John M Maris

List of Publications by Year in descending order

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2975 3487 38,196 336 93 182 citations h-index g-index papers 367 367 367 33909 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	<i>DNMT3A</i> overgrowth syndrome is associated with the development of hematopoietic malignancies in children and young adults. Blood, 2022, 139, 461-464.	1.4	9
2	Genomic predictors of response to PD-1 inhibition in children with germline DNA replication repair deficiency. Nature Medicine, 2022, 28, 125-135.	30.7	53
3	GPC2-CAR TÂcells tuned for low antigen density mediate potent activity against neuroblastoma without toxicity. Cancer Cell, 2022, 40, 53-69.e9.	16.8	60
4	Identification of Mitochondrial DNA Variants Associated With Risk of Neuroblastoma. Journal of the National Cancer Institute, 2022, 114, 910-913.	6.3	4
5	Epigenetic state determines inflammatory sensing in neuroblastoma. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, .	7.1	21
6	Genetic analysis in African American children supports ancestry specific neuroblastoma susceptibility. Cancer Epidemiology Biomarkers and Prevention, 2022, , cebp.EPI-21-0782-A.2021.	2.5	1
7	Paraneoplastic myasthenia gravis and pemphigus associated with follicular dendritic cell sarcoma leading to cardiorespiratory collapse in a 7â€yearâ€old. Pediatric Blood and Cancer, 2022, 69, e29723.	1.5	2
8	Abstract LB188: Identification of intrinsic molecular vulnerabilities in inherited and treatment-related hypermutant patient-derived glioma cell line models. Cancer Research, 2022, 82, LB188-LB188.	0.9	O
9	Survival of patients with neuroblastoma before versus after reduction of therapy due to the change in age cut-off from 12 to 18 months in Children's Oncology Group (COG) risk stratification Journal of Clinical Oncology, 2022, 40, 10013-10013.	1.6	O
10	Phase I trial of lorlatinib in combination with topotecan/cyclophosphamide in children with ALK-driven refractory or relapsed neuroblastoma: A new approaches to neuroblastoma therapy consortium study Journal of Clinical Oncology, 2022, 40, 10041-10041.	1.6	0
11	Evaluation of the DLL3-targeting Antibody–Drug Conjugate Rovalpituzumab Tesirine in Preclinical Models of Neuroblastoma. Cancer Research Communications, 2022, 2, 616-623.	1.7	1
12	Outcomes Following GD2-Directed Postconsolidation Therapy for Neuroblastoma After Cessation of Random Assignment on ANBL0032: A Report From the Children's Oncology Group. Journal of Clinical Oncology, 2022, 40, 4107-4118.	1.6	11
13	Refining immunotherapeutic approaches to highâ€risk neuroblastoma based on tumor genomic profiles. Molecular Oncology, 2021, 15, 347-349.	4.6	O
14	PARP Targeted Alpha-Particle Therapy Enhances Response to PD-1 Immune-Checkpoint Blockade in a Syngeneic Mouse Model of Glioblastoma. ACS Pharmacology and Translational Science, 2021, 4, 344-351.	4.9	16
15	Long-Term Follow-up of a Phase III Study of ch14.18 (Dinutuximab) + Cytokine Immunotherapy in Children with High-Risk Neuroblastoma: COG Study ANBL0032. Clinical Cancer Research, 2021, 27, 2179-2189.	7.0	95
16	Image-Guided Biopsy for Relapsed Neuroblastoma: Focus on Safety, Adequacy for Genetic Sequencing, and Correlation of Tumor Cell Percent With Quantitative Lesion MIBG Uptake. JCO Precision Oncology, 2021, 5, 275-285.	3.0	3
17	GAS7 Deficiency Promotes Metastasis in MYCN-Driven Neuroblastoma. Cancer Research, 2021, 81, 2995-3007.	0.9	15
18	Mutations in the RAS/MAPK Pathway Drive Replication Repair–Deficient Hypermutated Tumors and Confer Sensitivity to MEK Inhibition. Cancer Discovery, 2021, 11, 1454-1467.	9.4	19

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19	HACE1 blocks HIF1 \hat{l} ± accumulation under hypoxia in a RAC1 dependent manner. Oncogene, 2021, 40, 1988-2001.	5.9	5
20	The B7-H3–Targeting Antibody–Drug Conjugate m276-SL-PBD Is Potently Effective Against Pediatric Cancer Preclinical Solid Tumor Models. Clinical Cancer Research, 2021, 27, 2938-2946.	7.0	55
21	Epigenetic regulator BMI1 promotes alveolar rhabdomyosarcoma proliferation and constitutes a novel therapeutic target. Molecular Oncology, 2021, 15, 2156-2171.	4.6	11
22	Bromodomain and extra-terminalÂinhibitors—A consensus prioritisation after the Paediatric Strategy Forum for medicinal product development of epigenetic modifiers in children—ACCELERATE. European Journal of Cancer, 2021, 146, 115-124.	2.8	10
23	A G316A Polymorphism in the Ornithine Decarboxylase Gene Promoter Modulates MYCN-Driven Childhood Neuroblastoma. Cancers, 2021, 13, 1807.	3.7	4
24	Testing of B7-H3 targeting antibody-drug conjugate (ADC) MGC018 in models of pediatric solid tumors by the Pediatric Preclinical Testing Consortium (PPTC) Journal of Clinical Oncology, 2021, 39, 10037-10037.	1.6	2
25	IMMU-16. TARGETING GLYPICAN 2 (GPC2) ON PEDIATRIC MALIGNANT BRAIN TUMORS WITH MRNA CAR T CELLS. Neuro-Oncology, 2021, 23, i30-i30.	1.2	1
26	Revised Neuroblastoma Risk Classification System: A Report From the Children's Oncology Group. Journal of Clinical Oncology, 2021, 39, 3229-3241.	1.6	174
27	A GPC2 antibody-drug conjugate is efficacious against neuroblastoma and small-cell lung cancer via binding a conformational epitope. Cell Reports Medicine, 2021, 2, 100344.	6.5	14
28	Abstract 3028: Integrative genomics reveals lncRNAs associated with pediatric cancer. , 2021, , .		1
29	Abstract 1493: Discovery and CAR T targeting of lineage-restricted neuroblastoma oncoproteins. , 2021,		0
30	Randomized Phase II Trial of MIBG Versus MIBG, Vincristine, and Irinotecan Versus MIBG and Vorinostat for Patients With Relapsed or Refractory Neuroblastoma: A Report From NANT Consortium. Journal of Clinical Oncology, 2021, 39, 3506-3514.	1.6	38
31	Drugging the "Undruggable―MYCN Oncogenic Transcription Factor: Overcoming Previous Obstacles to Impact Childhood Cancers. Cancer Research, 2021, 81, 1627-1632.	0.9	25
32	Stage 4S Neuroblastoma. American Journal of Surgical Pathology, 2021, 45, 1075-1081.	3.7	10
33	Retention of CD19 intron 2 contributes to CART-19 resistance in leukemias with subclonal frameshift mutations in CD19. Leukemia, 2020, 34, 1202-1207.	7.2	61
34	PARP-1–Targeted Auger Emitters Display High-LET Cytotoxic Properties In Vitro but Show Limited Therapeutic Utility in Solid Tumor Models of Human Neuroblastoma. Journal of Nuclear Medicine, 2020, 61, 850-856.	5.0	30
35	Pan-neuroblastoma analysis reveals age- and signature-associated driver alterations. Nature Communications, 2020, 11, 5183.	12.8	87
36	SAT-163 Status at 10 Years: Long-Term Follow-Up for a Phase 2a Study of High-Specific-Activity (HSA) I 131 lobenguane in Patients (Pts) with Relapsed/Refractory High-Risk Neuroblastoma. Journal of the Endocrine Society, 2020, 4, .	0.2	2

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37	Somatic structural variation targets neurodevelopmental genes and identifies <i>SHANK2</i> as a tumor suppressor in neuroblastoma. Genome Research, 2020, 30, 1228-1242.	5.5	20
38	annoFuse: an R Package to annotate, prioritize, and interactively explore putative oncogenic RNA fusions. BMC Bioinformatics, 2020, 21, 577.	2.6	4
39	Identification of SARS-CoV-2 Vaccine Epitopes Predicted to Induce Long-Term Population-Scale Immunity. Cell Reports Medicine, 2020, 1, 100036.	6.5	65
40	Immune-Based Approaches for the Treatment of Pediatric Malignancies. Annual Review of Cancer Biology, 2020, 4, 353-370.	4.5	7
41	Crossing Oceans: Preclinical Collaboration to Improve Pediatric Drug Development. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2020, 40, 409-416.	3.8	1
42	Immunogenicity and Immune Silence in Human Cancer. Frontiers in Immunology, 2020, 11, 69.	4.8	22
43	CAMKV Is a Candidate Immunotherapeutic Target in MYCN Amplified Neuroblastoma. Frontiers in Oncology, 2020, 10, 302.	2.8	13
44	Accelerating drug development for neuroblastoma: Summary of the Second Neuroblastoma Drug Development Strategy forum from Innovative Therapies for Children with Cancer and International Society of Paediatric Oncology Europe Neuroblastoma. European Journal of Cancer, 2020, 136, 52-68.	2.8	42
45	Mitochondrial DNA Haplogroups and Susceptibility to Neuroblastoma. Journal of the National Cancer Institute, 2020, 112, 1259-1266.	6.3	10
46	Preclinical evaluation of the combination of AZD1775 and irinotecan against selected pediatric solid tumors: A Pediatric Preclinical Testing Consortium report. Pediatric Blood and Cancer, 2020, 67, e28098.	1.5	13
47	Tattonâ€Brownâ€Rahman syndrome: Six individuals with novel features. American Journal of Medical Genetics, Part A, 2020, 182, 673-680.	1.2	11
48	Rare copy number variants in over 100,000 European ancestry subjects reveal multiple disease associations. Nature Communications, 2020, 11, 255.	12.8	48
49	Phase I Clinical Trial of the Wee1 Inhibitor Adavosertib (AZD1775) with Irinotecan in Children with Relapsed Solid Tumors: A COG Phase I Consortium Report (ADVL1312). Clinical Cancer Research, 2020, 26, 1213-1219.	7.0	38
50	Irinotecan, Temozolomide, and Dinutuximab With GM-CSF in Children With Refractory or Relapsed Neuroblastoma: A Report From the Children's Oncology Group. Journal of Clinical Oncology, 2020, 38, 2160-2169.	1.6	98
51	Locoregional delivery of CAR T cells to the cerebrospinal fluid for treatment of metastatic medulloblastoma and ependymoma. Nature Medicine, 2020, 26, 720-731.	30.7	141
52	LIN28B promotes neuroblastoma metastasis and regulates PDZ binding kinase. Neoplasia, 2020, 22, 231-241.	5.3	21
53	The Human Tumor Atlas Network: Charting Tumor Transitions across Space and Time at Single-Cell Resolution. Cell, 2020, 181, 236-249.	28.9	334
54	Limited antitumor activity of combined BET and MEK inhibition in neuroblastoma. Pediatric Blood and Cancer, 2020, 67, e28267.	1.5	16

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55	High throughput pMHC-I tetramer library production using chaperone-mediated peptide exchange. Nature Communications, 2020, 11, 1909.	12.8	48
56	Epigenomic profiling of neuroblastoma cell lines. Scientific Data, 2020, 7, 116.	5.3	32
57	Telomere Maintenance Mechanisms Define Clinical Outcome in High-Risk Neuroblastoma. Cancer Research, 2020, 80, 2663-2675.	0.9	55
58	Phase I trial of lorlatinib in patients with ALK-driven refractory or relapsed neuroblastoma: A New Approaches to Neuroblastoma Consortium study Journal of Clinical Oncology, 2020, 38, 10504-10504.	1.6	20
59	Clinical significance of serial tumor next generation sequencing (NGS) in 155 pediatric cancer patients Journal of Clinical Oncology, 2020, 38, e13666-e13666.	1.6	1
60	Image-guided core needle biopsy for relapsed and refractory neuroblastoma: A focus on sample adequacy for genetic sequencing Journal of Clinical Oncology, 2020, 38, e22521-e22521.	1.6	1
61	A phase I study of Aurora kinase A inhibitor LY3295668 erbumine as a single agent and in combination in patients with relapsed/refractory neuroblastoma Journal of Clinical Oncology, 2020, 38, TPS10561-TPS10561.	1.6	2
62	Outcomes and toxicities in patients (pts) non-randomly assigned to immunotherapy Children's Oncology Group (COG) ANBL0032 Journal of Clinical Oncology, 2020, 38, 10523-10523.	1.6	0
63	Maintaining Outstanding Outcomes Using Response- and Biology-Based Therapy for Intermediate-Risk Neuroblastoma: A Report From the Children's Oncology Group Study ANBL0531. Journal of Clinical Oncology, 2019, 37, 3243-3255.	1.6	61
64	Neuroblastoma in relation to joint effects of vitamin A and maternal and offspring variants in vitamin A-related genes: A report of the Children's Oncology Group. Cancer Epidemiology, 2019, 61, 165-171.	1.9	6
65	ATRX In-Frame Fusion Neuroblastoma Is Sensitive to EZH2 Inhibition via Modulation of Neuronal Gene Signatures. Cancer Cell, 2019, 36, 512-527.e9.	16.8	44
66	Immunotherapy for pediatric brain tumors: past and present. Neuro-Oncology, 2019, 21, 1226-1238.	1.2	32
67	Genomic Profiling of Childhood Tumor Patient-Derived Xenograft Models to Enable Rational Clinical Trial Design. Cell Reports, 2019, 29, 1675-1689.e9.	6.4	103
68	YAP1 Mediates Resistance to MEK1/2 Inhibition in Neuroblastomas with Hyperactivated RAS Signaling. Cancer Research, 2019, 79, 6204-6214.	0.9	46
69	Antitumor Activity and Tolerability of hu14.18-IL2 with GMCSF and Isotretinoin in Recurrent or Refractory Neuroblastoma: A Children's Oncology Group Phase II Study. Clinical Cancer Research, 2019, 25, 6044-6051.	7.0	20
70	Effect of Tandem Autologous Stem Cell Transplant vs Single Transplant on Event-Free Survival in Patients With High-Risk Neuroblastoma. JAMA - Journal of the American Medical Association, 2019, 322, 746.	7.4	220
71	Exploring Shared Susceptibility between Two Neural Crest Cells Originating Conditions: Neuroblastoma and Congenital Heart Disease. Genes, 2019, 10, 663.	2.4	14
72	Clinical utility of custom-designed NGS panel testing in pediatric tumors. Genome Medicine, 2019, 11, 32.	8.2	79

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73	Targeting PARP-1 with Alpha-Particles Is Potently Cytotoxic to Human Neuroblastoma in Preclinical Models. Molecular Cancer Therapeutics, 2019, 18, 1195-1204.	4.1	36
74	IMMU-04. DEVELOPMENT OF GPC2-DIRECTED CHIMERIC ANTIGEN RECEPTOR THERAPY FOR PEDIATRIC BRAIN TUMORS WITH IN VITRO TRANSCRIBED mRNA. Neuro-Oncology, 2019, 21, ii93-ii93.	1.2	0
75	Outcomes After Proton Therapy for Treatment of Pediatric High-Risk Neuroblastoma. International Journal of Radiation Oncology Biology Physics, 2019, 104, 401-408.	0.8	19
76	Defining Risk Factors for Chemotherapeutic Intervention in Infants With Stage 4S Neuroblastoma: A Report From Children's Oncology Group Study ANBLO531. Journal of Clinical Oncology, 2019, 37, 115-124.	1.6	45
77	PRIMA-1MET-induced neuroblastoma cell death is modulated by p53 and mycn through glutathione level. Journal of Experimental and Clinical Cancer Research, 2019, 38, 69.	8.6	19
78	Dr. Giulio J. D'Angio (1922–2018). Pediatric Research, 2019, 85, 752-753.	2.3	1
79	When Cold Is Hot: Immune Checkpoint Inhibition Therapy for Rhabdoid Tumors. Cancer Cell, 2019, 36, 575-576.	16.8	10
80	Combined innate and adaptive immunotherapy overcomes resistance of immunologically cold syngeneic murine neuroblastoma to checkpoint inhibition., 2019, 7, 344.		45
81	ASCL1 is a MYCN- and LMO1-dependent member of the adrenergic neuroblastoma core regulatory circuitry. Nature Communications, 2019, 10, 5622.	12.8	56
82	Broad Spectrum Activity of the Checkpoint Kinase 1 Inhibitor Prexasertib as a Single Agent or Chemopotentiator Across a Range of Preclinical Pediatric Tumor Models. Clinical Cancer Research, 2019, 25, 2278-2289.	7.0	57
83	The challenge of defining "ultraâ€high―isk―neuroblastoma. Pediatric Blood and Cancer, 2019, 66, e27556.	1.5	43
84	CAR T Cells Targeting B7-H3, a Pan-Cancer Antigen, Demonstrate Potent Preclinical Activity Against Pediatric Solid Tumors and Brain Tumors. Clinical Cancer Research, 2019, 25, 2560-2574.	7.0	369
85	A revised Children's Oncology Group (COG) neuroblastoma risk classification system: Report from the COG biology study ANBLOOB1 Journal of Clinical Oncology, 2019, 37, 10012-10012.	1.6	1
86	Abstract LB-321: Re-evaluating sample sizes in preclinical testing of patient-derived xenografts., 2019,,.		1
87	Cross-Cohort Analysis Identifies a TEAD4–MYCN Positive Feedback Loop as the Core Regulatory Element of High-Risk Neuroblastoma. Cancer Discovery, 2018, 8, 582-599.	9.4	119
88	Genomic Amplifications and Distal 6q Loss: Novel Markers for Poor Survival in High-risk Neuroblastoma Patients. Journal of the National Cancer Institute, 2018, 110, 1084-1093.	6.3	73
89	Pan-cancer genome and transcriptome analyses of 1,699 paediatric leukaemias and solid tumours. Nature, 2018, 555, 371-376.	27.8	649
90	Rare <i> MYC</i> -amplified Neuroblastoma With Large Cell Histology. Pediatric and Developmental Pathology, 2018, 21, 461-466.	1.0	11

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91	QuantumClone: clonal assessment of functional mutations in cancer based on a genotype-aware method for clonal reconstruction. Bioinformatics, 2018, 34, 1808-1816.	4.1	20
92	Neuroblastoma Patients' KIR and KIR-Ligand Genotypes Influence Clinical Outcome for Dinutuximab-based Immunotherapy: A Report from the Children's Oncology Group. Clinical Cancer Research, 2018, 24, 189-196.	7.0	45
93	Intravenous immunoglobulin with prednisone and risk-adapted chemotherapy for children with opsoclonus myoclonus ataxia syndrome associated with neuroblastoma (ANBL00P3): a randomised, open-label, phase 3 trial. The Lancet Child and Adolescent Health, 2018, 2, 25-34.	5.6	38
94	Transverse myelitis as an unexpected complication following treatment with dinutuximab in pediatric patients with highâ€risk neuroblastoma: A case series. Pediatric Blood and Cancer, 2018, 65, e26732.	1.5	21
95	TBIO-29. PedcBioPortal, A CANCER DATA VISUALIZATION TOOL FOR INTEGRATIVE PEDIATRIC CANCER ANALYSES. Neuro-Oncology, 2018, 20, i186-i186.	1.2	0
96	MBRS-57. TARGETING METABOLIC ADAPTATION IN MYC/MYCN AMPLIFIED PEDIATRIC MEDULLOBLASTOMA AND NEUROBLASTOMA. Neuro-Oncology, 2018, 20, i140-i140.	1,2	0
97	Clinically Relevant Cytotoxic Immune Cell Signatures and Clonal Expansion of T-Cell Receptors in High-Risk <i>MYCN</i> -Not-Amplified Human Neuroblastoma. Clinical Cancer Research, 2018, 24, 5673-5684.	7.0	92
98	A Recurrent Mutation in Anaplastic Lymphoma Kinase with Distinct Neoepitope Conformations. Frontiers in Immunology, 2018, 9, 99.	4.8	35
99	A Comprehensive Safety Trial of Chimeric Antibody 14.18 With GM-CSF, IL-2, and Isotretinoin in High-Risk Neuroblastoma Patients Following Myeloablative Therapy: Children's Oncology Group Study ANBL0931. Frontiers in Immunology, 2018, 9, 1355.	4.8	66
100	Phase II Trial of Alisertib in Combination with Irinotecan and Temozolomide for Patients with Relapsed or Refractory Neuroblastoma. Clinical Cancer Research, 2018, 24, 6142-6149.	7.0	55
101	Fine mapping of 2q35 highâ€risk neuroblastoma locus reveals independent functional risk variants and suggests fullâ€length BARD1 as tumorâ€suppressor. International Journal of Cancer, 2018, 143, 2828-2837.	5.1	54
102	Phase II trial of irinotecan/temozolomide/dinutuximab/granulocyte macrophage colony stimulating factor (I/T/DIN/GMCSF) in children with relapsed/refractory neuroblastoma (NBL): A report from the Children's Oncology Group (COG) Journal of Clinical Oncology, 2018, 36, 10508-10508.	1.6	3
103	MYC-family protein overexpression and prominent nucleolar formation represent prognostic indicators and potential therapeutic targets for aggressive high-MKI neuroblastomas: a report from the children's oncology group. Oncotarget, 2018, 9, 6416-6432.	1.8	31
104	Genomic Profiling of a Large Set of Diverse Pediatric Cancers Identifies Known and Novel Mutations across Tumor Spectra. Cancer Research, 2017, 77, 509-519.	0.9	75
105	Association Between Telomere Length and Risk of Cancer and Non-Neoplastic Diseases. JAMA Oncology, 2017, 3, 636.	7.1	376
106	Genetic susceptibility to neuroblastoma. Current Opinion in Genetics and Development, 2017, 42, 81-90.	3.3	75
107	Irinotecan–temozolomide with temsirolimus or dinutuximab in children with refractory or relapsed neuroblastoma (COG ANBL1221): an open-label, randomised, phase 2 trial. Lancet Oncology, The, 2017, 18, 946-957.	10.7	205
108	MYCN amplified neuroblastoma requires the mRNA translation regulator eEF2 kinase to adapt to nutrient deprivation. Cell Death and Differentiation, 2017, 24, 1564-1576.	11.2	24

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109	Assessment of programmed deathâ€ligand 1 expression and tumorâ€associated immune cells in pediatric cancer tissues. Cancer, 2017, 123, 3807-3815.	4.1	135
110	MIBG avidity correlates with clinical features, tumor biology, and outcomes in neuroblastoma: A report from the Children's Oncology Group. Pediatric Blood and Cancer, 2017, 64, e26545.	1.5	30
111	Transcriptomic profiling of 39 commonly-used neuroblastoma cell lines. Scientific Data, 2017, 4, 170033.	5.3	113
112	Dexmedetomidine does not interfere with metaâ€iodobenzylguanidine (MIBG) uptake at clinically relevant concentrations. Pediatric Blood and Cancer, 2017, 64, e26268.	1.5	2
113	Comprehensive Analysis of Hypermutation in Human Cancer. Cell, 2017, 171, 1042-1056.e10.	28.9	596
114	LMO1 Synergizes with MYCN to Promote Neuroblastoma Initiation and Metastasis. Cancer Cell, 2017, 32, 310-323.e5.	16.8	80
115	Common variants in MMP20 at $11q22.2$ predispose to $11q$ deletion and neuroblastoma risk. Nature Communications, 2017, 8, 569.	12.8	22
116	Identification of GPC2 as an Oncoprotein and Candidate Immunotherapeutic Target in High-Risk Neuroblastoma. Cancer Cell, 2017, 32, 295-309.e12.	16.8	148
117	Evaluation of Genetic Predisposition for MYCN-Amplified Neuroblastoma. Journal of the National Cancer Institute, 2017, 109, .	6.3	20
118	11q deletion in neuroblastoma: a review of biological and clinical implications. Molecular Cancer, 2017, 16, 114.	19.2	96
119	Initial testing of VS-4718, a novel inhibitor of focal adhesion kinase (FAK), against pediatric tumor models by the Pediatric Preclinical Testing Program. Pediatric Blood and Cancer, 2017, 64, e26304.	1.5	20
120	Initial testing (stage 1) of the curaxin CBL0137 by the pediatric preclinical testing program. Pediatric Blood and Cancer, 2017, 64, e26263.	1.5	15
121	Preclinical Therapeutic Synergy of MEK1/2 and CDK4/6 Inhibition in Neuroblastoma. Clinical Cancer Research, 2017, 23, 1785-1796.	7.0	66
122	Revisions to the International Neuroblastoma Response Criteria: A Consensus Statement From the National Cancer Institute Clinical Trials Planning Meeting. Journal of Clinical Oncology, 2017, 35, 2580-2587.	1.6	219
123	A phase 1, open-label, dose escalation study of enoblituzumab (MGA271) in pediatric patients with B7-H3-expressing relapsed or refractory solid tumors Journal of Clinical Oncology, 2017, 35, TPS2596-TPS2596.	1.6	13
124	Serum-Based Quantification of MYCN Gene Amplification in Young Patients with Neuroblastoma: Potential Utility as a Surrogate Biomarker for Neuroblastoma. PLoS ONE, 2016, 11, e0161039.	2.5	21
125	Initial Testing (Stage 1) of MKâ€8242—A Novel MDM2 Inhibitor—by the Pediatric Preclinical Testing Program. Pediatric Blood and Cancer, 2016, 63, 1744-1752.	1.5	27
126	A Phase I New Approaches to Neuroblastoma Therapy Study of Buthionine Sulfoximine and Melphalan With Autologous Stem Cells for Recurrent/Refractory High-Risk Neuroblastoma. Pediatric Blood and Cancer, 2016, 63, 1349-1356.	1.5	66

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127	A family-based study of gene variants and maternal folate and choline in neuroblastoma: a report from the Children's Oncology Group. Cancer Causes and Control, 2016, 27, 1209-1218.	1.8	8
128	Advances in the translational genomics of neuroblastoma: From improving risk stratification and revealing novel biology to identifying actionable genomic alterations. Cancer, 2016, 122, 20-33.	4.1	175
129	Incidence and risk factors for secondary malignancy in patients with neuroblastoma after treatment with 131I-metaiodobenzylguanidine. European Journal of Cancer, 2016, 66, 144-152.	2.8	22
130	Evaluation of Alternative <i>In Vivo</i> Drug Screening Methodology: A Single Mouse Analysis. Cancer Research, 2016, 76, 5798-5809.	0.9	52
131	Neuroblastoma. Nature Reviews Disease Primers, 2016, 2, 16078.	30.5	907
132	Pharmacodynamic and genomic markers associated with response to the XPO1/CRM1 inhibitor selinexor (KPTâ€330): A report from the pediatric preclinical testing program. Pediatric Blood and Cancer, 2016, 63, 276-286.	1.5	28
133	Initial Testing of NSC 750854, a Novel Purine Analog, Against Pediatric Tumor Models by the Pediatric Preclinical Testing Program. Pediatric Blood and Cancer, 2016, 63, 443-450.	1.5	0
134	MYCN controls an alternative RNA splicing program in high-risk metastatic neuroblastoma. Cancer Letters, 2016, 371, 214-224.	7.2	46
135	Differential killing of CD56-expressing cells by drug-conjugated human antibodies targeting membrane-distal and membrane-proximal non-overlapping epitopes. MAbs, 2016, 8, 799-810.	5. 2	30
136	Imaging genomics in cancer research: limitations and promises. British Journal of Radiology, 2016, 89, 20151030.	2.2	90
137	Phase I Study of the Aurora A Kinase Inhibitor Alisertib in Combination With Irinotecan and Temozolomide for Patients With Relapsed or Refractory Neuroblastoma: A NANT (New Approaches to) Tj ETQq1	1 0. 78431	4 19В Т /Оve
138	Phase II randomized trial of irinotecan/temozolomide (I/T) with temsirolimus (TEM) or dinutuximab plus granulocyte colony stimulating factor (DIN/GMCSF) in children with refractory or relapsed neuroblastoma: A report from the Children's Oncology Group (COG) Journal of Clinical Oncology, 2016, 34, 10502-10502.	1.6	4
139	Assessment of PD-L1 expression and tumor associated immune cells in pediatric cancer tissues Journal of Clinical Oncology, 2016, 34, 11542-11542.	1.6	3
140	Impact of KIR/KIR ligand genotype for neuroblastoma patients in a phase III COG immunotherapy trial Journal of Clinical Oncology, 2016, 34, e14014-e14014.	1.6	1
141	A phase III randomized clinical trial (RCT) of tandem myeloablative autologous stem cell transplant (ASCT) using peripheral blood stem cell (PBSC) as consolidation therapy for high-risk neuroblastoma (HR-NB): A Children's Oncology Group (COG) study Journal of Clinical Oncology, 2016, 34, LBA3-LBA3.	1.6	17
142	A phase III randomized clinical trial (RCT) of tandem myeloablative autologous stem cell transplant (ASCT) using peripheral blood stem cell (PBSC) as consolidation therapy for high-risk neuroblastoma (HR-NB): A Children's Oncology Group (COG) study Journal of Clinical Oncology, 2016, 34, LBA3-LBA3.	1.6	31
143	Enrichment of Targetable Mutations in the Relapsed Neuroblastoma Genome. PLoS Genetics, 2016, 12, e1006501.	3.5	98
144	Targeting the mTOR Complex by Everolimus in NRAS Mutant Neuroblastoma. PLoS ONE, 2016, 11, e0147682.	2.5	32

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145	Clinical, biologic, and outcome differences according to MIBG avidity in children with neuroblastoma: A report from the Children's Oncology Group (COG) Journal of Clinical Oncology, 2016, 34, 10526-10526.	1.6	O
146	Phase II study of alisertib, irinotecan, and temozolomide in children with relapsed and refractory neuroblastoma: A report from the New Approaches to Neuroblastoma Therapy (NANT) consortium Journal of Clinical Oncology, 2016, 34, 10556-10556.	1.6	0
147	Initial testing (stage 1) of the antiâ€microtubule agents cabazitaxel and docetaxel, by the Pediatric Preclinical Testing Program. Pediatric Blood and Cancer, 2015, 62, 1897-1905.	1.5	14
148	Initial testing (stage 1) of the PARP inhibitor BMN 673 by the pediatric preclinical testing program: <i>PALB2</i> mutation predicts exceptional <i>in vivo</i> response to BMN 673. Pediatric Blood and Cancer, 2015, 62, 91-98.	1.5	65
149	Initial testing (stage 1) of the tubulin binding agent nanoparticle albuminâ€bound (<i>nab</i>) paclitaxel (Abraxane [®]) by the Pediatric Preclinical Testing Program (PPTP). Pediatric Blood and Cancer, 2015, 62, 1214-1221.	1.5	29
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