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	SHP-1 and SHP-2 Associate with Immunoreceptor Tyrosine-Based Switch Motif of Programmed Death 1 upon Primary Human T Cell Stimulation, but Only Receptor Ligation Prevents T Cell Activation. Journal of Immunology, 2004, 173, 945-954.	0.8	989
2	Germline Mutations in Predisposition Genes in Pediatric Cancer. New England Journal of Medicine, 2015, 373, 2336-2346.	27.0	949
3	Recommendations for the management of hemophagocytic lymphohistiocytosis in adults. Blood, 2019, 133, 2465-2477.	1.4	587
4	Cytokine release syndrome after blinatumomab treatment related to abnormal macrophage activation and ameliorated with cytokine-directed therapy. Blood, 2013, 121, 5154-5157.	1.4	524
5	Familial dyserythropoietic anaemia and thrombocytopenia due to an inherited mutation in GATA1. Nature Genetics, 2000, 24, 266-270.	21.4	474
6	Inactivating mutations in an SH2 domain-encoding gene in X-linked lymphoproliferative syndrome. Proceedings of the National Academy of Sciences of the United States of America, 1998, 95, 13765-13770.	7.1	459
7	The Immunology of Macrophage Activation Syndrome. Frontiers in Immunology, 2019, 10, 119.	4.8	448
8	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
9	Regulation of NKT cell development by SAP, the protein defective in XLP. Nature Medicine, 2005, 11, 340-345.	30.7	349
10	2016 Classification Criteria for Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Annals of the Rheumatic Diseases, 2016, 75, 481-489.	0.9	338
11	Mg ²⁺ Regulates Cytotoxic Functions of NK and CD8 T Cells in Chronic EBV Infection Through NKG2D. Science, 2013, 341, 186-191.	12.6	269
12	X-linked lymphoproliferative disease due to SAP/SH2D1A deficiency: a multicenter study on the manifestations, management and outcome of the disease. Blood, 2011, 117, 53-62.	1.4	268
13	Cutting Edge: Functional Requirement for SAP in 2B4-Mediated Activation of Human Natural Killer Cells as Revealed by the X-Linked Lymphoproliferative Syndrome. Journal of Immunology, 2000, 165, 2932-2936.	0.8	245
14	The genetic basis and cell of origin of mixed phenotype acute leukaemia. Nature, 2018, 562, 373-379.	27.8	236
15	Regulation of Cellular and Humoral Immune Responses by the SLAM and SAP Families of Molecules. Annual Review of Immunology, 2007, 25, 337-379.	21.8	229
16	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
17	SAP Regulates TH2 Differentiation and PKC-Î¸-Mediated Activation of NF-Î²B1. Immunity, 2004, 21, 693-706.	14.3	215
18	Janus kinase inhibition lessens inflammation and ameliorates disease in murine models of hemophagocytic lymphohistiocytosis. Blood, 2016, 127, 1666-1675.	1.4	207

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19	CD150 Association with Either the SH2-Containing Inositol Phosphatase or the SH2-Containing Protein Tyrosine Phosphatase Is Regulated by the Adaptor Protein SH2D1A. Journal of Immunology, 2001, 166, 5480-5487.	0.8	201
20	Enhanced T cell responses due to diacylglycerol kinase Î¶ deficiency. Nature Immunology, 2003, 4, 882-890.	14.5	201
21	Molecular and cellular pathogenesis of X-linked lymphoproliferative disease. Immunological Reviews, 2005, 203, 180-199.	6.0	200
22	Treatment of <scp>E</scp>pstein <scp>B</scp>arr virusâ€induced haemophagocytic lymphohistiocytosis with rituximabâ€containing chemoâ€immunotherapeutic regimens. British Journal of Haematology, 2013, 162, 376-382.	2.5	191
23	Expansion of Functionally Immature Transitional B Cells Is Associated with Human-Immunodeficient States Characterized by Impaired Humoral Immunity. Journal of Immunology, 2006, 176, 1506-1516.	0.8	169
24	The EWS-WT1 translocation product induces PDGFA in desmoplastic small round-cell tumour. Nature Genetics, 1997, 17, 309-313.	21.4	166
25	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
26	Outcome of hematopoietic stem cell transplantation for adenosine deaminaseâ€“deficient severe combined immunodeficiency. Blood, 2012, 120, 3615-3624.	1.4	151
27	Hemophagocytic lymphohistiocytosis due to germline mutations inSH2D1A, the X-linked lymphoproliferative disease gene. Blood, 2001, 97, 1131-1133.	1.4	148
28	The genomic landscape of pediatric myelodysplastic syndromes. Nature Communications, 2017, 8, 1557.	12.8	143
29	Germline Genetic IKZF1 Variation and Predisposition to Childhood Acute Lymphoblastic Leukemia. Cancer Cell, 2018, 33, 937-948.e8.	16.8	142
30	Primary immunodeficiency diseases associated with increased susceptibility to viral infections and malignancies. Journal of Allergy and Clinical Immunology, 2011, 127, 1329-1341.e2.	2.9	140
31	Surveillance Recommendations for Children with Overgrowth Syndromes and Predisposition to Wilms Tumors and Hepatoblastoma. Clinical Cancer Research, 2017, 23, e115-e122.	7.0	140
32	Development of Promyelocytic Zinc Finger and ThPOK-Expressing Innate Î³Î´ T Cells Is Controlled by Strength of TCR Signaling and Id3. Journal of Immunology, 2010, 184, 1268-1279.	0.8	139
33	Impaired humoral immunity in X-linked lymphoproliferative disease is associated with defective IL-10 production by CD4+ T cells. Journal of Clinical Investigation, 2005, 115, 1049-1059.	8.2	139
34	Characteristics and Outcomes of Children With the Wilms Tumor-Aniridia Syndrome: A Report From the National Wilms Tumor Study Group. Journal of Clinical Oncology, 2003, 21, 4579-4585.	1.6	136
35	Consensus recommendations for the diagnosis and management of hemophagocytic lymphohistiocytosis associated with malignancies. Haematologica, 2015, 100, 997-1004.	3.5	135
36	Treatment of primary Epstein-Barr virus infection in patients with X-linked lymphoproliferative disease using B-cell-directed therapy. Blood, 2004, 105, 994-996.	1.4	132

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37	Somatic and germline genomics in paediatric acute lymphoblastic leukaemia. Nature Reviews Clinical Oncology, 2019, 16, 227-240.	27.6	132
38	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
39	Germline ETV6 Mutations Confer Susceptibility to Acute Lymphoblastic Leukemia and Thrombocytopenia. PLoS Genetics, 2015, 11, e1005262.	3.5	128
40	Restimulation-induced apoptosis of T cells is impaired in patients with X-linked lymphoproliferative disease caused by SAP deficiency. Journal of Clinical Investigation, 2009, 119, 2976-89.	8.2	126
41	Selective generation of functional somatically mutated IgM+CD27+, but not Ig isotype-switched, memory B cells in X-linked lymphoproliferative disease. Journal of Clinical Investigation, 2006, 116, 322-333.	8.2	122
42	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
43	Malignancyâ€associated haemophagocytic lymphohistiocytosis in children and adolescents. British Journal of Haematology, 2015, 170, 539-549.	2.5	118
44	Familial clustering of Langerhans cell histiocytosis. British Journal of Haematology, 1999, 107, 883-888.	2.5	116
45	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
46	ClinGen Myeloid Malignancy Variant Curation Expert Panel recommendations for germline RUNX1 variants. Blood Advances, 2019, 3, 2962-2979.	5.2	110
47	St. Jude Cloud: A Pediatric Cancer Genomic Data-Sharing Ecosystem. Cancer Discovery, 2021, 11, 1082-1099.	9.4	109
48	Genetic Risk for Subsequent Neoplasms Among Long-Term Survivors of Childhood Cancer. Journal of Clinical Oncology, 2018, 36, 2078-2087.	1.6	105
49	Use of Rituximab in Conjunction With Immunosuppressive Chemotherapy as a Novel Therapy for Epstein Barr Virus-associated Hemophagocytic Lymphohistiocytosis. Journal of Pediatric Hematology/Oncology, 2007, 29, 569-573.	0.6	103
50	Stepwise phosphorylation of p65 promotes NF-ÎºB activation and NK cell responses during target cell recognition. Nature Communications, 2016, 7, 11686.	12.8	101
51	Hemophagocytic lymphohistiocytosis caused by dominant-negative mutations in STXBP2 that inhibit SNARE-mediated membrane fusion. Blood, 2015, 125, 1566-1577.	1.4	99
52	Mechanisms of action of ruxolitinib in murine models of hemophagocytic lymphohistiocytosis. Blood, 2019, 134, 147-159.	1.4	99
53	PLZF Induces the Spontaneous Acquisition of Memory/Effector Functions in T Cells Independently of NKT Cell-Related Signals. Journal of Immunology, 2010, 184, 6746-6755.	0.8	94
54	Germline Elongator mutations in Sonic Hedgehog medulloblastoma. Nature, 2020, 580, 396-401.	27.8	94

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55	Complementation In Trans of Altered ThymocyteÂDevelopment in Mice Expressing MutantÂForms of the Adaptor Molecule SLP76. Immunity, 2008, 28, 359-369.	14.3	93
56	Mutational Landscape and Patterns of Clonal Evolution in Relapsed Pediatric Acute Lymphoblastic Leukemia. Blood Cancer Discovery, 2020, 1, 96-111.	5.0	93
57	Xâ€linked lymphoproliferative syndrome: a genetic condition typified by the triad of infection, immunodeficiency and lymphoma. British Journal of Haematology, 2011, 152, 13-30.	2.5	92
58	Graded repression of PU.1/Sfpi1 gene transcription by GATA factors regulates hematopoietic cell fate. Blood, 2009, 114, 983-994.	1.4	89
59	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
60	Liâ€Fraumeni syndrome: a paradigm for the understanding of hereditary cancer predisposition. British Journal of Haematology, 2017, 176, 539-552.	2.5	83
61	Transcriptional control of invariant NKT cell development. Immunological Reviews, 2010, 238, 195-215.	6.0	82
62	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
63	Impaired humoral immunity in X-linked lymphoproliferative disease is associated with defective IL-10 production by CD4+ T cells. Journal of Clinical Investigation, 2005, 115, 1049-1059.	8.2	81
64	Recommendations for Surveillance for Children with Leukemia-Predisposing Conditions. Clinical Cancer Research, 2017, 23, e14-e22.	7.0	80
65	Use of the JAK Inhibitor Ruxolitinib in the Treatment of Hemophagocytic Lymphohistiocytosis. Frontiers in Immunology, 2021, 12, 614704.	4.8	77
66	Invariant natural killer T cells from children with versus without food allergy exhibit differential responsiveness to milk-derived sphingomyelin. Journal of Allergy and Clinical Immunology, 2011, 128, 102-109.e13.	2.9	75
67	Coincident expression of the chemokine receptors CCR6 and CCR7 by pathologic Langerhans cells in Langerhans cell histiocytosis. Blood, 2003, 101, 2473-2475.	1.4	73
68	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
69	Cohort Profile: The St. Jude Lifetime Cohort Study (SJLIFE) for paediatric cancer survivors. International Journal of Epidemiology, 2021, 50, 39-49.	1.9	70
70	Recommended Guidelines for Validation, Quality Control, and Reporting of TP53 Variants in Clinical Practice. Cancer Research, 2017, 77, 1250-1260.	0.9	68
71	Pathology of the Liver in Familial Hemophagocytic Lymphohistiocytosis. American Journal of Surgical Pathology, 2010, 34, 852-867.	3.7	64
72	Functional Requirements for Interactions Between CD84 and Src Homology 2 Domain-Containing Proteins and Their Contribution to Human T Cell Activation. Journal of Immunology, 2003, 171, 2485-2495.	0.8	63

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73	Successful use of the antiâ€CD25 antibody daclizumab in an adult patient with hemophagocytic lymphohistiocytosis. American Journal of Hematology, 2008, 83, 747-749.	4.1	63
74	Clinical features of three girls with mosaic genomeâ€wide paternal uniparental isodisomy. American Journal of Medical Genetics, Part A, 2013, 161, 1929-1939.	1.2	63
75	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
76	Hemophagocytic lymphohistiocytosisâ€like toxicity (carHLH) after CD19â€specific CAR Tâ€cell therapy. British Journal of Haematology, 2021, 194, 701-707.	2.5	61
77	Diacylglycerol kinase Î¶ regulates microbial recognition and host resistance to Toxoplasma gondii. Journal of Experimental Medicine, 2007, 204, 781-792.	8.5	60
78	iNKT Cell Cytotoxic Responses Control T-Lymphoma Growth In Vitro and In Vivo. Cancer Immunology Research, 2014, 2, 59-69.	3.4	60
79	Sensitive multistep clinical molecular screening of 180 unrelated individuals with retinoblastoma detects 36 novel mutations in theRB1 gene. Human Mutation, 2005, 25, 566-574.	2.5	59
80	Expansion of somatically reverted memory CD8+ T cells in patients with X-linked lymphoproliferative disease caused by selective pressure from Epstein-Barr virus. Journal of Experimental Medicine, 2012, 209, 913-924.	8.5	59
81	Heterozygous Germline ATM Mutations Do Not Contribute to Radiation-Associated Malignancies After Hodgkin's Disease. Journal of Clinical Oncology, 1999, 17, 1259-1259.	1.6	58
82	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
83	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
84	Germline Lysine-Specific Demethylase 1 (LSD1/KDM1A) Mutations Confer Susceptibility to Multiple Myeloma. Cancer Research, 2018, 78, 2747-2759.	0.9	56
85	Analysis of the Li-Fraumeni Spectrum Based on an International Germline TP53 Variant Data Set. JAMA Oncology, 2021, 7, 1800.	7.1	55
86	Loss of circulating CD27+ memory B cells and CCR4+ T cells occurring in association with elevated EBV loads in XLP patients surviving primary EBV infection. Blood, 2004, 103, 1625-1631.	1.4	53
87	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
88	Educational paper. European Journal of Pediatrics, 2011, 170, 285-294.	2.7	50
89	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
90	JAK/STAT pathway inhibition sensitizes CD8 T cells to dexamethasone-induced apoptosis in hyperinflammation. Blood, 2020, 136, 657-668.	1.4	50

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91	Central nervous system juvenile xanthogranuloma with malignant transformation. Pediatric Blood and Cancer, 2008, 50, 927-930.	1.5	49
92	Primary studentsâ€™ scientific reasoning and discourse during cooperative inquiry-based science activities. International Journal of Educational Research, 2014, 63, 127-140.	2.2	48
93	Differential Requirement for the SAP-Fyn Interaction during NK T Cell Development and Function. Journal of Immunology, 2008, 181, 2311-2320.	0.8	46
94	Representational Classroom Practices that Contribute to Studentsâ€™ Conceptual and Representational Understanding of Chemical Bonding. International Journal of Science Education, 2011, 33, 2215-2246.	1.9	45
95	SAP-Mediated Inhibition of Diacylglycerol Kinase Î± Regulates TCR-Induced Diacylglycerol Signaling. Journal of Immunology, 2011, 187, 5941-5951.	0.8	43
96	Salvage therapy for refractory hemophagocytic lymphohistiocytosis: A review of the published experience. Pediatric Blood and Cancer, 2017, 64, e26308.	1.5	43
97	Distinct Interactions of the X-Linked Lymphoproliferative Syndrome Gene Product SAP with Cytoplasmic Domains of Members of the CD2 Receptor Family. Clinical Immunology, 2001, 100, 15-23.	3.2	41
98	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
99	Parent decisionâ€making around the genetic testing of children for germline TP53 mutations. Cancer, 2015, 121, 286-293.	4.1	41
100	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
101	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
102	Survival Benefit for Individuals With Constitutional Mismatch Repair Deficiency Undergoing Surveillance. Journal of Clinical Oncology, 2021, 39, 2779-2790.	1.6	40
103	Genetic testing and tumor surveillance for children with cancer predisposition syndromes. Current Opinion in Pediatrics, 2008, 20, 1-7.	2.0	39
104	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
105	The adaptor protein SH2D1A regulates signaling through CD150 (SLAM) in B cells. Blood, 2004, 104, 4063-4070.	1.4	37
106	XAF1 as a modifier of p53 function and cancer susceptibility. Science Advances, 2020, 6, eaba3231.	10.3	37
107	Molecular basis of ETV6-mediated predisposition to childhood acute lymphoblastic leukemia. Blood, 2021, 137, 364-373.	1.4	37
108	Strategies to reduce medication errors with reference to older adults. International Journal of Evidence-Based Healthcare, 2006, 4, 2-41.	0.5	36

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109	Impaired SLAM-SLAM Homotypic Interaction between Invariant NKT Cells and Dendritic Cells Affects Differentiation of IL-4/IL-10-Secreting NKT2 Cells in Nonobese Diabetic Mice. Journal of Immunology, 2008, 181, 869-877.	0.8	36
110	A Common Molecular Mechanism Underlies Two Phenotypically Distinct 17p13.1 Microdeletion Syndromes. American Journal of Human Genetics, 2010, 87, 631-642.	6.2	36
111	Microbiota-dependent signals are required to sustain TLR-mediated immune responses. JCI Insight, 2019, 4, .	5.0	36
112	Recent advances in genetic predisposition to pediatric acute lymphoblastic leukemia. Expert Review of Hematology, 2020, 13, 55-70.	2.2	35
113	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
114	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
115	Prevalence of germline truncating mutations inATM in women with a second breast cancer after radiation therapy for a contralateral tumor. , 2000, 27, 124-129.		33
116	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
117	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
118	Lymphocytic vasculitis involving the central nervous system occurs in patients with Xâ€linked lymphoproliferative disease in the absence of Epsteinâ€“Barr virus infection. Pediatric Blood and Cancer, 2009, 53, 1120-1123.	1.5	32
119	Recent advances in retinoblastoma genetic research. Current Opinion in Ophthalmology, 2009, 20, 351-355.	2.9	32
120	Murine natural killer immunoreceptors use distinct proximal signaling complexes to direct cell function. Blood, 2013, 121, 3135-3146.	1.4	32
121	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
122	TP53, BRCA1, and BRCA2 Tumor Suppressor Genes Are Not Commonly Mutated in Survivors of Hodgkinâ€™s Disease With Second Primary Neoplasms. Journal of Clinical Oncology, 2003, 21, 4505-4509.	1.6	31
123	The acquisition of molecular drivers in pediatric therapy-related myeloid neoplasms. Nature Communications, 2021, 12, 985.	12.8	31
124	X-linked lymphoproliferative disease (XLP): a model of impaired anti-viral, anti-tumor and humoral immune responses. Immunologic Research, 2008, 42, 145-159.	2.9	29
125	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
126	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

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127	The adaptor molecule SAP plays essential roles during invariant NKT cell cytotoxicity and lytic synapse formation. Blood, 2013, 121, 3386-3395.	1.4	28
128	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
129	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
130	A multicenter study of patients with multisystem Langerhans cell histiocytosis who develop secondary hemophagocytic lymphohistiocytosis. Cancer, 2019, 125, 963-971.	4.1	26
131	Recent advances in Wilmsâ€™ tumor predisposition. Human Molecular Genetics, 2020, 29, R138-R149.	2.9	26
132	Loss of SLP-76 Expression within Myeloid Cells Confers Resistance to Neutrophil-Mediated Tissue Damage while Maintaining Effective Bacterial Killing. Journal of Immunology, 2007, 178, 4606-4614.	0.8	25
133	Introduction to cancer genetic susceptibility syndromes. Hematology American Society of Hematology Education Program, 2016, 2016, 293-301.	2.5	25
134	Primary immunodeficiencies associated with EBV-Induced lymphoproliferative disorders. Critical Reviews in Oncology/Hematology, 2016, 108, 109-127.	4.4	25
135	Identification, Management, and Evaluation of Children with Cancer-Predisposition Syndromes. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2012, , 576-584.	3.8	24
136	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
137	The Parallels Between Philosophical Inquiry and Scientific Inquiry: Implications for science education. Educational Philosophy and Theory, 2012, 44, 1045-1059.	1.8	23
138	Ethical considerations surrounding germline next-generation sequencing of children with cancer. Expert Review of Molecular Diagnostics, 2017, 17, 523-534.	3.1	23
139	Clinical Outcomes and Complications of Pituitary Blastoma. Journal of Clinical Endocrinology and Metabolism, 2021, 106, 351-363.	3.6	23
140	Ruxolitinib, a JAK1/2 Inhibitor, Ameliorates Cytokine Storm in Experimental Models of Hyperinflammation Syndrome. Frontiers in Pharmacology, 2021, 12, 650295.	3.5	23
141	The Adaptor Molecule Signaling Lymphocytic Activation Molecule-Associated Protein (SAP) Regulates IFN-Î³ and IL-4 Production in Vî±14 Transgenic NKT Cells via Effects on GATA-3 and T-bet Expression. Journal of Immunology, 2009, 182, 1370-1378.	0.8	22
142	Performance of the McGill Interactive Pediatric OncoGenetic Guidelines for Identifying Cancer Predisposition Syndromes. JAMA Oncology, 2021, 7, 1806.	7.1	22
143	Childhoood cancer predisposition: Applications of molecular testing and future implications. Journal of Pediatrics, 1998, 132, 389-397.	1.8	21
144	A novel CD93 polymorphism in non-obese diabetic (NOD) and NZB/W F1 mice is linked to a CD4+ iNKT cell deficient state. Immunogenetics, 2010, 62, 397-407.	2.4	21

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145	Medical radiation exposure and risk of retinoblastoma resulting from new germline RB1 mutation. International Journal of Cancer, 2011, 128, 2393-2404.	5.1	21
146	Bilateral Pheochromocytomas, Hemihyperplasia, and Subtle Somatic Mosaicism: The Importance of Detecting Lowâ€‹scp›L‹/scp›evel Uniparental Disomy. American Journal of Medical Genetics, Part A, 2013, 161, 993-1001.	1.2	21
147	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
148	Invariant NKT cells. OncoImmunology, 2013, 2, e27440.	4.6	20
149	Germline Mutations in Predisposition Genes in Pediatric Cancer. New England Journal of Medicine, 2016, 374, 1390-1391.	27.0	20
150	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
151	Germline RUNX1 variation and predisposition to childhood acute lymphoblastic leukemia. Journal of Clinical Investigation, 2021, 131, .	8.2	20
152	Genotype Versus Phenotype: The Yin and Yang of Germline ‹i›TP53‹/i› Mutations in Li-Fraumeni Syndrome. Journal of Clinical Oncology, 2015, 33, 2331-2333.	1.6	19
153	Association of Germline ‹i›BRCA‹/i›2 Mutations With the Risk of Pediatric or Adolescent Nonâ€“Hodgkin Lymphoma. JAMA Oncology, 2019, 5, 1362.	7.1	19
154	SAP: natural inhibitor or grand SLAM of T cell activation?. Nature Immunology, 2001, 2, 665-666.	14.5	18
155	BRAF, a piece of the LCH puzzle. Blood, 2010, 116, 1825-1827.	1.4	18
156	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
157	Macrophage activation and FcÎ³ receptor-mediated signaling do not require expression of the SLP-76 and SLP-65 adaptors. Journal of Leukocyte Biology, 2004, 75, 541-552.	3.3	16
158	Mutation of tyrosine 145 of lymphocyte cytosolic protein 2 protects mice from anaphylaxis and arthritis. Journal of Allergy and Clinical Immunology, 2009, 124, 1088-1098.	2.9	16
159	MOLECULAR KARYOTYPE OF SPORADIC UNILATERAL RETINOBLASTOMA TUMORS. Retina, 2009, 29, 1002-1012.	1.7	16
160	Physical Map and Cosmid Contig Encompassing a New Interstitial Deletion of the X-Linked LymphoprolifÃ©rative Syndrome Region. European Journal of Human Genetics, 1996, 4, 342-351.	2.8	16
161	Missense mutations in SH2D1A identified in patients with X-linked lymphoproliferative disease differentially affect the expression and function of SAP. International Immunology, 2006, 18, 1055-1065.	4.0	15
162	Transforming the Social Practices of Learning with Representations: A Study of Disciplinary Discourse. Research in Science Education, 2013, 43, 179-208.	2.3	15

#	Article	IF	Citations
163	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
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