

Rajeev Vibhakar

List of Publications by Year in descending order

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139
papers

5,924
citations

116194

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#	ARTICLE	IF	CITATIONS
1	Neoplastic and immune single-cell transcriptomics define subgroup-specific intra-tumoral heterogeneity of childhood medulloblastoma. <i>Neuro-Oncology</i> , 2022, 24, 273-286.	0.6	52
2	A novel PLK1 inhibitor onvansertib effectively sensitizes MYC-driven medulloblastoma to radiotherapy. <i>Neuro-Oncology</i> , 2022, 24, 414-426.	0.6	15
3	Single-cell transcriptional analysis of human endothelial colony-forming cells from patients with low VWF levels. <i>Blood</i> , 2022, 139, 2240-2251.	0.6	9
4	Targeting the TP53/MDM2 axis enhances radiation sensitivity in atypical teratoid rhabdoid tumors. <i>International Journal of Oncology</i> , 2022, 60, .	1.4	4
5	SMYD3 Promotes Cell Cycle Progression by Inducing Cyclin D3 Transcription and Stabilizing the Cyclin D1 Protein in Medulloblastoma. <i>Cancers</i> , 2022, 14, 1673.	1.7	5
6	Venetoclax Cooperates with Ionizing Radiation to Attenuate Diffuse Midline Glioma Tumor Growth. <i>Clinical Cancer Research</i> , 2022, 28, 2409-2424.	3.2	6
7	ATRT-23. SIRT2 cooperates with SMARCB1 to induce a differentiation block in ATRT. <i>Neuro-Oncology</i> , 2022, 24, i8-i8.	0.6	0
8	HGG-20. PRMT5 promotes the formation and growth of pediatric high-grade glioma by maintaining tumor stem cell populations. <i>Neuro-Oncology</i> , 2022, 24, i64-i65.	0.6	0
9	DIPG-56. Development and application of a novel antibody against CD99 as a therapeutic strategy in Diffuse Midline Glioma. <i>Neuro-Oncology</i> , 2022, 24, i31-i31.	0.6	0
10	MEDB-80. CDK8 promotes stemness of MYC-driven medulloblastoma. <i>Neuro-Oncology</i> , 2022, 24, i125-i125.	0.6	0
11	IMMU-23. Novel gene-edited CAR-T cell therapy against Diffuse Intrinsic Pontine Glioma. <i>Neuro-Oncology</i> , 2022, 24, i86-i87.	0.6	0
12	RONC-05. Peri-transplant Radiation Therapy for Young Children Treated with High-Dose Chemotherapy for Primary Brain Tumors. <i>Neuro-Oncology</i> , 2022, 24, i177-i177.	0.6	0
13	MEDB-70. Metabolism mediated radiation resistance in MYC-driven Medulloblastoma. <i>Neuro-Oncology</i> , 2022, 24, i122-i123.	0.6	0
14	IMMU-21. Targeting the adenosinergic immune suppression pathway in high grade glioma synergizes with innate immune checkpoint blockade. <i>Neuro-Oncology</i> , 2022, 24, i86-i86.	0.6	0
15	ATRT-10. Single-cell transcriptional profiling of ATRTs reveals heterogeneous signatures of tumor and non-malignant cell populations. <i>Neuro-Oncology</i> , 2022, 24, i4-i5.	0.6	0
16	MEDB-81. Combined inhibition of CDK11 and EZH2 results in regression of MYC-amplified medulloblastoma. <i>Neuro-Oncology</i> , 2022, 24, i125-i125.	0.6	0
17	DIPG-61. Preclinical efficacy of combined radiotherapy with venetoclax treatment in targeting diffuse midline gliomas. <i>Neuro-Oncology</i> , 2022, 24, i32-i33.	0.6	0
18	MEDB-44. Transcriptomic resolution of subgroup-specific medulloblastoma architecture. <i>Neuro-Oncology</i> , 2022, 24, i115-i116.	0.6	0

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19	A Regulatory Loop of FBXW7-MYC-PLK1 Controls Tumorigenesis of MYC-Driven Medulloblastoma. <i>Cancers</i> , 2021, 13, 387.	1.7	11
20	Bromodomain and extra-terminal inhibitors: A consensus prioritisation after the Paediatric Strategy Forum for medicinal product development of epigenetic modifiers in children: ACCELERATE. <i>European Journal of Cancer</i> , 2021, 146, 115-124.	1.3	10
21	NTRK Fusions Can Co-Occur With H3K27M Mutations and May Define Druggable Subclones Within Diffuse Midline Gliomas. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 345-353.	0.9	5
22	The transcriptional landscape of Shh medulloblastoma. <i>Nature Communications</i> , 2021, 12, 1749.	5.8	47
23	Transcriptional control of DNA repair networks by CDK7 regulates sensitivity to radiation in MYC-driven medulloblastoma. <i>Cell Reports</i> , 2021, 35, 109013.	2.9	18
24	Converging evidence for inhibition of transcriptional control in high-grade gliomas. <i>Neuro-Oncology</i> , 2021, 23, 1225-1227.	0.6	1
25	Cryptic developmental events determine medulloblastoma radiosensitivity and cellular heterogeneity without altering transcriptomic profile. <i>Communications Biology</i> , 2021, 4, 616.	2.0	13
26	EMBR-27. NEOPLASTIC AND IMMUNE SINGLE CELL TRANSCRIPTOMICS DEFINE SUBGROUP-SPECIFIC INTRA-TUMORAL HETEROGENEITY OF CHILDHOOD MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2021, 23, i11-i12.	0.6	0
27	RARE-18. NF1-MUTATED TUMORS EXHIBIT INCREASED SENSITIVITY TO AUTOPHAGY INHIBITION ALONE AND IN COMBINATION WITH MEK INHIBITION. <i>Neuro-Oncology</i> , 2021, 23, i44-i44.	0.6	0
28	EMBR-30. A NOVEL PLK1 INHIBITOR ONVANSERTIB EFFECTIVELY SENSITIZES GROUP 3 MEDULLOBLASTOMA TO RADIOTHERAPY. <i>Neuro-Oncology</i> , 2021, 23, i12-i12.	0.6	0
29	HGG-25. PRMT5 PROMOTES TUMOR GROWTH BY MAINTAINING STEMNESS OF PEDIATRIC HIGH-GRADE GLIOMA CELLS. <i>Neuro-Oncology</i> , 2021, 23, i22-i22.	0.6	0
30	HGG-29. VENETOCLAX SYNERGIZES WITH RADIATION THERAPY FOR THE TREATMENT OF DIPG. <i>Neuro-Oncology</i> , 2021, 23, i23-i23.	0.6	0
31	Subgroup and subtype-specific outcomes in adult medulloblastoma. <i>Acta Neuropathologica</i> , 2021, 142, 859-871.	3.9	34
32	Comprehensive molecular characterization of pediatric radiation-induced high-grade glioma. <i>Nature Communications</i> , 2021, 12, 5531.	5.8	31
33	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.	2.7	12
34	The long noncoding RNA <i>lnc-HLX-2-7</i> is oncogenic in Group 3 medulloblastomas. <i>Neuro-Oncology</i> , 2021, 23, 572-585.	0.6	23
35	Exportin 1 Inhibition Induces Nerve Growth Factor Receptor Expression to Inhibit the NF- κ B Pathway in Preclinical Models of Pediatric High-Grade Glioma. <i>Molecular Cancer Therapeutics</i> , 2020, 19, 540-551.	1.9	14
36	Targeted fusion analysis can aid in the classification and treatment of pediatric glioma, ependymoma, and glioneuronal tumors. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28028.	0.8	33

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37	BPTF regulates growth of adult and pediatric high-grade glioma through the MYC pathway. <i>Oncogene</i> , 2020, 39, 2305-2327.	2.6	31
38	Medulloblastoma has a global impact on health related quality of life: Findings from an international cohort. <i>Cancer Medicine</i> , 2020, 9, 447-459.	1.3	11
39	Clinical and molecular characterization of a multi-institutional cohort of pediatric spinal cord low-grade gliomas. <i>Neuro-Oncology Advances</i> , 2020, 2, vdaa103.	0.4	6
40	Senescence Induced by BMI1 Inhibition Is a Therapeutic Vulnerability in H3K27M-Mutant DIPG. <i>Cell Reports</i> , 2020, 33, 108286.	2.9	39
41	Super Elongation Complex as a Targetable Dependency in Diffuse Midline Glioma. <i>Cell Reports</i> , 2020, 31, 107485.	2.9	27
42	Preclinical and clinical investigation of intratumoral chemotherapy pharmacokinetics in DIPG using gemcitabine. <i>Neuro-Oncology Advances</i> , 2020, 2, vdaa021.	0.4	10
43	Proteasome inhibition as a therapeutic approach in atypical teratoid/rhabdoid tumors. <i>Neuro-Oncology Advances</i> , 2020, 2, vdaa051.	0.4	8
44	Single-Cell RNA Sequencing of Childhood Ependymoma Reveals Neoplastic Cell Subpopulations That Impact Molecular Classification and Etiology. <i>Cell Reports</i> , 2020, 32, 108023.	2.9	47
45	The effects of ephrinB2 signaling on proliferation and invasion in glioblastoma multiforme. <i>Molecular Carcinogenesis</i> , 2020, 59, 1064-1075.	1.3	9
46	Targeting MYC-driven replication stress in medulloblastoma with AZD1775 and gemcitabine. <i>Journal of Neuro-Oncology</i> , 2020, 147, 531-545.	1.4	10
47	MiR-1253 exerts tumor-suppressive effects in medulloblastoma via inhibition of CDK6 and CD276 (B7-H3). <i>Brain Pathology</i> , 2020, 30, 732-745.	2.1	35
48	Increased HDAC Activity and c-MYC Expression Mediate Acquired Resistance to WEE1 Inhibition in Acute Leukemia. <i>Frontiers in Oncology</i> , 2020, 10, 296.	1.3	14
49	Advancing biology-based therapeutic approaches for atypical teratoid rhabdoid tumors. <i>Neuro-Oncology</i> , 2020, 22, 944-954.	0.6	25
50	Pattern of Relapse and Treatment Response in WNT-Activated Medulloblastoma. <i>Cell Reports Medicine</i> , 2020, 1, 100038.	3.3	24
51	MBRS-26. CDK7 MEDIATED TRANSCRIPTIONAL PROCESSIVITY OF DNA REPAIR NETWORKS REGULATES SENSITIVITY TO RADIATION IN MYC DRIVEN MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2020, 22, iii403-iii403.	0.6	0
52	MBRS-46. CHARTING NEOPLASTIC AND IMMUNE CELL HETEROGENEITY IN HUMAN AND GEM MODELS OF MEDULLOBLASTOMA USING scRNAseq. <i>Neuro-Oncology</i> , 2020, 22, iii406-iii406.	0.6	0
53	ATRT-24. CELL SURFACE PROTEOME ANALYSIS OF ATRT IDENTIFIES TARGETS FOR IMMUNOTHERAPY. <i>Neuro-Oncology</i> , 2020, 22, iii280-iii280.	0.6	0
54	ATRT-20. CDK7 INHIBITION IN AT/RT. <i>Neuro-Oncology</i> , 2020, 22, iii279-iii280.	0.6	0

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55	EPEN-31. SINGLE-CELL RNAseq OF CHILDHOOD EPENDYMOMA REVEALS DISTINCT NEOPLASTIC CELL SUBPOPULATIONS THAT IMPACT ETIOLOGY, MOLECULAR CLASSIFICATION AND OUTCOME. <i>Neuro-Oncology</i> , 2020, 22, iii314-iii314.	0.6	0
56	IMG-17. RADIOMICS CHARACTERIZATION OF FOUR PEDIATRIC BRAIN TUMOR SUBTYPES IN PDX MOUSE MODELS. <i>Neuro-Oncology</i> , 2020, 22, iii358-iii358.	0.6	0
57	MBRS-65. FBXW7 ACTS A TUMOR SUPPRESSOR IN MYC-DRIVEN MEDULLOBLASTOMA BY CONTROLLING A FEED-FORWARD REGULATORY LOOP OF PLK1 AND MYC. <i>Neuro-Oncology</i> , 2020, 22, iii409-iii409.	0.6	0
58	ATRT-06. SMARCB1 LOSS DRIVEN NON-CANONICAL PRC1 ACTIVITY REGULATES DIFFERENTIATION IN ATYPICAL TERATOID RHABDOID TUMORS (ATRT). <i>Neuro-Oncology</i> , 2020, 22, iii276-iii277.	0.6	0
59	Role of MYC-miR-29-B7-H3 in Medulloblastoma Growth and Angiogenesis. <i>Journal of Clinical Medicine</i> , 2019, 8, 1158.	1.0	30
60	Establishment of patient-derived orthotopic xenograft model of 1q+ posterior fossa group A ependymoma. <i>Neuro-Oncology</i> , 2019, 21, 1540-1551.	0.6	11
61	MEDU-19. EZH2-REGULATED INHIBITION OF HIPK2 SUPPRESSES TREATMENT-INDUCED APOPTOSIS IN GROUP 3 MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2019, 21, ii107-ii107.	0.6	0
62	MEDU-13. FUNCTIONAL CRISPR-CAS9 SCREEN IDENTIFIES DRUGGABLE DEPENDENCIES IN MYC-DRIVEN MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2019, 21, ii105-ii106.	0.6	1
63	Effect of early-stage autophagy inhibition in BRAFV600E autophagy-dependent brain tumor cells. <i>Cell Death and Disease</i> , 2019, 10, 679.	2.7	24
64	DIPG-07. EPIGENOME SCREENING IDENTIFIES TRANSCRIPTIONAL ELONGATION AS THERAPEUTIC VULNERABILITY IN H3K27M-MUTANT DIFFUSE INTRINSIC PONTINE GLIOMA. <i>Neuro-Oncology</i> , 2019, 21, ii69-ii69.	0.6	0
65	DIPG-28. NTRK FUSIONS IN PEDIATRIC DIFFUSE INTRINSIC PONTINE GLIOMAS. <i>Neuro-Oncology</i> , 2019, 21, ii74-ii75.	0.6	0
66	EPEN-09. PRECLINICAL MODELS REVEAL SUBGROUP-STRATIFIED TARGETED THERAPY OPTIONS FOR CHILDHOOD SUPRATENTORIAL EPENDYMOMA. <i>Neuro-Oncology</i> , 2019, 21, ii79-ii79.	0.6	0
67	ATRT-05. PRC1 IS AN ESSENTIAL DEPENDENCY AND THERAPEUTIC TARGET IN SMARCB1 DEFICIENT ATYPICAL TERATOID RHABDOID TUMORS. <i>Neuro-Oncology</i> , 2019, 21, ii63-ii64.	0.6	0
68	scRNA-seq in medulloblastoma shows cellular heterogeneity and lineage expansion support resistance to SHH inhibitor therapy. <i>Nature Communications</i> , 2019, 10, 5829.	5.8	77
69	Recurrent noncoding U1 snRNA mutations drive cryptic splicing in SHH medulloblastoma. <i>Nature</i> , 2019, 574, 707-711.	13.7	129
70	Inhibition of MYC attenuates tumor cell self-renewal and promotes senescence in SMARCB1-deficient Group 2 atypical teratoid rhabdoid tumors to suppress tumor growth in vivo. <i>International Journal of Cancer</i> , 2019, 144, 1983-1995.	2.3	43
71	Combined functional genomic and chemical screens identify SETD8 as a therapeutic target in MYC-driven medulloblastoma. <i>JCI Insight</i> , 2019, 4, .	2.3	20
72	Therapeutic targeting of immune checkpoints with small molecule inhibitors. <i>American Journal of Translational Research (discontinued)</i> , 2019, 11, 529-541.	0.0	9

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73	Preclinical analysis of MTOR complex 1/2 inhibition in diffuse intrinsic pontine glioma. <i>Oncology Reports</i> , 2018, 39, 455-464.	1.2	19
74	EPEN-21. SINGLE CELL RNASEQ IDENTIFIES A PUTATIVE CANCER STEM CELL POPULATION IN POSTERIOR FOSSA EPN. <i>Neuro-Oncology</i> , 2018, 20, i77-i77.	0.6	0
75	DIPG-66. THE H3K27M MUTATION CAUSES WIDE-RANGING CHANGES MEDIATING DIPG TUMORIGENESIS. <i>Neuro-Oncology</i> , 2018, 20, i62-i62.	0.6	0
76	DIPG-77. INTRATUMORAL PHARMACOKINETICS OF CHEMOTHERAPY IN DIPG: XENOGRAFT AND INITIAL PHASE 0 CLINICAL TRIAL RESULTS. <i>Neuro-Oncology</i> , 2018, 20, i64-i65.	0.6	0
77	QOL-58. IMPROVEMENT IN VISUAL ACUITY OF PEDIATRIC PATIENTS WITH BRAIN TUMORS WITH BEVACIZUMAB. <i>Neuro-Oncology</i> , 2018, 20, i169-i169.	0.6	0
78	Exosomal microRNA profiling to identify hypoxia-related biomarkers in prostate cancer. <i>Oncotarget</i> , 2018, 9, 13894-13910.	0.8	47
79	ATRT-18. VALIDATION OF PROTEASOME INHIBITION AS A THERAPEUTIC TARGET IN ATYPICAL TERATOID/RHABDOID TUMORS. <i>Neuro-Oncology</i> , 2018, 20, i31-i31.	0.6	0
80	MBRS-23. EFFECT OF KNOCKDOWN OF KDM6A BY CRISPR/CAS9 EDITING IN MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2018, 20, i133-i133.	0.6	0
81	EPEN-16. PATIENT-DERIVED PFA EPENDYMOMA XENOGRAFT MODEL. <i>Neuro-Oncology</i> , 2018, 20, i76-i76.	0.6	0
82	DIPG-10. THE ONCOGENIC ROLE OF THE SUPER ELONGATION COMPLEX IN H3K27M-MUTANT DIFFUSE MIDLINE GLIOMAS. <i>Neuro-Oncology</i> , 2018, 20, i50-i50.	0.6	0
83	HGG-45. COMPREHENSIVE MOLECULAR CHARACTERIZATION OF PEDIATRIC TREATMENT-INDUCED HIGH-GRADE GLIOMA: GERMLINE DNA REPAIR DEFECTS AS A POTENTIAL ETIOLOGY. <i>Neuro-Oncology</i> , 2018, 20, i98-i98.	0.6	0
84	PHRM-02. DELIVERY OF CHEMOTHERAPEUTICS TO PEDIATRIC BRAIN TUMORS USING CITRATE-CAPPED GOLD NANOPARTICLES. <i>Neuro-Oncology</i> , 2018, 20, i157-i157.	0.6	1
85	PDTM-41. SUPER ELONGATION COMPLEX-MEDIATED TRANSCRIPTIONAL DEPENDENCY IN H3K27M-MUTANT DIFFUSE MIDLINE GLIOMAS. <i>Neuro-Oncology</i> , 2018, 20, vi212-vi212.	0.6	0
86	EPEN-14. SUBGROUP-SPECIFIC THERAPY OPTIONS FOR CHILDHOOD SUPRATENTORIAL EPENDYMOMA. <i>Neuro-Oncology</i> , 2018, 20, i76-i76.	0.6	0
87	MBRS-22. EZH2 OVEREXPRESSION INCREASES THE ONCOGENIC CHARACTER OF CEREBELLAR PROGENITORS AND ISOGRAFTS IN MICE RESULT IN TUMORS RESEMBLING GROUP 3 MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2018, 20, i132-i133.	0.6	0
88	Identification of FDA-Approved Oncology Drugs with Selective Potency in High-Risk Childhood Ependymoma. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 1984-1994.	1.9	19
89	EPEN-15. RETINOIDS AS POTENTIAL CHEMOTHERAPEUTIC OPTIONS FOR POSTERIOR FOSSA EPENDYMOMA OF CHILDHOOD. <i>Neuro-Oncology</i> , 2018, 20, i76-i76.	0.6	0
90	Tumor treating fields in pediatric high-grade glioma. <i>Child's Nervous System</i> , 2017, 33, 1043-1045.	0.6	12

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91	NF- κ B upregulation through epigenetic silencing of LDOC1 drives tumor biology and specific immunophenotype in Group A ependymoma. <i>Neuro-Oncology</i> , 2017, 19, 1350-1360.	0.6	32
92	Loss and E2F/cell cycle deregulation in infant posterior fossa ependymoma. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26656.	0.8	7
93	Intertumoral Heterogeneity within Medulloblastoma Subgroups. <i>Cancer Cell</i> , 2017, 31, 737-754.e6.	7.7	836
94	Combined EphB2 receptor knockdown with radiation decreases cell viability and invasion in medulloblastoma. <i>Cancer Cell International</i> , 2017, 17, 41.	1.8	16
95	A Small-Molecule Inhibitor of WEE1, AZD1775, Synergizes with Olaparib by Impairing Homologous Recombination and Enhancing DNA Damage and Apoptosis in Acute Leukemia. <i>Molecular Cancer Therapeutics</i> , 2017, 16, 2058-2068.	1.9	61
96	Characterization of 2 Novel Ependymoma Cell Lines With Chromosome 1q Gain Derived From Posterior Fossa Tumors of Childhood. <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 595-604.	0.9	19
97	Vincristine and Vinblastine: Is checking bilirubin mandatory in children with Brain Tumors?. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26329.	0.8	1
98	Targeting Polo-like kinase 1 in SMARCB1 deleted atypical teratoid rhabdoid tumor. <i>Oncotarget</i> , 2017, 8, 97290-97303.	0.8	15
99	Autophagy inhibition overcomes multiple mechanisms of resistance to BRAF inhibition in brain tumors. <i>ELife</i> , 2017, 6, .	2.8	128
100	AT-07PLK1 AS A THERAPEUTIC TARGET IN ATRT. <i>Neuro-Oncology</i> , 2016, 18, iii2.2-iii2.	0.6	0
101	HG-78SYNTHETIC LETHAL EPIGENETIC INTERACTIONS IN K27M MUTATED DIPG. <i>Neuro-Oncology</i> , 2016, 18, iii66.3-iii66.	0.6	1
102	MB-71THE ROLE OF WEE1 IN Myc-DRIVEN MEDULLOBLASTOMA. <i>Neuro-Oncology</i> , 2016, 18, iii113.2-iii113.	0.6	2
103	MPS1 kinase as a potential therapeutic target in medulloblastoma. <i>Oncology Reports</i> , 2016, 36, 2633-2640.	1.2	23
104	Polo-like Kinase \hat{A} 1 as a potential therapeutic target in Diffuse Intrinsic Pontine Glioma. <i>BMC Cancer</i> , 2016, 16, 647.	1.1	31
105	Prognostic value of medulloblastoma extent of resection after accounting for molecular subgroup: a retrospective integrated clinical and molecular analysis. <i>Lancet Oncology</i> , The, 2016, 17, 484-495.	5.1	274
106	A WEE1 Inhibitor Analog of AZD1775 Maintains Synergy with Cisplatin and Demonstrates Reduced Single-Agent Cytotoxicity in Medulloblastoma Cells. <i>ACS Chemical Biology</i> , 2016, 11, 921-930.	1.6	42
107	Checkpoint kinase 1 expression is an adverse prognostic marker and therapeutic target in MYC-driven medulloblastoma. <i>Oncotarget</i> , 2016, 7, 53881-53894.	0.8	17
108	EP-04 * ACTIVATION OF THE IL6/STAT3 PATHWAY IN CHILDHOOD EPENDYMOMA IS ASSOCIATED WITH A PRO-INFLAMMATORY TUMOR MICROENVIRONMENT AND A POOR PROGNOSIS. <i>Neuro-Oncology</i> , 2015, 17, iii6-iii6.	0.6	0

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109	Interleukin-6/STAT3 Pathway Signaling Drives an Inflammatory Phenotype in Group A Ependymoma. <i>Cancer Immunology Research</i> , 2015, 3, 1165-1174.	1.6	61
110	Inhibition of BRD4 attenuates tumor cell self-renewal and suppresses stem cell signaling in MYC driven medulloblastoma. <i>Oncotarget</i> , 2014, 5, 2355-2371.	0.8	103
111	Fractionated stereotactic radiosurgery for recurrent ependymoma in children. <i>Journal of Neuro-Oncology</i> , 2014, 116, 107-111.	1.4	45
112	Preclinical high-dose acetaminophen with N-acetylcysteine rescue enhances the efficacy of cisplatin chemotherapy in atypical teratoid rhabdoid tumors. <i>Pediatric Blood and Cancer</i> , 2014, 61, 120-127.	0.8	12
113	Cytogenetic Prognostication Within Medulloblastoma Subgroups. <i>Journal of Clinical Oncology</i> , 2014, 32, 886-896.	0.8	263
114	Integrated genomic analysis identifies the mitotic checkpoint kinase WEE1 as a novel therapeutic target in medulloblastoma. <i>Molecular Cancer</i> , 2014, 13, 72.	7.9	62
115	Epigenetic Regulation of Von Willebrand Factor Gene Expression May Contribute to Von Willebrand Disease Severity. <i>Blood</i> , 2014, 124, 470-470.	0.6	1
116	B7-H3, a potential therapeutic target, is expressed in diffuse intrinsic pontine glioma. <i>Journal of Neuro-Oncology</i> , 2013, 111, 257-264.	1.4	101
117	Aberrant patterns of H3K4 and H3K27 histone lysine methylation occur across subgroups in medulloblastoma. <i>Acta Neuropathologica</i> , 2013, 125, 373-384.	3.9	169
118	Pediatric rhabdoid tumors of kidney and brain show many differences in gene expression but share dysregulation of cell cycle and epigenetic effector genes. <i>Pediatric Blood and Cancer</i> , 2013, 60, 1095-1102.	0.8	40
119	MicroRNA 218 Acts as a Tumor Suppressor by Targeting Multiple Cancer Phenotype-associated Genes in Medulloblastoma. <i>Journal of Biological Chemistry</i> , 2013, 288, 1918-1928.	1.6	100
120	Inhibition of EZH2 suppresses self-renewal and induces radiation sensitivity in atypical rhabdoid teratoid tumor cells. <i>Neuro-Oncology</i> , 2013, 15, 149-160.	0.6	115
121	Inhibition of cyclin-dependent kinase 6 suppresses cell proliferation and enhances radiation sensitivity in medulloblastoma cells. <i>Journal of Neuro-Oncology</i> , 2013, 111, 113-121.	1.4	59
122	TERT promoter mutations are highly recurrent in SHH subgroup medulloblastoma. <i>Acta Neuropathologica</i> , 2013, 126, 917-929.	3.9	146
123	Subgroup-specific structural variation across 1,000 medulloblastoma genomes. <i>Nature</i> , 2012, 488, 49-56.	13.7	761
124	Histone deacetylase inhibition decreases proliferation and potentiates the effect of ionizing radiation in atypical teratoid/rhabdoid tumor cells. <i>Neuro-Oncology</i> , 2012, 14, 175-183.	0.6	51
125	Targeting the enhancer of zeste homologue 2 in medulloblastoma. <i>International Journal of Cancer</i> , 2012, 131, 1800-1809.	2.3	71
126	Targeting Aurora Kinase A enhances radiation sensitivity of atypical teratoid rhabdoid tumor cells. <i>Journal of Neuro-Oncology</i> , 2012, 107, 517-526.	1.4	56

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127	Polo-like kinase 1 (PLK1) inhibition suppresses cell growth and enhances radiation sensitivity in medulloblastoma cells. <i>BMC Cancer</i> , 2012, 12, 80.	1.1	69
128	Survey of MicroRNA expression in pediatric brain tumors. <i>Pediatric Blood and Cancer</i> , 2011, 56, 211-216.	0.8	89
129	High expression of BMP pathway genes distinguishes a subset of atypical teratoid/rhabdoid tumors associated with shorter survival. <i>Neuro-Oncology</i> , 2011, 13, 1296-1307.	0.6	52
130	Inhibition of Aurora Kinase A enhances chemosensitivity of medulloblastoma cell lines. <i>Pediatric Blood and Cancer</i> , 2010, 55, 35-41.	0.8	39
131	MicroRNA 128a Increases Intracellular ROS Level by Targeting Bmi-1 and Inhibits Medulloblastoma Cancer Cell Growth by Promoting Senescence. <i>PLoS ONE</i> , 2010, 5, e10748.	1.1	158
132	Aurora kinase A as a rational target for therapy in glioblastoma. <i>Journal of Neurosurgery: Pediatrics</i> , 2010, 6, 98-105.	0.8	39
133	Regulation of cyclin dependent kinase 6 by microRNA 124 in medulloblastoma. <i>Journal of Neuro-Oncology</i> , 2008, 90, 1-7.	1.4	230
134	Dickkopf-1 is an epigenetically silenced candidate tumor suppressor gene in medulloblastoma1. <i>Neuro-Oncology</i> , 2007, 9, 135-144.	0.6	64
135	Genome-Wide Analysis of Epigenetic Silencing Identifies BEX1 and BEX2 as Candidate Tumor Suppressor Genes in Malignant Glioma. <i>Cancer Research</i> , 2006, 66, 6665-6674.	0.4	135
136	Successful Unrelated Umbilical Cord Blood Transplantation for Shwachman-Diamond Syndrome.. <i>Blood</i> , 2004, 104, 5179-5179.	0.6	0
137	Aberrant T-Cell Antigen Receptor-Mediated Responses in Autoimmune Lymphoproliferative Syndrome. <i>Clinical Immunology</i> , 2002, 104, 31-39.	1.4	15
138	Activation-Induced Expression of Human Programmed Death-1 Gene in T-Lymphocytes. <i>Experimental Cell Research</i> , 1997, 232, 25-28.	1.2	133
139	The human PD-1 gene: complete cDNA, genomic organization, and developmentally regulated expression in B cell progenitors. <i>Gene</i> , 1997, 197, 177-187.	1.0	99