Integrated (epi)-Genomic Analyses Identify Subgroup-S Rhabdoid Tumors

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Citation Report

#	Article	IF	CITATIONS
1	PGBD5 promotes site-specific oncogenic mutations in human tumors. Nature Genetics, 2017, 49, 1005-1014.	9.4	69
2	Sellar Atypical Teratoid/Rhabdoid Tumor (AT/RT). American Journal of Surgical Pathology, 2017, 41, 932-940.	2.1	38
3	Opportunities and challenges in the immunological therapy of pediatric malignancy: a concise snapshot. European Journal of Pediatrics, 2017, 176, 1163-1172.	1.3	11
4	High-Throughput Drug Screening Identifies Pazopanib and Clofilium Tosylate as Promising Treatments for Malignant Rhabdoid Tumors. Cell Reports, 2017, 21, 1737-1745.	2.9	32
5	Integrating RNA sequencing into neuro-oncology practice. Translational Research, 2017, 189, 93-104.	2.2	10
6	SWI/SNF-Komplex-assoziierte Tumordispositions-Syndrome. Medizinische Genetik, 2017, 29, 296-305.	0.1	3
8	Genomic Analysis of Childhood Brain Tumors: Methods for Genome-Wide Discovery and Precision Medicine Become Mainstream. Journal of Clinical Oncology, 2017, 35, 2346-2354.	0.8	25
9	Incorporating Advances in Molecular Pathology Into Brain Tumor Diagnostics. Advances in Anatomic Pathology, 2018, 25, 143-171.	2.4	31
10	Fitting the epigenome into the picture: methylation classification for paediatric brain tumours. Neuropathology and Applied Neurobiology, 2018, 44, 543-547.	1.8	0
11	Primary diffuse leptomeningeal atypical teratoid/rhabdoid tumor diagnosed by cerebrospinal fluid cytology: case report with molecular genetic analysis. Human Pathology, 2018, 77, 116-120.	1.1	7
12	Sellar Region Atypical Teratoid/Rhabdoid Tumors (ATRT) in Adults Display DNA Methylation Profiles of the ATRT-MYC Subgroup. American Journal of Surgical Pathology, 2018, 42, 506-511.	2.1	43
13	Emerging therapeutic targets for the treatment of malignant rhabdoid tumors. Expert Opinion on Therapeutic Targets, 2018, 22, 365-379.	1.5	46
14	<scp>CNS</scp> embryonal tumours: <scp>WHO</scp> 2016 and beyond. Neuropathology and Applied Neurobiology, 2018, 44, 151-162.	1.8	33
15	Improving Diagnostic and Therapeutic Outcomes in Pediatric Brain Tumors. Molecular Diagnosis and Therapy, 2018, 22, 25-39.	1.6	8
16	Review of molecular classification and treatment implications of pediatric brain tumors. Current Opinion in Pediatrics, 2018, 30, 3-9.	1.0	38
17	Malignant Brain Tumours in Children : Present and Future Perspectives. Journal of Korean Neurosurgical Society, 2018, 61, 402-406.	0.5	3
18	Rare Embryonal Brain Tumours. , 2018, , 289-316.		5
19	Advances in the classification of pediatric brain tumors through DNA methylation profiling: From research tool to frontline diagnostic. Cancer, 2018, 124, 4168-4180.	2.0	64

#	Article	IF	CITATIONS
20	Sellar Region Atypical Teratoid/Rhabdoid Tumors in Adults: Clinicopathological Characterization of Five Cases and Review of the Literature. Journal of Neuropathology and Experimental Neurology, 2018, 77, 1115-1121.	0.9	21
21	A biobank of patient-derived pediatric brain tumor models. Nature Medicine, 2018, 24, 1752-1761.	15.2	124
22	Magnetic resonance imaging surrogates of molecular subgroups in atypical teratoid/rhabdoid tumor. Neuro-Oncology, 2018, 20, 1672-1679.	0.6	40
23	Targeting the MTF2–MDM2 Axis Sensitizes Refractory Acute Myeloid Leukemia to Chemotherapy. Cancer Discovery, 2018, 8, 1376-1389.	7.7	40
24	Cerebellar tumors. Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn, 2018, 155, 289-299.	1.0	8
25	Molecular characteristics and therapeutic vulnerabilities across paediatric solid tumours. Nature Reviews Cancer, 2019, 19, 420-438.	12.8	98
26	A C19MC-LIN28A-MYCN Oncogenic Circuit Driven by Hijacked Super-enhancers Is a Distinct Therapeutic Vulnerability in ETMRs: A Lethal Brain Tumor. Cancer Cell, 2019, 36, 51-67.e7.	7.7	69
27	Identification and Analyses of Extra-Cranial and Cranial Rhabdoid Tumor Molecular Subgroups Reveal Tumors with Cytotoxic T Cell Infiltration. Cell Reports, 2019, 29, 2338-2354.e7.	2.9	74
28	Clonally Expanded T Cells Reveal Immunogenicity of Rhabdoid Tumors. Cancer Cell, 2019, 36, 597-612.e8.	7.7	100
29	Small-Molecule and CRISPR Screening Converge to Reveal Receptor Tyrosine Kinase Dependencies in Pediatric Rhabdoid Tumors. Cell Reports, 2019, 28, 2331-2344.e8.	2.9	24
30	NPM1 as a potential therapeutic target for atypical teratoid/rhabdoid tumors. BMC Cancer, 2019, 19, 848.	1.1	13
31	Unbiased Metabolic Profiling Predicts Sensitivity of High MYC-Expressing Atypical Teratoid/Rhabdoid Tumors to Glutamine Inhibition with 6-Diazo-5-Oxo-L-Norleucine. Clinical Cancer Research, 2019, 25, 5925-5936.	3.2	22
32	Dangerous liaisons: interplay between SWI/SNF, NuRD, and Polycomb in chromatin regulation and cancer. Genes and Development, 2019, 33, 936-959.	2.7	127
33	Atypical Teratoid Rhabdoid Tumors. , 2019, , 615-629.		0
34	Molecular pathology of tumors of the central nervous system. Annals of Oncology, 2019, 30, 1265-1278.	0.6	129
35	Tumor Mesenchymal Stromal Cells Regulate Cell Migration of Atypical Teratoid Rhabdoid Tumor through Exosome-Mediated miR155/SMARCA4 Pathway. Cancers, 2019, 11, 720.	1.7	21
36	GenPipes: an open-source framework for distributed and scalable genomic analyses. GigaScience, 2019, 8, .	3.3	121
37	Atypical Teratoid/Rhabdoid Sellar Tumor in an Adult with a Familial History of a Germline SMARCB1 Mutation: Case Report and Review of the Literature. World Neurosurgery, 2019, 127, 336-345.	0.7	12

#	Article	IF	CITATIONS
38	Evaluation of Protein Kinase Inhibitors with PLK4 Cross-Over Potential in a Pre-Clinical Model of Cancer. International Journal of Molecular Sciences, 2019, 20, 2112.	1.8	33
39	Two molecularly distinct atypical teratoid/rhabdoid tumors (or tumor components) occurring in an infant with rhabdoid tumor predisposition syndrome 1. Acta Neuropathologica, 2019, 137, 847-850.	3.9	7
40	Pediatric Atypical Teratoid/Rhabdoid Tumors of the Brain: Identification of Metabolic Subgroups Using In Vivo ¹ H-MR Spectroscopy. American Journal of Neuroradiology, 2019, 40, 872-877.	1.2	6
41	p53 Is a Master Regulator of Proteostasis in SMARCB1-Deficient Malignant Rhabdoid Tumors. Cancer Cell, 2019, 35, 204-220.e9.	7.7	62
42	Stalled developmental programs at the root of pediatric brain tumors. Nature Genetics, 2019, 51, 1702-1713.	9.4	136
43	Inhibition of <i>MYC</i> attenuates tumor cell selfâ€renewal and promotes senescence in SMARCB1â€deficient Group 2 atypical teratoid rhabdoid tumors to suppress tumor growth <i>in vivo</i> . International Journal of Cancer, 2019, 144, 1983-1995.	2.3	43
44	Comprehensive Analysis of Chromatin States in Atypical Teratoid/Rhabdoid Tumor Identifies Diverging Roles for SWI/SNF and Polycomb in Gene Regulation. Cancer Cell, 2019, 35, 95-110.e8.	7.7	65
45	Functional relevance of genes predicted to be affected by epigenetic alterations in atypical teratoid/rhabdoid tumors. Journal of Neuro-Oncology, 2019, 141, 43-55.	1.4	7
46	Prognostic and Predictive Epigenetic Biomarkers in Oncology. Molecular Diagnosis and Therapy, 2019, 23, 83-95.	1.6	57
47	Case-based review: atypical teratoid/rhabdoid tumor. Neuro-Oncology Practice, 2019, 6, 163-178.	1.0	18
48	Tyrosinase immunohistochemistry can be employed for the diagnosis of atypical teratoid/rhabdoid tumours of the tyrosinase subgroup (ATRTâ€TYR). Neuropathology and Applied Neurobiology, 2020, 46, 186-189.	1.8	9
49	Exploiting epigenetic vulnerabilities in solid tumors: Novel therapeutic opportunities in the treatment of SWI/SNF-defective cancers. Seminars in Cancer Biology, 2020, 61, 180-198.	4.3	28
50	Subgroup-specific outcomes of children with malignant childhood brain tumors treated with an irradiation-sparing protocol. Child's Nervous System, 2020, 36, 133-144.	0.6	3
51	MEK/MELK inhibition and blood–brain barrier deficiencies in atypical teratoid/rhabdoid tumors. Neuro-Oncology, 2020, 22, 58-69.	0.6	21
52	Secondary INI1-deficient rhabdoid tumors of the central nervous system: analysis of four cases and literature review. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2020, 476, 763-772.	1.4	8
53	Targeting MEK/MELK in atypical teratoid rhabdoid tumor: a treatment approach aimed at exploiting blood–brain barrier deficiencies. Neuro-Oncology, 2020, 22, 3-4.	0.6	3
54	Translational genomics of malignant rhabdoid tumours: Current impact and future possibilities. Seminars in Cancer Biology, 2020, 61, 30-41.	4.3	17
55	Pediatric embryonal brain tumors in the molecular era. Expert Review of Molecular Diagnostics, 2020, 20, 293-303.	1.5	6

#	Article	IF	CITATIONS
56	SWI/SNF complex heterogeneity is related to polyphenotypic differentiation, prognosis, and immune response in rhabdoid tumors. Neuro-Oncology, 2020, 22, 785-796.	0.6	18
57	Age and DNA methylation subgroup as potential independent risk factors for treatment stratification in children with atypical teratoid/rhabdoid tumors. Neuro-Oncology, 2020, 22, 1006-1017.	0.6	72
58	Outcomes with respect to extent of surgical resection for pediatric atypical teratoid rhabdoid tumors. Child's Nervous System, 2020, 36, 713-719.	0.6	10
59	Desmoplastic myxoid tumor, SMARCB1-mutant: clinical, histopathological and molecular characterization of a pineal region tumor encountered in adolescents and adults. Acta Neuropathologica, 2020, 139, 277-286.	3.9	36
60	Cancer of the Central Nervous System. , 2020, , 906-967.e12.		9
61	Risk-adapted therapy and biological heterogeneity in pineoblastoma: integrated clinico-pathological analysis from the prospective, multi-center SJMB03 and SJYC07 trials. Acta Neuropathologica, 2020, 139, 259-271.	3.9	36
62	Macrophage-tumor cell interaction promotes ATRT progression and chemoresistance. Acta Neuropathologica, 2020, 139, 913-936.	3.9	24
63	Pineoblastoma segregates into molecular sub-groups with distinct clinico-pathologic features: a Rare Brain Tumor Consortium registry study. Acta Neuropathologica, 2020, 139, 223-241.	3.9	65
64	Principles and methods of integrative chromatin analysis in primary tissues and tumors. Biochimica Et Biophysica Acta: Reviews on Cancer, 2020, 1873, 188333.	3.3	7
65	Molecular subgrouping of atypical teratoid/rhabdoid tumors—a reinvestigation and current consensus. Neuro-Oncology, 2020, 22, 613-624.	0.6	133
66	Embryonal tumors of the central nervous system. Current Opinion in Oncology, 2020, 32, 623-630.	1.1	5
67	Atypical teratoid rhabdoid tumor: molecular insights and translation to novel therapeutics. Journal of Neuro-Oncology, 2020, 150, 47-56.	1.4	30
68	Proteasome inhibition as a therapeutic approach in atypical teratoid/rhabdoid tumors. Neuro-Oncology Advances, 2020, 2, vdaa051.	0.4	8
69	Histopathologic and Molecular Features of Central Nervous System Embryonal Tumors for Integrated Diagnosis Reporting. Surgical Pathology Clinics, 2020, 13, 783-800.	0.7	1
71	Pediatric pan-central nervous system tumor analysis of immune-cell infiltration identifies correlates of antitumor immunity. Nature Communications, 2020, 11, 4324.	5.8	75
72	CAR T Cell Therapy for Pediatric Brain Tumors. Frontiers in Oncology, 2020, 10, 1582.	1.3	37
73	SMARCB1 loss interacts with neuronal differentiation state to block maturation and impact cell stability. Genes and Development, 2020, 34, 1316-1329.	2.7	30
74	MEK Inhibition Suppresses Growth of Atypical Teratoid/Rhabdoid Tumors. Journal of Neuropathology and Experimental Neurology, 2020, 79, 746-753.	0.9	4

CITATION REPORT

#	Article	IF	CITATIONS
75	Patient-derived orthotopic xenografts of pediatric brain tumors: a St. Jude resource. Acta Neuropathologica, 2020, 140, 209-225.	3.9	45
76	<scp>SMARCB1</scp> loss induces druggable cyclin <scp>D1</scp> deficiency via upregulation of <scp><i>MIR17HG</i></scp> in atypical teratoid rhabdoid tumors. Journal of Pathology, 2020, 252, 77-87.	2.1	11
77	Clinical characteristics, treatment, and survival outcome in pediatric patients with atypical teratoid/rhabdoid tumors: a retrospective study by the Japan Children's Cancer Group. Journal of Neurosurgery: Pediatrics, 2020, 25, 111-120.	0.8	16
78	Analysis of Dual Class I Histone Deacetylase and Lysine Demethylase Inhibitor Domatinostat (4SC-202) on Growth and Cellular and Genomic Landscape of Atypical Teratoid/Rhabdoid. Cancers, 2020, 12, 756.	1.7	25
79	Advancing biology-based therapeutic approaches for atypical teratoid rhabdoid tumors. Neuro-Oncology, 2020, 22, 944-954.	0.6	25
80	A lncRNA-SWI/SNF complex crosstalk controls transcriptional activation at specific promoter regions. Nature Communications, 2020, 11, 936.	5.8	69
81	Efficacy of High-Dose Chemotherapy and Three-Dimensional Conformal Radiation for Atypical Teratoid/Rhabdoid Tumor: A Report From the Children's Oncology Group Trial ACNS0333. Journal of Clinical Oncology, 2020, 38, 1175-1185.	0.8	102
82	Invited Review: DNA methylationâ€based classification of paediatric brain tumours. Neuropathology and Applied Neurobiology, 2020, 46, 28-47.	1.8	33
83	SWI/SNF complex differences promote cellular heterogeneity in rhabdoid tumors. Neuro-Oncology, 2020, 22, 741-742.	0.6	1
84	Upregulation of Protein Synthesis and Proteasome Degradation Confers Sensitivity to Proteasome Inhibitor Bortezomib in Myc-Atypical Teratoid/Rhabdoid Tumors. Cancers, 2020, 12, 752.	1.7	6
85	Locoregionally administered B7-H3-targeted CAR T cells for treatment of atypical teratoid/rhabdoid tumors. Nature Medicine, 2020, 26, 712-719.	15.2	172
86	The SWI/SNF complex in cancer — biology, biomarkers and therapy. Nature Reviews Clinical Oncology, 2020, 17, 435-448.	12.5	297
87	Invited Review: Dysregulation of chromatin remodellers in paediatric brain tumours – SMARCB1 and beyond. Neuropathology and Applied Neurobiology, 2020, 46, 57-72.	1.8	10
89	Drug screening with a novel tumor-derived cell line identified alternative therapeutic options for patients with atypical teratoid/rhabdoid tumor. Human Cell, 2021, 34, 271-278.	1.2	2
90	Atypical teratoid/rhabdoid tumors (ATRTs) with SMARCA4 mutation are molecularly distinct from SMARCB1-deficient cases. Acta Neuropathologica, 2021, 141, 291-301.	3.9	47
91	Understanding the trajectory of research efforts in atypical teratoid rhabdoid tumors: a bibliometric analysis of the 50 most impactful studies to date. Child's Nervous System, 2021, 37, 419-425.	0.6	3
92	An Adult Case of Sellar Atypical Teratoid/Rhabdoid Tumor Presenting with Lung Metastasis, Harboring a Compound Heterozygous Mutation in INI1. NMC Case Report Journal, 2021, 8, 267-274.	0.2	2
93	DNA methylation profiling as a model for discovery and precision diagnostics in neuro-oncology. Neuro-Oncology, 2021, 23, S16-S29.	0.6	34

CITATION REPORT

#	Article	IF	CITATIONS
94	DIMEimmune: Robust estimation of infiltrating lymphocytes in CNS tumors from DNA methylation profiles. Oncolmmunology, 2021, 10, 1932365.	2.1	17
95	Rhabdoid Tumor Predisposition Syndrome: From Clinical Suspicion to General Management. Frontiers in Oncology, 2021, 11, 586288.	1.3	20
96	Neonatal Central Nervous System Tumors. Clinics in Perinatology, 2021, 48, 35-51.	0.8	3
97	Recent Advances in Understanding the Role of Autophagy in Paediatric Brain Tumours. Diagnostics, 2021, 11, 481.	1.3	5
98	Somatic mutations and single-cell transcriptomes reveal the root of malignant rhabdoid tumours. Nature Communications, 2021, 12, 1407.	5.8	41
99	Histopathological patterns in atypical teratoid/rhabdoid tumors are related to molecular subgroup. Brain Pathology, 2021, 31, e12967.	2.1	16
100	Inhibition of nuclear export restores nuclear localization and residual tumor suppressor function of truncated SMARCB1/INI1 protein in a molecular subset of atypical teratoid/rhabdoid tumors. Acta Neuropathologica, 2021, 142, 361-374.	3.9	6
101	SMARCB1 deletion in atypical teratoid rhabdoid tumors results in human endogenous retrovirus K (HML-2) expression. Scientific Reports, 2021, 11, 12893.	1.6	17
102	Optical genome mapping identifies a germline retrotransposon insertion in <scp><i>SMARCB1</i></scp> in two siblings with atypical teratoid rhabdoid tumors. Journal of Pathology, 2021, 255, 202-211.	2.1	23
103	Pathogenic noncoding variants in the neurofibromatosis and schwannomatosis predisposition genes. Human Mutation, 2021, 42, 1187-1207.	1.1	5
104	Haplotype-resolved germline and somatic alterations in renal medullary carcinomas. Genome Medicine, 2021, 13, 114.	3.6	5
105	Combination Treatment of CI-994 With Etoposide Potentiates Anticancer Effects Through a Topoisomerase II-Dependent Mechanism in Atypical Teratoid/Rhabdoid Tumor (AT/RT). Frontiers in Oncology, 2021, 11, 648023.	1.3	2
106	Clinical evidence for a biological effect of epigenetically active decitabine in relapsed or progressive rhabdoid tumors. Pediatric Blood and Cancer, 2021, 68, e29267.	0.8	7
107	Recent Advances in Pediatric Cancer Research. Cancer Research, 2021, 81, 5783-5799.	0.4	8
108	Polycomb repressive complex 2 in the driver's seat of childhood and young adult brain tumours. Trends in Cell Biology, 2021, 31, 814-828.	3.6	17
109	Pioneer factors in development and cancer. IScience, 2021, 24, 103132.	1.9	15
110	Chromatin accessibility profiling methods. Nature Reviews Methods Primers, 2021, 1, .	11.8	95
113	Advances in the classification and treatment of pediatric brain tumors. Current Opinion in Pediatrics, 2021, 33, 26-32.	1.0	11

		CITATION REI	PORT	
#	Article		IF	CITATIONS
115	Pediatric Brain Tumors. CONTINUUM Lifelong Learning in Neurology, 2017, 23, 1727-175	57.	0.4	19
116	Mithramycin induces promoter reprogramming and differentiation of rhabdoid tumor. EN Molecular Medicine, 2021, 13, e12640.	180	3.3	7
117	Ribavirin as a potential therapeutic for atypical teratoid/rhabdoid tumors. Oncotarget, 20 8054-8067.	118, 9,	0.8	15
118	Atypical Teratoid Rhabdoid Tumour : From Tumours to Therapies. Journal of Korean Neuro Society, 2018, 61, 302-311.	osurgical	0.5	31
119	Renal medullary carcinomas depend upon SMARCB1 loss and are sensitive to proteasom ELife, 2019, 8, .	e inhibition.	2.8	32
120	Endogenous Retroelements and the Viral Mimicry Response in Cancer Therapy and Cellul Homeostasis. Cancer Discovery, 2021, 11, 2707-2725.	ar	7.7	65
121	Atypical Teratoid Rhabdoid Tumours Are Susceptible to Panobinostat-Mediated Different Therapy. Cancers, 2021, 13, 5145.	iation	1.7	3
123	Embryonal Tumors: Atypical Teratoid/Rhabdoid Tumor (ATRT). , 2019, , 1643-1650.			1
124	Embryonale Tumoren. , 2019, , 573-592.			0
127	Rhabdoid Tumor, Soft Tissue. Encyclopedia of Pathology, 2020, , 1-9.		0.0	0
129	Atypical teratoid-rhabdoid tumors: molecular genetics feutures, perspectives of treatmer literature. Russian Journal of Pediatric Hematology and Oncology, 2020, 7, 41-50.	t. Review of	0.1	0
130	Diverse outcomes in extra-cranial rhabdoid tumors: A single institute experience. Pediatri Hematology and Oncology, 2021, , 1-8.	c	0.3	0
131	Embryonal Tumors of the Central Nervous System: The WHO 2016 Classification and Ner Journal of Pediatric Hematology/Oncology, 2021, 43, 79-89.	ν Insights.	0.3	2
132	Novel Two MRT Cell Lines Established from Multiple Sites of a Synchronous MRT Patient. Research, 2020, 40, 6159-6170.	Anticancer	0.5	0
133	The epidemiology of primary and metastatic brain tumors in infancy through childhood. J Neuro-Oncology, 2022, 156, 419-429.	ournal of	1.4	6
135	Atypical teratoid rhabdoid tumor (ATRT): disease mechanisms and potential drug targets Opinion on Therapeutic Targets, 2022, 26, 187-192.	. Expert	1.5	4
136	Evaluation and Diagnosis of Central Nervous System Embryonal Tumors (Non-Medullobla Pediatric and Developmental Pathology, 2022, 25, 34-45.	stoma).	0.5	5
137	Epigenetic mechanisms in paediatric brain tumours: regulators lose control. Biochemical Transactions, 2022, 50, 167-185.	Society	1.6	3

CITATION REPORT

#	Article	IF	CITATIONS
138	Current and Emerging Therapeutic Approaches for Extracranial Malignant Rhabdoid Tumors. Cancer Management and Research, 2022, Volume 14, 479-498.	0.9	11
139	Targeting the TP53/MDM2 axis enhances radiation sensitivity in atypical teratoid rhabdoid tumors. International Journal of Oncology, 2022, 60, .	1.4	4
141	Single-cell transcriptomics identifies potential cells of origin of MYC rhabdoid tumors. Nature Communications, 2022, 13, 1544.	5.8	9
142	Atypical Teratoid Rhabdoid Tumor: A Possible Oriented Female Pathology?. Frontiers in Oncology, 2022, 12, 854437.	1.3	4
143	Therapeutic Targeting of EZH2 and BET BRD4 in Pediatric Rhabdoid Tumors. Molecular Cancer Therapeutics, 2022, 21, 715-726.	1.9	11
145	Comprehensive Genomic Profiling of High-Risk Pediatric Cancer Patients Has a Measurable Impact on Clinical Care. JCO Precision Oncology, 2022, 6, e2100451.	1.5	3
146	ATRT–SHH comprises three molecular subgroups with characteristic clinical and histopathological features and prognostic significance. Acta Neuropathologica, 2022, 143, 697-711.	3.9	13
147	Dual mTORC1/2 inhibition compromises cell defenses against exogenous stress potentiating Obatoclax-induced cytotoxicity in atypical teratoid/rhabdoid tumors. Cell Death and Disease, 2022, 13, 410.	2.7	4
148	Molecular targeted therapies for pediatric atypical teratoid/rhabdoid tumors. Pediatric Investigation, 2022, 6, 111-122.	0.6	3
149	Pediatric CNS cancer genomics and immunogenomics. Current Opinion in Genetics and Development, 2022, 75, 101918.	1.5	0
150	Bromodomain and Extra-Terminal Protein Inhibitors: Biologic Insights and Therapeutic Potential in Pediatric Brain Tumors. Pharmaceuticals, 2022, 15, 665.	1.7	5
151	The HHIP-AS1 lncRNA promotes tumorigenicity through stabilization of dynein complex 1 in human SHH-driven tumors. Nature Communications, 2022, 13, .	5.8	16
152	Molecular Heterogeneity in Pediatric Malignant Rhabdoid Tumors in Patients With Multi-Organ Involvement. Frontiers in Oncology, 0, 12, .	1.3	3
153	SMARCB1-Deficient Cancers: Novel Molecular Insights and Therapeutic Vulnerabilities. Cancers, 2022, 14, 3645.	1.7	18
154	Establishment and characterization of NCC-MRT1-C1: a novel cell line of malignant rhabdoid tumor. Human Cell, 0, , .	1.2	0
155	Primary cilia contribute to the aggressiveness of atypical teratoid/rhabdoid tumors. Cell Death and Disease, 2022, 13, .	2.7	2
157	Dual targeting of EZH1 and EZH2 for the treatment of malignant rhabdoid tumors. Molecular Therapy - Oncolytics, 2022, 27, 14-25.	2.0	10
158	Inherited Genetics Syndromes Associated with Central Nervous System Tumors. , 2022, , .		0

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#	Article	IF	CITATIONS
159	Molecular genetics of paediatric brain tumours and opportunities for precision medicine – a focus on infant tumours. Current Opinion in Neurology, 2022, 35, 772-778.	1.8	1
160	Artificial intelligence platform, RADR®, aids in the discovery of DNA damaging agent for the ultra-rare cancer Atypical Teratoid Rhabdoid Tumors. Frontiers in Drug Discovery, 0, 2, .	1.1	2
161	Pediatric Brain Tumors in the Molecular Era: Updates for the Radiologist. Seminars in Roentgenology, 2023, 58, 47-66.	0.2	3
162	Clustered regularly interspaced short palindromic repeats screens in pediatric tumours: A review. Clinical and Translational Discovery, 2022, 2, .	0.2	0
163	Atypical Teratoid/Rhabdoid Tumor of the Sellar Region in an Adult Male: A Case Report. Cureus, 2023, , .	0.2	0
164	Recent progress and novel approaches to treating atypical teratoid rhabdoid tumor. Neoplasia, 2023, 37, 100880.	2.3	5
165	Pediatric brain tumors: A neuropathologist's approach to the integrated diagnosis. Frontiers in Pediatrics, 0, 11, .	0.9	0
166	Diagnostic classification of childhood cancer using multiscale transcriptomics. Nature Medicine, 2023, 29, 656-666.	15.2	6
167	Generation and multi-dimensional profiling of a childhood cancer cell line atlas defines new therapeutic opportunities. Cancer Cell, 2023, 41, 660-677.e7.	7.7	7
168	Atypical teratoid/rhabdoid tumoroids reveal subgroup-specific drug vulnerabilities. Oncogene, 2023, 42, 1661-1671.	2.6	5
169	Comparative treatment results of children with atypical teratoid/rhabdoid tumor of the central nervous system in the younger age group. Russian Journal of Pediatric Hematology and Oncology, 2023, 10, 11-24.	0.1	1
174	Embryonal Tumors of the Central Nervous System with Multilayered Rosettes and Atypical Teratoid/Rhabdoid Tumors. Advances in Experimental Medicine and Biology, 2023, , 225-252.	0.8	0
177	Medulloblastomas, CNS embryonal tumors, and cerebellar mutism syndrome: advances in care and future directions. Child's Nervous System, 0, , .	0.6	0
184	Bromodomain and extraterminal (BET) proteins: biological functions, diseases, and targeted therapy. Signal Transduction and Targeted Therapy, 2023, 8, .	7.1	7