, 84-90	0.7	1
35	Genomic analysis of diffuse intrinsic pontine gliomas identifies three molecular subgroups and recurrent activating ACVR1 mutations. <i>Nature Genetics</i> , 2014 , 46, 451-6	36.3	411
34	Paediatric and adult glioblastoma: multiform (epi)genomic culprits emerge. <i>Nature Reviews Cancer</i> , 2014 , 14, 92-107	31.3	383
33	DETAILED MOLECULAR CHARACTERISATION OF DIFFUSE INTRINSIC PONTINE GLIOMAS IDENTIFIES THREE MOLECULAR SUBGROUPS AND A NOVEL CANCER DRIVER, ACVR1. Neuro-Oncology, 2014, 16, iii26-iii27	1	78
32	International Society Of NeuropathologyHaarlem consensus guidelines for nervous system tumor classification and grading. <i>Brain Pathology</i> , 2014 , 24, 429-35	6	408
31	Genetic and clinical determinants of constitutional mismatch repair deficiency syndrome: report from the constitutional mismatch repair deficiency consortium. <i>European Journal of Cancer</i> , 2014 , 50, 987-96	7.5	149
30	Telomerase inhibition abolishes the tumorigenicity of pediatric ependymoma tumor-initiating cells. <i>Acta Neuropathologica</i> , 2014 , 128, 863-77	14.3	30
29	Histopathological spectrum of paediatric diffuse intrinsic pontine glioma: diagnostic and therapeutic implications. <i>Acta Neuropathologica</i> , 2014 , 128, 573-81	14.3	203
28	ATM regulates 3-methylpurine-DNA glycosylase and promotes therapeutic resistance to alkylating agents. <i>Cancer Discovery</i> , 2014 , 4, 1198-213	24.4	43
27	Loss of p53 cooperates with K-ras activation to induce glioma formation in a region-independent manner. <i>Glia</i> , 2013 , 61, 1862-72	9	15
26	Recurrence patterns across medulloblastoma subgroups: an integrated clinical and molecular analysis. <i>Lancet Oncology, The</i> , 2013 , 14, 1200-7	21.7	226
25	Methylation of the TERT promoter and risk stratification of childhood brain tumours: an integrative genomic and molecular study. <i>Lancet Oncology, The</i> , 2013 , 14, 534-42	21.7	169

24	Differential transformation capacity of neuro-glial progenitors during development. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 14378-83	11.5	8
23	Genomic analysis of diffuse pediatric low-grade gliomas identifies recurrent oncogenic truncating rearrangements in the transcription factor MYBL1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 8188-93	11.5	156
22	Weekly vinblastine in chemotherapy-naive children with unresectable or progressive low grade glioma: A Canadian cooperative study <i>Journal of Clinical Oncology</i> , 2013 , 31, 10029-10029	2.2	2
21	K27M mutation in histone H3.3 defines clinically and biologically distinct subgroups of pediatric diffuse intrinsic pontine gliomas. <i>Acta Neuropathologica</i> , 2012 , 124, 439-47	14.3	629
20	Genome sequencing of pediatric medulloblastoma links catastrophic DNA rearrangements with TP53 mutations. <i>Cell</i> , 2012 , 148, 59-71	56.2	600
19	Phase II study of weekly vinblastine in recurrent or refractory pediatric low-grade glioma. <i>Journal of Clinical Oncology</i> , 2012 , 30, 1358-63	2.2	158
18	Alkylpurine-DNA-N-glycosylase confers resistance to temozolomide in xenograft models of glioblastoma multiforme and is associated with poor survival in patients. <i>Journal of Clinical Investigation</i> , 2012 , 122, 253-66	15.9	108
17	GLI2 is a potential therapeutic target in pediatric medulloblastoma. <i>Journal of Neuropathology and Experimental Neurology</i> , 2011 , 70, 430-7	3.1	17
16	Cribriform neuroepithelial tumour: novel clinicopathological, ultrastructural and cytogenetic findings. <i>Acta Neuropathologica</i> , 2011 , 122, 511-4	14.3	19
15	BRAF-KIAA1549 fusion predicts better clinical outcome in pediatric low-grade astrocytoma. <i>Clinical Cancer Research</i> , 2011 , 17, 4790-8	12.9	178
14	Reply to J.C. Lindsey et al. <i>Journal of Clinical Oncology</i> , 2011 , 29, e347-e347	2.2	1
13	Genetic aberrations leading to MAPK pathway activation mediate oncogene-induced senescence in sporadic pilocytic astrocytomas. <i>Clinical Cancer Research</i> , 2011 , 17, 4650-60	12.9	103
12	A GATA4-regulated tumor suppressor network represses formation of malignant human astrocytomas. <i>Journal of Experimental Medicine</i> , 2011 , 208, 689-702	16.6	65
11	Viruses and human brain tumors: cytomegalovirus enters the fray. <i>Journal of Clinical Investigation</i> , 2011 , 121, 3831-3	15.9	11
10	TP53 alterations determine clinical subgroups and survival of patients with choroid plexus tumors. Journal of Clinical Oncology, 2010 , 28, 1995-2001	2.2	144
9	Whole-genome profiling of pediatric diffuse intrinsic pontine gliomas highlights platelet-derived growth factor receptor alpha and poly (ADP-ribose) polymerase as potential therapeutic targets. Journal of Clinical Oncology, 2010 , 28, 1337-44	2.2	251
8	A Canadian paediatric brain tumour consortium (CPBTC) phase II molecularly targeted study of imatinib in recurrent and refractory paediatric central nervous system tumours. <i>European Journal of Cancer</i> , 2009 , 45, 2352-9	7.5	26
7	Tyrosine kinase expression in pediatric high grade astrocytoma. <i>Journal of Neuro-Oncology</i> , 2008 , 87, 247-53	4.8	47

LIST OF PUBLICATIONS

6	The neuroprotective effects of anti-cd18 therapy after transient global ischemic brain injury in the mouse. <i>Canadian Journal of Anaesthesia</i> , 2006 , 53, 26428-26428	3	
5	Human telomere reverse transcriptase expression predicts progression and survival in pediatric intracranial ependymoma. <i>Journal of Clinical Oncology</i> , 2006 , 24, 1522-8	2.2	100
4	The role of telomere maintenance in the spontaneous growth arrest of pediatric low-grade gliomas. <i>Neoplasia</i> , 2006 , 8, 136-42	6.4	61
3	Identification of human brain tumour initiating cells. <i>Nature</i> , 2004 , 432, 396-401	50.4	5869
2	Identification of human brain tumour initiating cells. <i>Nature</i> , 2004 , 432, 396-401 Identification of a cancer stem cell in human brain tumors. <i>Cancer Research</i> , 2003 , 63, 5821-8	50.4	5869 3368