Kim E Nichols

List of Publications by Year
in descending order

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270 papers	17,262 citations	14655 66 h-index	17105 122 g-index
277 all docs	277 docs citations	277 times ranked	19560 citing authors

#	Article	IF	Citations
1	Clinical and Functional Significance of TP53 Exon 4â€“Intron 4 Splice Junction Variants. Molecular Cancer Research, 2022, 20, 207-216.	3.4	4
2	CPX-351 induces remission in newly diagnosed pediatric secondary myeloid malignancies. Blood Advances, 2022, 6, 521-527.	5.2	10
3	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
4	A Quality Improvement Bundle to Improve Outcomes in Extremely Preterm Infants in the First Week. Pediatrics, 2022, 149, .	2.1	6
5	Development of BRAFV600E-positive acute myeloid leukemia in a patient on long-term dabrafenib for multisystem LCH. Blood Advances, 2022, , .	5.2	5
6	Transient Inhibition of the JAK/STAT Pathway Prevents B-ALL Development in Genetically Predisposed Mice. Cancer Research, 2022, 82, 1098-1109.	0.9	9
7	Managing Pandoraâ€™s Box: Familial Expectations around the Return of (Future) Germline Results. AJOB Empirical Bioethics, 2022, 13, 152-165.	1.6	7
8	Towards the prevention of childhood leukemia. Oncoscience, 2022, 9, 17-19.	2.2	0
9	ATRT-22. Outcomes for children with recurrent atypical teratoid rhabdoid tumor: A single institution study with updated molecular and germline analysis. Neuro-Oncology, 2022, 24, i8-i8.	1.2	1
10	Molecular basis of ETV6-mediated predisposition to childhood acute lymphoblastic leukemia. Blood, 2021, 137, 364-373.	1.4	37
11	Clinical Outcomes and Complications of Pituitary Blastoma. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 351-363.	3.6	23
12	Is neutralization of IFNâ€Î³ sufficient to control inflammation in HLH?. Pediatric Blood and Cancer, 2021, 68, e28886.	1.5	7
13	Creating a cancer genomics curriculum for pediatric hematologyâ€oncology fellows: A national needs assessment. Cancer Medicine, 2021, 10, 2026-2034.	2.8	2
14	Use of the JAK Inhibitor Ruxolitinib in the Treatment of Hemophagocytic Lymphohistiocytosis. Frontiers in Immunology, 2021, 12, 614704.	4.8	77
15	The acquisition of molecular drivers in pediatric therapy-related myeloid neoplasms. Nature Communications, 2021, 12, 985.	12.8	31
16	Relevance of Molecular Groups in Children with Newly Diagnosed Atypical Teratoid Rhabdoid Tumor: Results from Prospective St. Jude Multi-institutional Trials. Clinical Cancer Research, 2021, 27, 2879-2889.	7.0	35
17	Ruxolitinib, a JAK1/2 Inhibitor, Ameliorates Cytokine Storm in Experimental Models of Hyperinflammation Syndrome. Frontiers in Pharmacology, 2021, 12, 650295.	3.5	23
18	HLH or sepsis: the truth is in the T cells. Blood, 2021, 137, 2279-2280.	1.4	1

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19	RHOG: Rac1-ing up another HLH gene. Blood, 2021, 137, 1990-1991.	1.4	1
20	Knowledge Is Power: Benefits, Risks, Hopes, and Decision-Making Reported by Parents Consenting to Next-Generation Sequencing for Children and Adolescents with Cancer. Seminars in Oncology Nursing, 2021, 37, 151167.	1.5	11
21	Genomes for Kids: The Scope of Pathogenic Mutations in Pediatric Cancer Revealed by Comprehensive DNA and RNA Sequencing. Cancer Discovery, 2021, 11, 3008-3027.	9.4	88
22	Hemophagocytic lymphohistiocytosisâ€like toxicity (carHLH) after CD19â€specific CAR Tâ€cell therapy. British Journal of Haematology, 2021, 194, 701-707.	2.5	61
23	Abstract 642: Genomes for Kids: Comprehensive DNA and RNA sequencing defining the scope of actionable mutations in pediatric cancer. , 2021, , .		0
24	Identifying Childhood Cancer Survivors at Increased Genetic Risk for Second Malignant Neoplasms: Use of a Novel Screening Tool. Journal of Clinical Oncology, 2021, 39, JCO.21.01817.	1.6	1
25	Survival Benefit for Individuals With Constitutional Mismatch Repair Deficiency Undergoing Surveillance. Journal of Clinical Oncology, 2021, 39, 2779-2790.	1.6	40
26	Germline RUNX1 variation and predisposition to childhood acute lymphoblastic leukemia. Journal of Clinical Investigation, 2021, 131, .	8.2	20
27	St. Jude Cloud: A Pediatric Cancer Genomic Data-Sharing Ecosystem. Cancer Discovery, 2021, 11, 1082-1099.	9.4	109
28	Cohort Profile: The St. Jude Lifetime Cohort Study (SJLIFE) for paediatric cancer survivors. International Journal of Epidemiology, 2021, 50, 39-49.	1.9	70
29	Performance of the McGill Interactive Pediatric OncoGenetic Guidelines for Identifying Cancer Predisposition Syndromes. JAMA Oncology, 2021, 7, 1806.	7.1	22
30	Analysis of the Li-Fraumeni Spectrum Based on an International Germline TP53 Variant Data Set. JAMA Oncology, 2021, 7, 1800.	7.1	55
31	Liposome-Encapsulated Cytarabine and Daunorubicin (CPX-351) Induces Remission in Newly Diagnosed Pediatric Secondary Myeloid Malignancies. Blood, 2021, 138, 4415-4415.	1.4	0
32	Digenic Inheritance: Evidence and Gaps in Hemophagocytic Lymphohistiocytosis. Frontiers in Immunology, 2021, 12, 777851.	4.8	12
33	Molecular Mechanism of Telomere Length Dynamics and Its Prognostic Value in Pediatric Cancers. Journal of the National Cancer Institute, 2020, 112, 756-764.	6.3	11
34	Recent advances in genetic predisposition to pediatric acute lymphoblastic leukemia. Expert Review of Hematology, 2020, 13, 55-70.	2.2	35
35	Synergistic Signaling of TLR and IFNÎ±/Î² Facilitates Escape of IL-18 Expression from Endotoxin Tolerance. American Journal of Respiratory and Critical Care Medicine, 2020, 201, 526-539.	5.6	38
36	Estimated number of adult survivors of childhood cancer in United States with cancerâ€predisposing germline variants. Pediatric Blood and Cancer, 2020, 67, e28047.	1.5	13

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37	Dasatinib induces a dramatic response in a child with refractory juvenile xanthogranuloma with a novel MRC1-PDGFRB fusion. Blood Advances, 2020, 4, 2991-2995.	5.2	10
38	Combination of NKT14m and Low Dose IL-12 Promotes Invariant Natural Killer T Cell IFN-Î³ Production and Tumor Control. International Journal of Molecular Sciences, 2020, 21, 5085.	4.1	2
39	NK cells: energized yet exhausted in adult HLH. Blood, 2020, 136, 524-525.	1.4	1
40	Cancer surveillance for individuals with Li-Fraumeni syndrome. European Journal of Human Genetics, 2020, 28, 1481-1482.	2.8	7
41	A pedigree-based prediction model identifies carriers of deleterious de novo mutations in families with Li-Fraumeni syndrome. Genome Research, 2020, 30, 1170-1180.	5.5	4
42	Neuroimaging Findings in Children with Constitutional Mismatch Repair Deficiency Syndrome. American Journal of Neuroradiology, 2020, 41, 904-910.	2.4	2
43	Expansion and CD2/CD3/CD28 stimulation enhance Th2 cytokine secretion of human invariant NKT cells with retained anti-tumor cytotoxicity. Cytotherapy, 2020, 22, 276-290.	0.7	7
44	JAK/STAT pathway inhibition sensitizes CD8 T cells to dexamethasone-induced apoptosis in hyperinflammation. Blood, 2020, 136, 657-668.	1.4	50
45	Li-Fraumeni Exploration Consortium Data Coordinating Center: Building an Interactive Web-Based Resource for Collaborative International Cancer Epidemiology Research for a Rare Condition. Cancer Epidemiology Biomarkers and Prevention, 2020, 29, 927-935.	2.5	7
46	Pathogenic Germline Mutations in DNA Repair Genes in Combination With Cancer Treatment Exposures and Risk of Subsequent Neoplasms Among Long-Term Survivors of Childhood Cancer. Journal of Clinical Oncology, 2020, 38, 2728-2740.	1.6	34
47	Recent advances in Wilmsâ€™ tumor predisposition. Human Molecular Genetics, 2020, 29, R138-R149.	2.9	26
48	Cancer Immunotherapeutic Potential of NKTT320, a Novel, Invariant, Natural Killer T Cell-Activating, Humanized Monoclonal Antibody. International Journal of Molecular Sciences, 2020, 21, 4317.	4.1	7
49	Secondary Sarcomas: Biology, Presentation, and Clinical Care. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2020, 40, 463-474.	3.8	4
50	Germline Elongator mutations in Sonic Hedgehog medulloblastoma. Nature, 2020, 580, 396-401.	27.8	94
51	Benign infiltrative myofibroblastic neoplasms of childhood with USP6 gene rearrangement. Histopathology, 2020, 77, 760-768.	2.9	15
52	XAF1 as a modifier of p53 function and cancer susceptibility. Science Advances, 2020, 6, eaba3231.	10.3	37
53	EBV susceptibility. , 2020, , 591-616.		0
54	Evolution of histomorphologic, cytogenetic, and genetic abnormalities in an untreated patient with MIRAGE syndrome. Cancer Genetics, 2020, 245, 42-48.	0.4	7

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55	Mutational Landscape and Patterns of Clonal Evolution in Relapsed Pediatric Acute Lymphoblastic Leukemia. Blood Cancer Discovery, 2020, 1, 96-111.	5.0	93
56	Factors Associated With Declining to Participate in a Pediatric Oncology Next-Generation Sequencing Study. JCO Precision Oncology, 2020, 4, 202-211.	3.0	15
57	Germline Gain-of-Function JAK3 Mutation in Familial Chronic Lymphoproliferative Disorder of NK Cells. Blood, 2020, 136, 9-10.	1.4	9
58	Inborn Errors of Immunity and Cancers. , 2020, , 545-583.		0
59	Translationally Relevant Oral Ruxolitinib Dosing Reduces Inflammation and Ameliorates Disease in Murine Models of Hemophagocytic Lymphohistiocytosis. Blood, 2020, 136, 21-21.	1.4	0
60	Association of Germline BRCA2 Mutations With the Risk of Pediatric or Adolescent Nonâ€“Hodgkin Lymphoma. JAMA Oncology, 2019, 5, 1362.	7.1	19
61	Challenges in the diagnosis of hemophagocytic lymphohistiocytosis: Recommendations from the North American Consortium for Histiocytosis (NACHO). Pediatric Blood and Cancer, 2019, 66, e27929.	1.5	220
62	Enrichment of heterozygous germline RECQL4 loss-of-function variants in pediatric osteosarcoma. Journal of Physical Education and Sports Management, 2019, 5, a004218.	1.2	26
63	Pediatric Cancer Variant Pathogenicity Information Exchange (PeCanPIE): a cloud-based platform for curating and classifying germline variants. Genome Research, 2019, 29, 1555-1565.	5.5	28
64	SLAM-SAP-Fyn: Old Players with New Roles in iNKT Cell Development and Function. International Journal of Molecular Sciences, 2019, 20, 4797.	4.1	2
65	The Immunology of Macrophage Activation Syndrome. Frontiers in Immunology, 2019, 10, 119.	4.8	448
66	Stopping Leukemia in Its Tracks: Should Preemptive Hematopoietic Stem-Cell Transplantation be Offered to Patients at Increased Genetic Risk for Acute Myeloid Leukemia?. Journal of Clinical Oncology, 2019, 37, 2098-2104.	1.6	21
67	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. Neuro-Oncology, 2019, 21, ii96-ii97.	1.2	0
68	Mechanisms of action of ruxolitinib in murine models of hemophagocytic lymphohistiocytosis. Blood, 2019, 134, 147-159.	1.4	99
69	Recommendations for the management of hemophagocytic lymphohistiocytosis in adults. Blood, 2019, 133, 2465-2477.	1.4	587
70	Speaking genomics to parents offered germline testing for cancer predisposition: Use of a 2â€visit consent model. Cancer, 2019, 125, 2455-2464.	4.1	29
71	From uncertainty to pathogenicity: clinical and functional interrogation of a rare TP53 in-frame deletion. Journal of Physical Education and Sports Management, 2019, 5, a003921.	1.2	4
72	Clinical genome sequencing uncovers potentially targetable truncations and fusions of MAP3K8 in spitzoid and other melanomas. Nature Medicine, 2019, 25, 597-602.	30.7	61

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73	Enhancing the antitumor functions of invariant natural killer T cells using a soluble CD1d-CD19 fusion protein. Blood Advances, 2019, 3, 813-824.	5.2	13
74	ClinGen Myeloid Malignancy Variant Curation Expert Panel recommendations for germline RUNX1 variants. Blood Advances, 2019, 3, 2962-2979.	5.2	110
75	Somatic and germline genomics in paediatric acute lymphoblastic leukaemia. Nature Reviews Clinical Oncology, 2019, 16, 227-240.	27.6	132
76	A multicenter study of patients with multisystem Langerhans cell histiocytosis who develop secondary hemophagocytic lymphohistiocytosis. Cancer, 2019, 125, 963-971.	4.1	26
77	Abstract 731: Clinical genome sequencing uncovers potentially targetable truncations and fusions of MAP3K8 in spitzoid and other melanomas. , 2019, , .		1
78	Microbiota-dependent signals are required to sustain TLR-mediated immune responses. JCI Insight, 2019, 4, .	5.0	36
79	Gliomas in the context of Li-Fraumeni syndrome: An international cohort.. Journal of Clinical Oncology, 2019, 37, 1517-1517.	1.6	6
80	Cytokine Storm Syndromes Associated with Epsteinâ€“Barr Virus. , 2019, , 253-276.		2
81	SAT-LB058 Effect of a Genetic Modifier of Cancer Risk in TP53 Mutation Carriers. Journal of the Endocrine Society, 2019, 3, .	0.2	0
82	Real-time sharing of comprehensive clinical genomics sequencing data in St. Jude Cloud.. Journal of Clinical Oncology, 2019, 37, 10019-10019.	1.6	0
83	Abstract 3671: Visualize 10,000 whole-genomes from pediatric cancer patients on St. Jude Cloud. , 2019, , .		0
84	Abstract 4178: Germline mutations inBRCA2and pediatric/adolescent non-Hodgkin's lymphoma: A report from the St. Jude Lifetime (SJLIFE) and Childhood Cancer Survivor Study (CCSS) cohorts. , 2019, , .		0
85	Abstract 3651: Increased prevalence of germline monoallelicRECQL4mutations in children with cancer. , 2019, , .		0
86	The Combination of Dexamethasone and Ruxolitinib Synergistically Attenuates Disease Manifestations in a Preclinical Model of Hemophagocytic Lymphohistiocytosis. Blood, 2019, 134, 81-81.	1.4	1
87	Abstract 2406: A general probabilistic algorithm to predict de novo mutations in familial diseases as demonstrated in Li-Fraumeni Syndrome. , 2019, , .		0
88	Germline Genetic IKZF1 Variation and Predisposition to Childhood Acute Lymphoblastic Leukemia. Cancer Cell, 2018, 33, 937-948.e8.	16.8	142
89	Affinity purification mass spectrometry analysis of PD-1 uncovers SAP as a new checkpoint inhibitor. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E468-E477.	7.1	72
90	Germline Lysine-Specific Demethylase 1 (LSD1/KDM1A) Mutations Confer Susceptibility to Multiple Myeloma. Cancer Research, 2018, 78, 2747-2759.	0.9	56

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91	Impact of professional learning on teachersâ€™ representational strategies and studentsâ€™ cognitive engagement with molecular genetics concepts. Journal of Biological Education, 2018, 52, 31-46.	1.5	3
92	Hemophagocytic Lymphohistiocytosis Associated with Malignancies and with Epstein-Barr Virus. , 2018, , 215-231.		0
93	Genetic Risk for Subsequent Neoplasms Among Long-Term Survivors of Childhood Cancer. Journal of Clinical Oncology, 2018, 36, 2078-2087.	1.6	105
94	TP53 Germline Variations Influence the Predisposition and Prognosis of B-Cell Acute Lymphoblastic Leukemia in Children. Journal of Clinical Oncology, 2018, 36, 591-599.	1.6	121
95	Predisposition Syndromes to Central Nervous System Cancers. , 2018, , 91-116.		0
96	Polygenic Determinants for Subsequent Breast Cancer Risk in Survivors of Childhood Cancer: The St Jude Lifetime Cohort Study (SJLIFE). Clinical Cancer Research, 2018, 24, 6230-6235.	7.0	18
97	Twoâ€yearâ€old female with EBVâ€positive diffuse large Bâ€cell lymphoma and subsequent CNS involvement with neurolymphomatosis. Pediatric Blood and Cancer, 2018, 65, e27415.	1.5	2
98	The genetic basis and cell of origin of mixed phenotype acute leukaemia. Nature, 2018, 562, 373-379.	27.8	236
99	Recommendations for the Use of Etoposide-Based Therapy and Bone Marrow Transplantation for the Treatment of HLH: Consensus Statements by the HLH Steering Committee of the Histiocyte Society. Journal of Allergy and Clinical Immunology: in Practice, 2018, 6, 1508-1517.	3.8	112
100	Parentâ€“child communication surrounding genetic testing for Liâ€“Fraumeni syndrome: Living under the cloud of cancer. Pediatric Blood and Cancer, 2018, 65, e27350.	1.5	15
101	The Influence of Adolescence on Parentsâ€™ Perspectives of Testing and Discussing Inherited Cancer Predisposition. Journal of Genetic Counseling, 2018, 27, 1395-1404.	1.6	6
102	Abstract 922: Access, visualize and analyze 5,000 whole-genomes from pediatric cancer patients on St. Jude Cloud. , 2018, , .		1
103	Abstract 3007: Monogenic and polygenic associations with subsequent breast cancer risk in survivors of childhood cancer: The St. Jude Lifetime Cohort Study (SJLIFE). , 2018, , .		0
104	Myeloid Malignancy Variant Curation Expert Panel: An ASH-Sponsored Clingen Expert Panel to Optimize and Validate Acmg/AMP Variant Interpretation Guidelines for Genes Associated with Inherited Myeloid Neoplasms. Blood, 2018, 132, 5849-5849.	1.4	0
105	Therapeutic Candidate Alpn-101, a Dual ICOS/CD28 Antagonist, Potently Suppresses Human/NSG Mouse Xenograft Graft Vs. Host Disease (GvHD) in a Dose Ranging Study and Reduces Disease Activity in a Mouse Model of Hemophagocytic Lymphohistiocytosis (HLH). Blood, 2018, 132, 2037-2037.	1.4	0
106	Recommended Guidelines for Validation, Quality Control, and Reporting of TP53 Variants in Clinical Practice. Cancer Research, 2017, 77, 1250-1260.	0.9	68
107	Ethical considerations surrounding germline next-generation sequencing of children with cancer. Expert Review of Molecular Diagnostics, 2017, 17, 523-534.	3.1	23
108	Integrating nextâ€generation sequencing into pediatric oncology practice: An assessment of physician confidence and understanding of clinical genomics. Cancer, 2017, 123, 2352-2359.	4.1	58

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109	Clear cell sarcoma of kidney involving a horseshoe kidney and harboring EGFR internal tandem duplication. Pediatric Blood and Cancer, 2017, 64, e26602.	1.5	14
110	Pediatric Cancer Predisposition and Surveillance: An Overview, and a Tribute to Alfred G. Knudson Jr. Clinical Cancer Research, 2017, 23, e1-e5.	7.0	130
111	Recommendations for Surveillance for Children with Leukemia-Predisposing Conditions. Clinical Cancer Research, 2017, 23, e14-e22.	7.0	80
112	Go with the flow: perforin and CD107a in HLH. Blood, 2017, 129, 2954-2955.	1.4	12
113	Should Genetic Testing be Offered for Children? The Perspectives of Adolescents and Emerging Adults in Families with Liâ€Fraumeni Syndrome. Journal of Genetic Counseling, 2017, 26, 1106-1115.	1.6	32
114	Cancer predisposition syndromes associated with myeloid malignancy. Seminars in Hematology, 2017, 54, 115-122.	3.4	8
115	Liâ€Fraumeni syndrome: a paradigm for the understanding of hereditary cancer predisposition. British Journal of Haematology, 2017, 176, 539-552.	2.5	83
116	Development and Initial Validation of the Macrophage Activation Syndrome/Primary Hemophagocytic Lymphohistiocytosis Score, a Diagnostic Tool that Differentiates Primary Hemophagocytic Lymphohistiocytosis from Macrophage Activation Syndrome. Journal of Pediatrics, 2017, 189, 72-78.e3.	1.8	50
117	The Future of Surveillance in the Context of Cancer Predisposition: Through the Murky Looking Glass. Clinical Cancer Research, 2017, 23, e133-e137.	7.0	29
118	The genomic landscape of pediatric myelodysplastic syndromes. Nature Communications, 2017, 8, 1557.	12.8	143
119	Surveillance Recommendations for Children with Overgrowth Syndromes and Predisposition to Wilms Tumors and Hepatoblastoma. Clinical Cancer Research, 2017, 23, e115-e122.	7.0	140
120	Salvage therapy for refractory hemophagocytic lymphohistiocytosis: A review of the published experience. Pediatric Blood and Cancer, 2017, 64, e26308.	1.5	43
121	Abstract 3001: Germline mutations in cancer predisposition genes and risk for subsequent neoplasms among long-term survivors of childhood cancer in the St. Jude Lifetime Cohort. Cancer Research, 2017, 77, 3001-3001.	0.9	2
122	Abstract NG05: TP53-mediated human cancer susceptibility is defined by epigenetic dysregulation of microRNA-34A. , 2017, , .		0
123	The Advantages and Challenges of Testing Children for Heritable Predisposition to Cancer. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2016, 35, 251-269.	3.8	24
124	Metachronous T-Lymphoblastic Lymphoma and Burkitt Lymphoma in a Child With Constitutional Mismatch Repair Deficiency Syndrome. Pediatric Blood and Cancer, 2016, 63, 1454-1456.	1.5	4
125	Stepwise phosphorylation of p65 promotes NF-ÎºB activation and NK cell responses during target cell recognition. Nature Communications, 2016, 7, 11686.	12.8	101
126	Introduction to cancer genetic susceptibility syndromes. Hematology American Society of Hematology Education Program, 2016, 2016, 293-301.	2.5	25

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127	A Novel Collaborative Community-Based Hepatitis B Screening and Linkage to Care Program for African Immigrants. Clinical Infectious Diseases, 2016, 62, S289-S297.	5.8	40
128	Germline Mutations in Predisposition Genes in Pediatric Cancer. New England Journal of Medicine, 2016, 374, 1390-1391.	27.0	20
129	Janus kinase inhibition lessens inflammation and ameliorates disease in murine models of hemophagocytic lymphohistiocytosis. Blood, 2016, 127, 1666-1675.	1.4	207
130	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis: A European League Against Rheumatism/American College of Rheumatology/Paediatric Rheumatology International Trials Organisation Collaborative Initiative. Arthritis and Rheumatology, 2016, 68, 566-576.	5.6	427
131	Genome-Wide DNA Methylation Analysis Reveals Epigenetic Dysregulation of MicroRNA-34A in TP53-Associated Cancer Susceptibility. Journal of Clinical Oncology, 2016, 34, 3697-3704.	1.6	33
132	Primary immunodeficiencies associated with EBV-Induced lymphoproliferative disorders. Critical Reviews in Oncology/Hematology, 2016, 108, 109-127.	4.4	25
133	Unique Familial MLL(KMT2A)â€Rearranged Precursor Bâ€Cell Infant Acute Lymphoblastic Leukemia in Nonâ€twin Siblings. Pediatric Blood and Cancer, 2016, 63, 1175-1180.	1.5	5
134	Expert consensus on dynamics of laboratory tests for diagnosis of macrophage activation syndrome complicating systemic juvenile idiopathic arthritis. RMD Open, 2016, 2, e000161.	3.8	57
135	Inhibition of diacylglycerol kinase Î± restores restimulation-induced cell death and reduces immunopathology in XLP-1. Science Translational Medicine, 2016, 8, 321ra7.	12.4	41
136	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Annals of the Rheumatic Diseases, 2016, 75, 481-489.	0.9	338
137	Reply to â€œWhole-Body MRI Screening in Children With Li-Fraumeni and Other Cancer-Predisposition Syndromes.â€• American Journal of Roentgenology, 2016, 206, W53-W53.	2.2	3
138	A professional learning model that cultivates primary science classroomsâ€™ representational profiles. International Journal of Educational Research, 2016, 76, 12-33.	2.2	4
139	Primary teachersâ€™ representational practices: from competency to fluency. Cambridge Journal of Education, 2016, 46, 509-531.	2.4	4
140	Argumentation-Based Collaborative Inquiry in Science Through Representational Work: Impact on Primary Studentsâ€™ Representational Fluency. Research in Science Education, 2016, 46, 343-364.	2.3	20
141	Comprehensive Functional Characterization of Germline ETV6 Variants Associated with Inherited Predisposition to Acute Lymphoblastic Leukemia in Children. Blood, 2016, 128, 1085-1085.	1.4	1
142	Hemophagocytic Lymphohistiocytosis (HLH) in Langerhans Cell Histiocytosis (LCH): A Multicenter Retrospective Descriptional Study. Blood, 2016, 128, 707-707.	1.4	2
143	Germline Genetic Variation in IKZF1 and Predisposition to Childhood Acute Lymphoblastic Leukemia. Blood, 2016, 128, LBA-2-LBA-2.	1.4	3
144	The Advantages and Challenges of Testing Children for Heritable Predisposition to Cancer. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2016, 36, 251-269.	3.8	10

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145	Combined Treatment with Ruxolitinib and Dexamethasone Curtails Inflammation and Lessens Disease in Preclinical Studies of Hemophagocytic Lymphohistiocytosis. Blood, 2016, 128, 4894-4894.	1.4	1
146	Diagnosis of 9q22.3 microdeletion syndrome in utero following identification of craniosynostosis, overgrowth, and skeletal anomalies. American Journal of Medical Genetics, Part A, 2015, 167, 862-865.	1.2	11
147	Malignancyâ€associated haemophagocytic lymphohistiocytosis in children and adolescents. British Journal of Haematology, 2015, 170, 539-549.	2.5	118
148	Germline ETV6 Mutations Confer Susceptibility to Acute Lymphoblastic Leukemia and Thrombocytopenia. PLoS Genetics, 2015, 11, e1005262.	3.5	128
149	Germline Mutations in Predisposition Genes in Pediatric Cancer. New England Journal of Medicine, 2015, 373, 2336-2346.	27.0	949
150	The effects of scientific representations on primary studentsâ€™ development of scientific discourse and conceptual understandings during cooperative contemporary inquiry-science. Cambridge Journal of Education, 2015, 45, 427-449.	2.4	14
151	Diagnostic Performance of Whole-Body MRI as a Tool for Cancer Screening in Children With Genetic Cancer-Predisposing Conditions. American Journal of Roentgenology, 2015, 205, 400-408.	2.2	82
152	Parent decisionâ€making around the genetic testing of children for germline TP53 mutations. Cancer, 2015, 121, 286-293.	4.1	41
153	Hemophagocytic lymphohistiocytosis caused by dominant-negative mutations in STXBP2 that inhibit SNARE-mediated membrane fusion. Blood, 2015, 125, 1566-1577.	1.4	99
154	Germline genetic variation in ETV6 and risk of childhood acute lymphoblastic leukaemia: a systematic genetic study. Lancet Oncology, The, 2015, 16, 1659-1666.	10.7	161
155	Genotype Versus Phenotype: The Yin and Yang of Germline TP53 Mutations in Li-Fraumeni Syndrome. Journal of Clinical Oncology, 2015, 33, 2331-2333.	1.6	19
156	Abstract 2033: Germline mutations inETV6confer risk of thrombocytopenia and acute lymphocytic leukemia. , 2015, , .		1
157	Genetic Predisposition to Neonatal Tumors. Current Pediatric Reviews, 2015, 11, 164-178.	0.8	3
158	Consensus recommendations for the diagnosis and management of hemophagocytic lymphohistiocytosis associated with malignancies. Haematologica, 2015, 100, 997-1004.	3.5	135
159	Primary Immunodeficiencies and Cancers. , 2015, , 343-375.		0
160	Abstract 4294: Cancer immunotherapeutic potential of NKTT320, a novel human invariant natural killer T-cell activating monoclonal antibody. , 2015, , .		0
161	Germline Genetic Variation in ETV6 and Predisposition to Childhood Acute Lymphoblastic Leukemia. Blood, 2015, 126, 695-695.	1.4	2
162	Predisposition to Pediatric and Hematologic Cancers: A Moving Target. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2014, , e44-e55.	3.8	41

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163	The X-Linked Lymphoproliferative Syndromes. , 2014, , 475-495.		0
164	iNKT Cell Cytotoxic Responses Control T-Lymphoma Growth In Vitro and In Vivo. Cancer Immunology Research, 2014, 2, 59-69.	3.4	60
165	Older Mice Intranasally Sensitized with Aspergillus Fumigatus Develop Stronger Eosinophilic Esophageal Inflammation Compared to Their Younger Counterparts. Journal of Allergy and Clinical Immunology, 2014, 133, AB257.	2.9	0
166	Signaling lymphocytic activation molecule (SLAM)/SLAM-associated protein pathway regulates human B-cell tolerance. Journal of Allergy and Clinical Immunology, 2014, 133, 1149-1161.	2.9	33
167	Primary studentsâ€™ scientific reasoning and discourse during cooperative inquiry-based science activities. International Journal of Educational Research, 2014, 63, 127-140.	2.2	48
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