Susan L Slager

List of Publications by Year
in descending order

430 papers	19,666 citations	11608 70 h-index	14702 127 g-index
432 all docs	432 docs citations	432 times ranked	24074 citing authors

#	Article	IF	Citations
1	Familial Primary Pulmonary Hypertension (Gene PPH1) Is Caused by Mutations in the Bone Morphogenetic Protein Receptorâ€“II Gene. American Journal of Human Genetics, 2000, 67, 737-744.	2.6	1,089
2	Large-scale genotyping identifies 41 new loci associated with breast cancer risk. Nature Genetics, 2013, 45, 353-361.	9.4	960
3	Discovery and prioritization of somatic mutations in diffuse large B-cell lymphoma (DLBCL) by whole-exome sequencing. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 3879-3884.	3.3	853
4	Polygenic Risk Scores for Prediction of Breast Cancer and Breast Cancer Subtypes. American Journal of Human Genetics, 2019, 104, 21-34.	2.6	711
5	Inherited Mutations in 17 Breast Cancer Susceptibility Genes Among a Large Triple-Negative Breast Cancer Cohort Unselected for Family History of Breast Cancer. Journal of Clinical Oncology, 2015, 33, 304-311.	0.8	521
6	Genome-wide association analysis of more than 120,000 individuals identifies 15 new susceptibility loci for breast cancer. Nature Genetics, 2015, 47, 373-380.	9.4	513
7	Multiple independent variants at the TERT locus are associated with telomere length and risks of breast and ovarian cancer. Nature Genetics, 2013, 45, 371-384.	9.4	493
8	Genome-wide association studies identify four ER negativeâ€“specific breast cancer risk loci. Nature Genetics, 2013, 45, 392-398.	9.4	374
9	A Genomewide Association Study of Citalopram Response in Major Depressive Disorder. Biological Psychiatry, 2010, 67, 133-138.	0.7	289
10	A common variant at the TERT-CLPTM1L locus is associated with estrogen receptorâ€“negative breast cancer. Nature Genetics, 2011, 43, 1210-1214.	9.4	279
11	Etiologic Heterogeneity Among Non-Hodgkin Lymphoma Subtypes: The InterLymph Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 130-144.	0.9	265
12	Mutations in CHEK2 Associated with Prostate Cancer Risk. American Journal of Human Genetics, 2003, 72, 270-280.	2.6	264
13	Rates and Outcomes of Follicular Lymphoma Transformation in the Immunochemotherapy Era: A Report From the University of Iowa/Mayo Clinic Specialized Program of Research Excellence Molecular Epidemiology Resource. Journal of Clinical Oncology, 2013, 31, 3272-3278.	0.8	259
14	Genome-Wide Association Study in BRCA1 Mutation Carriers Identifies Novel Loci Associated with Breast and Ovarian Cancer Risk. PLoS Genetics, 2013, 9, e1003212.	1.5	244
15	Sequence Analysis of the Serotonin Transporter and Associations with Antidepressant Response. Biological Psychiatry, 2005, 58, 374-381.	0.7	203
16	Diffuse large <scp>B</scp>â€cell lymphoma (<scp>R</scp>ichter syndrome) in patients with chronic lymphocytic leukaemia (CLL): a cohort study of newly diagnosed patients. British Journal of Haematology, 2013, 162, 774-782.	1.2	187
17	Vitamin D Insufficiency and Prognosis in Non-Hodgkin's Lymphoma. Journal of Clinical Oncology, 2010, 28, 4191-4198.	0.8	184
18	Early event status informs subsequent outcome in newly diagnosed follicular lymphoma. American Journal of Hematology, 2016, 91, 1096-1101.	2.0	180

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19	Genome-wide association study identifies multiple risk loci for chronic lymphocytic leukemia. Nature Genetics, 2013, 45, 868-876.	9.4	179
20	Comorbid conditions and survival in unselected, newly diagnosed patients with chronic lymphocytic leukemia. Leukemia and Lymphoma, 2008, 49, 49-56.	0.6	176
21	PALB2,CHEK2andATMrare variants and cancer risk: data from COGS. Journal of Medical Genetics, 2016, 53, 800-811.	1.5	174
22	A meta-analysis of genome-wide association studies of breast cancer identifies two novel susceptibility loci at 6q14 and 20q11. Human Molecular Genetics, 2012, 21, 5373-5384.	1.4	168
23	Family history of hematopoietic malignancies and risk of non-Hodgkin lymphoma (NHL): a pooled analysis of 10â€‰211 cases and 11â€‰905 controls from the International Lymphoma Epidemiology Consortium (InterLymph). Blood, 2007, 109, 3479-3488.	0.6	159
24	Case-Control Studies of Genetic Markers: Power and Sample Size Approximations for Armitageâ€™s Test for Trend. Human Heredity, 2001, 52, 149-153.	0.4	154
25	Genome-wide association study of follicular lymphoma identifies a risk locus at 6p21.32. Nature Genetics, 2010, 42, 661-664.	9.4	152
26	Analysis of Association Between the Serotonin Transporter and Antidepressant Response in a Large Clinical Sample. Biological Psychiatry, 2007, 61, 734-742.	0.7	148
27	Genome-wide association study identifies multiple susceptibility loci for diffuse large B cell lymphoma. Nature Genetics, 2014, 46, 1233-1238.	9.4	147
28	Genome-wide association study identifies 25 known breast cancer susceptibility loci as risk factors for triple-negative breast cancer. Carcinogenesis, 2014, 35, 1012-1019.	1.3	145
29	Evidence for Genetic Linkage Between a Polymorphism in the Adenosine 2A Receptor and Panic Disorder. Neuropsychopharmacology, 2004, 29, 558-565.	2.8	144
30	Genetic variation in 1253 immune and inflammation genes and risk of non-Hodgkin lymphoma. Blood, 2007, 110, 4455-4463.	0.6	144
31	Pharmacokinetic Genes Do Not Influence Response or Tolerance to Citalopram in the STAR*D Sample. PLoS ONE, 2008, 3, e1872.	1.1	144
32	Cause of Death in Follicular Lymphoma in the First Decade of the Rituximab Era: A Pooled Analysis of French and US Cohorts. Journal of Clinical Oncology, 2019, 37, 144-152.	0.8	142
33	A simplified scoring system in de novo follicular lymphoma treated initially with immunochemotherapy. Blood, 2018, 132, 49-58.	0.6	130
34	Comparison of Microsatellites Versus Single-Nucleotide Polymorphisms in a Genome Linkage Screen for Prostate Cancerâ€“Susceptibility Loci. American Journal of Human Genetics, 2004, 75, 948-965.	2.6	129
35	Tumor Necrosis Factor (TNF) and Lymphotoxin-Â (LTA) Polymorphisms and Risk of Non-Hodgkin Lymphoma in the InterLymph Consortium. American Journal of Epidemiology, 2010, 171, 267-276.	1.6	128
36	Brief Report: Natural History of Individuals With Clinically Recognized Monoclonal B-Cell Lymphocytosis Compared With Patients With Rai 0 Chronic Lymphocytic Leukemia. Journal of Clinical Oncology, 2009, 27, 3959-3963.	0.8	123

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37	Familial predisposition and genetic risk factors for lymphoma. Blood, 2015, 126, 2265-2273.	0.6	122
38	Evidence for a susceptibility locus for panic disorder near the catechol-O-methyltransferase gene on chromosome 22. Biological Psychiatry, 2002, 51, 591-601.	0.7	118
39	Genome-wide association study identifies a novel susceptibility locus at 6p21.3 among familial CLL. Blood, 2011, 117, 1911-1916.	0.6	118
40	Evaluation of Candidate Genes in Case-Control Studies: A Statistical Method to Account for Related Subjects. American Journal of Human Genetics, 2001, 68, 1457-1462.	2.6	114
41	Quality of life in chronic lymphocytic leukemia: an international survey of 1482 patients. British Journal of Haematology, 2007, 139, 255-264.	1.2	112
42	Vitamin D insufficiency and prognosis in chronic lymphocytic leukemia. Blood, 2011, 117, 1492-1498.	0.6	110
43	Common Breast Cancer Susceptibility Loci Are Associated with Triple-Negative Breast Cancer. Cancer Research, 2011, 71, 6240-6249.	0.4	109
44	BCL2 mutations are associated with increased risk of transformation and shortened survival in follicular lymphoma. Blood, 2015, 125, 658-667.	0.6	108
45	Age at diagnosis and the utility of prognostic testing in patients with chronic lymphocytic leukemia. Cancer, 2010, 116, 4777-4787.	2.0	107
46	Analysis of the RNASEL Gene in Familial and Sporadic Prostate Cancer. American Journal of Human Genetics, 2002, 71, 116-123.	2.6	105
47	Evidence that breast cancer risk at the 2q35 locus is mediated through IGFBP5 regulation. Nature Communications, 2014, 5, 4999.	5.8	105
48	Nonâ€Hodgkin lymphoma and obesity: A pooled analysis from the InterLymph Consortium. International Journal of Cancer, 2008, 122, 2062-2070.	2.3	104
49	B-cell count and survival: differentiating chronic lymphocytic leukemia from monoclonal B-cell lymphocytosis based on clinical outcome. Blood, 2009, 113, 4188-4196.	0.6	104
50	Atrial fibrillation in patients with chronic lymphocytic leukemia (CLL). Leukemia and Lymphoma, 2017, 58, 1630-1639.	0.6	102
51	The prognostic significance of cytopenia in chronic lymphocytic leukaemia/small lymphocytic lymphoma. British Journal of Haematology, 2008, 141, 615-621.	1.2	101
52	19p13.1 Is a Triple-Negativeâ€“Specific Breast Cancer Susceptibility Locus. Cancer Research, 2012, 72, 1795-1803.	0.4	100
53	Fine-Scale Mapping of the FGFR2 Breast Cancer Risk Locus: Putative Functional Variants Differentially Bind FOXA1 and E2F1. American Journal of Human Genetics, 2013, 93, 1046-1060.	2.6	98
54	Medical History, Lifestyle, Family History, and Occupational Risk Factors for Diffuse Large B-Cell Lymphoma: The InterLymph Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 15-25.	0.9	98

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55	Base resolution methylome profiling: considerations in platform selection, data preprocessing and analysis. Epigenomics, 2015, 7, 813-828.	1.0	97
56	A BAFF-R mutation associated with non-Hodgkin lymphoma alters TRAF recruitment and reveals new insights into BAFF-R signaling. Journal of Experimental Medicine, 2010, 207, 2569-2579.	4.2	96
57	Genome-wide Association Study Identifies Five Susceptibility Loci for Follicular Lymphoma outside the HLA Region. American Journal of Human Genetics, 2014, 95, 462-471.	2.6	96
58	Tumor Budding in Colorectal Carcinoma. American Journal of Surgical Pathology, 2015, 39, 1340-1346.	2.1	95
59	Meta-analysis of genome-wide association studies discovers multiple loci for chronic lymphocytic leukemia. Nature Communications, 2016, 7, 10933.	5.8	94
60	GWAS of Follicular Lymphoma Reveals Allelic Heterogeneity at 6p21.32 and Suggests Shared Genetic Susceptibility with Diffuse Large B-cell Lymphoma. PLoS Genetics, 2011, 7, e1001378.	1.5	93
61	Identification of four novel susceptibility loci for oestrogen receptor negative breast cancer. Nature Communications, 2016, 7, 11375.	5.8	93
62	Diagnosis-to-Treatment Interval Is an Important Clinical Factor in Newly Diagnosed Diffuse Large B-Cell Lymphoma and Has Implication for Bias in Clinical Trials. Journal of Clinical Oncology, 2018, 36, 1603-1610.	0.8	93
63	Impact of Ibrutinib and Idelalisib on the Pharmaceutical Cost of Treating Chronic Lymphocytic Leukemia at the Individual and Societal Levels. Journal of Oncology Practice, 2015, 11, 252-258.	2.5	92
64	Elevated Serum B-Lymphocyte Stimulator Levels in Patients With Familial Lymphoproliferative Disorders. Journal of Clinical Oncology, 2006, 24, 983-987.	0.8	85
65	The efficacy of ibrutinib in the treatment of Richter syndrome. Blood, 2015, 125, 1676-1678.	0.6	83
66	Medical History, Lifestyle, Family History, and Occupational Risk Factors for Chronic Lymphocytic Leukemia/Small Lymphocytic Lymphoma: The InterLymph Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 41-51.	0.9	82
67	A high-density SNP genome-wide linkage search of 206 families identifies susceptibility loci for chronic lymphocytic leukemia. Blood, 2007, 110, 3326-3333.	0.6	79
68	A comprehensive evaluation of the prognostic significance of 13q deletions in patients with Bâ€chronic lymphocytic leukaemia. British Journal of Haematology, 2010, 148, 544-550.	1.2	79
69	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breastâ€“ovarian cancer susceptibility locus. Nature Communications, 2016, 7, 12675.	5.8	78
70	Hypogammaglobulinemia in newly diagnosed chronic lymphocytic leukemia: Natural history, clinical correlates, and outcomes. Cancer, 2015, 121, 2883-2891.	2.0	77
71	Risk factors for development of a second lymphoid malignancy in patients with chronic lymphocytic leukaemia. British Journal of Haematology, 2007, 139, 398-404.	1.2	76
72	Common variation at 6p21.31 (BAK1) influences the risk of chronic lymphocytic leukemia. Blood, 2012, 120, 843-846.	0.6	76

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73	Fine-Scale Mapping of the 5q11.2 Breast Cancer Locus Reveals at Least Three Independent Risk Variants Regulating MAP3K1. American Journal of Human Genetics, 2015, 96, 5-20.	2.6	76
74	Statin Use and Prognosis in Patients With Diffuse Large B-Cell Lymphoma and Follicular Lymphoma in the Rituximab Era. Journal of Clinical Oncology, 2010, 28, 412-417.	0.8	75
75	BRCA2 Hypomorphic Missense Variants Confer Moderate Risks of Breast Cancer. Cancer Research, 2017, 77, 2789-2799.	0.4	75
76	Genome-wide association analysis implicates dysregulation of immunity genes in chronic lymphocytic leukaemia. Nature Communications, 2017, 8, 14175.	5.8	75
77	Assessment of polygenic architecture and risk prediction based on common variants across fourteen cancers. Nature Communications, 2020, 11, 3353.	5.8	75
78	Validation of a new prognostic index for patients with chronic lymphocytic leukemia. Cancer, 2009, 115, 363-372.	2.0	72
79	Medical History, Lifestyle, Family History, and Occupational Risk Factors for Marginal Zone Lymphoma: The InterLymph Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 52-65.	0.9	70
80	Autoimmune cytopenia in chronic lymphocytic leukemia/small lymphocytic lymphoma: changes in clinical presentation and prognosis. Leukemia and Lymphoma, 2009, 50, 1261-1268.	0.6	69
81	<scp>H</scp>odgkin transformation of chronic lymphocytic leukemia: <scp>I</scp>ncidence, outcomes, and comparison to de novo <scp>H</scp>odgkin lymphoma. American Journal of Hematology, 2015, 90, 334-338.	2.0	69
82	The oncogenic transcription factor IRF4 is regulated by a novel CD30/NF-Î°B positive feedback loop in peripheral T-cell lymphoma. Blood, 2015, 125, 3118-3127.	0.6	68
83	A genome-wide meta-analysis of nodular sclerosing Hodgkin lymphoma identifies risk loci at 6p21.32. Blood, 2012, 119, 469-475.	0.6	66
84	Relationship between coâ€morbidities at diagnosis, survival and ultimate cause of death in patients with chronic lymphocytic leukaemia (<scp>CLL</scp>): a prospective cohort study. British Journal of Haematology, 2017, 178, 394-402.	1.2	66
85	Prognostic Significance of Pretreatment Serum Cytokines in Classical Hodgkin Lymphoma. Clinical Cancer Research, 2013, 19, 6812-6819.	3.2	64
86	Clinical characteristics and outcomes of Richter transformation: experience of 204 patients from a single center. Haematologica, 2020, 105, 765-773.	1.7	64
87	Patterns of Dietary Fluoride Supplement Use During Infancy. Journal of Public Health Dentistry, 1998, 58, 228-233.	0.5	63
88	Common occurrence of monoclonal Bâ€cell lymphocytosis among members of highâ€risk CLL families. British Journal of Haematology, 2010, 151, 152-158.	1.2	61
89	No association of germline alteration of MSR1 with prostate cancer risk. Nature Genetics, 2003, 35, 128-129.	9.4	60
90	Chronic lymphocytic leukemia in young (<= 55 years) patients: a comprehensive analysis of prognostic factors and outcomes. Haematologica, 2014, 99, 140-147.	1.7	60

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91	Genetic Variation in B-Cellâ€“Activating Factor Is Associated with an Increased Risk of Developing B-Cell Nonâ€“Hodgkin Lymphoma. Cancer Research, 2009, 69, 4217-4224.	0.4	59
92	Treatment of autoimmune cytopenia complicating progressive chronic lymphocytic leukemia/small lymphocytic lymphoma with rituximab, cyclophosphamide, vincristine, and prednisone. Leukemia and Lymphoma, 2010, 51, 620-627.	0.6	59
93	Evidence that the 5p12 Variant rs10941679 Confers Susceptibility to Estrogen-Receptor-Positive Breast Cancer through FGF10 and MRPS30 Regulation. American Journal of Human Genetics, 2016, 99, 903-911.	2.6	59
94	Association of Mu-Opioid Receptor Variants and Response to Citalopram Treatment in Major Depressive Disorder. American Journal of Psychiatry, 2010, 167, 565-573.	4.0	58
95	The Functional Assessment of Cancer Therapy - General (FACT-G) is valid for monitoring quality of life in patients with non-Hodgkin lymphoma. Leukemia and Lymphoma, 2013, 54, 290-297.	0.6	58
96	A genome-wide association study of marginal zone lymphoma shows association to the HLA region. Nature Communications, 2015, 6, 5751.	5.8	58
97	Cohort Profile: The Lymphoma Specialized Program of Research Excellence (SPORE) Molecular Epidemiology Resource (MER) Cohort Study. International Journal of Epidemiology, 2017, 46, 1753-1754i.	0.9	57
98	Overall and Cancer-Specific Survival of Patients With Breast, Colon, Kidney, and Lung Cancers With and Without Chronic Lymphocytic Leukemia: A SEER Population-Based Study. Journal of Clinical Oncology, 2013, 31, 930-937.	0.8	56
99	Germline Lysine-Specific Demethylase 1 (LSD1/KDM1A) Mutations Confer Susceptibility to Multiple Myeloma. Cancer Research, 2018, 78, 2747-2759.	0.4	56
100	Recurrent MSCE116K mutations in ALK-negative anaplastic large cell lymphoma. Blood, 2019, 133, 2776-2789.	0.6	55
101	Associations of Non-Hodgkin Lymphoma (NHL) Risk With Autoimmune Conditions According to Putative NHL Loci. American Journal of Epidemiology, 2015, 181, 406-421.	1.6	54
102	Identification of recurrent truncated <scp>DDX</scp>3X mutations in chronic lymphocytic leukaemia. British Journal of Haematology, 2015, 169, 445-448.	1.2	54
103	Rationale and Design of the International Lymphoma Epidemiology Consortium (InterLymph) Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 1-14.	0.9	52
104	Genetically predicted longer telomere length is associated with increased risk of B-cell lymphoma subtypes. Human Molecular Genetics, 2016, 25, 1663-1676.	1.4	52
105	Pretreatment circulating serum cytokines associated with follicular and diffuse large B-cell lymphoma: A clinic-based case-control study. Cytokine, 2012, 60, 882-889.	1.4	50
106	MicroRNA Related Polymorphisms and Breast Cancer Risk. PLoS ONE, 2014, 9, e109973.	1.1	49
107	Genome linkage screen for prostate cancer susceptibility loci: Results from the Mayo Clinic familial prostate cancer study. Prostate, 2003, 57, 335-346.	1.2	48
108	DNA Glycosylases Involved in Base Excision Repair May Be Associated with Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. PLoS Genetics, 2014, 10, e1004256.	1.5	47

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109	Deep sequencing identifies genetic heterogeneity and recurrent convergent evolution in chronic lymphocytic leukemia. Blood, 2015, 125, 492-498.	0.6	47
110	Elevated serum levels of IL-2R, IL-1RA, and CXCL9 are associated with a poor prognosis in follicular lymphoma. Blood, 2015, 125, 992-998.	0.6	47
111	PatternCNV: a versatile tool for detecting copy number changes from exome sequencing data. Bioinformatics, 2014, 30, 2678-2680.	1.8	43
112	Genome-wide linkage scan for prostate cancer aggressiveness loci using families from the University of Michigan Prostate Cancer Genetics Project. Prostate, 2006, 66, 173-179.	1.2	42
113	Use of positron emission tomography-computed tomography in the management of patients with chronic lymphocytic leukemia/small lymphocytic lymphoma. Leukemia and Lymphoma, 2014, 55, 2079-2084.	0.6	42
114	Rapid disease progression following discontinuation of ibrutinib in patients with chronic lymphocytic leukemia treated in routine clinical practice. Leukemia and Lymphoma, 2019, 60, 2712-2719.	0.6	42
115	Personalized risk prediction for eventâ€free survival at 24 months in patients with diffuse large Bâ€cell lymphoma. American Journal of Hematology, 2016, 91, 179-184.	2.0	41
116	Fine-mapping identifies two additional breast cancer susceptibility loci at 9q31.2. Human Molecular Genetics, 2015, 24, 2966-2984.	1.4	40
117	Germline Variation in Apoptosis Pathway Genes and Risk of Nonâ€“Hodgkin's Lymphoma. Cancer Epidemiology Biomarkers and Prevention, 2010, 19, 2847-2858.	1.1	39
118	Breast Cancer Polygenic Risk Score and Contralateral Breast Cancer Risk. American Journal of Human Genetics, 2020, 107, 837-848.	2.6	39
119	Hematologist/oncologist diseaseâ€specific expertise and survival: Lessons from chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL). Cancer, 2012, 118, 1827-1837.	2.0	38
120	Genome-wide association study identifies variants at 16p13 associated with survival in multiple myeloma patients. Nature Communications, 2015, 6, 7539.	5.8	38
121	Identification and characterization of novel associations in the CASP8/ALS2CR12 region on chromosome 2 with breast cancer risk. Human Molecular Genetics, 2015, 24, 285-298.	1.4	38
122	Functional and Clinical Significance of Variants Localized to 8q24 in Colon Cancer. Cancer Epidemiology Biomarkers and Prevention, 2009, 18, 2492-2500.	1.1	37
123	The physicianâ€“patient relationship and quality of life: Lessons from chronic lymphocytic leukemia. Leukemia Research, 2009, 33, 263-270.	0.4	37
124	Foodâ€frequency questionnaireâ€based estimates of total antioxidant capacity and risk of nonâ€Hodgkin lymphoma. International Journal of Cancer, 2012, 131, 1158-1168.	2.3	37
125	Elevated pretreatment serum levels of interferonâ€inducible proteinâ€10 (CXCL10) predict disease relapse and prognosis in diffuse large Bâ€cell lymphoma patients. American Journal of Hematology, 2012, 87, 865-869.	2.0	37
126	Autoimmune cytopenias in patients with chronic lymphocytic leukaemia treated with ibrutinib in routine clinical practice at an academic medical centre. British Journal of Haematology, 2018, 183, 421-427.	1.2	37

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127	Amplification of 9p24.1 in diffuse large B-cell lymphoma identifies a unique subset of cases that resemble primary mediastinal large B-cell lymphoma. Blood Cancer Journal, 2019, 9, 73.	2.8	37
128	Design and validity of a clinic-based case-control study on the molecular epidemiology of lymphoma. International Journal of Molecular Epidemiology and Genetics, 2011, 2, 95-113.	0.4	37
129	Chronic lymphocytic leukaemia genetics overview. British Journal of Haematology, 2007, 139, 630-634.	1.2	36
130	Postmenopausal hormone therapy and colorectal cancer risk by molecularly defined subtypes among older women. Gut, 2012, 61, 1299-1305.	6.1	36
131	Germline variation in complement genes and eventâ€free survival in follicular and diffuse large Bâ€cell lymphoma. American Journal of Hematology, 2012, 87, 880-885.	2.0	36
132	trans Fatty Acid Intake Is Associated with Increased Risk and n3 Fatty Acid Intake with Reduced Risk of Non-Hodgkin Lymphoma. Journal of Nutrition, 2013, 143, 672-681.	1.3	36
133	Loss of TNFAIP3 enhances MYD88L265P-driven signaling in non-Hodgkin lymphoma. Blood Cancer Journal, 2018, 8, 97.	2.8	36
134	The impact of dose modification and temporary interruption of ibrutinib on outcomes of chronic lymphocytic leukemia patients in routine clinical practice. Cancer Medicine, 2020, 9, 3390-3399.	1.3	36
135	A comprehensive study of polymorphisms in the ABCB1, ABCC2, ABCG2, NR1I2 genes and lymphoma risk. International Journal of Cancer, 2012, 131, 803-812.	2.3	35
136	Patients with chronic lymphocytic leukaemia and clonal deletion of both 17p13.1 and 11q22.3 have a very poor prognosis. British Journal of Haematology, 2013, 163, 326-333.	1.2	35
137	PRRC2A and BCL2L11 gene variants influence risk of non-Hodgkin lymphoma: results from the InterLymph consortium. Blood, 2012, 120, 4645-4648.	0.6	34
138	Early life sun exposure, vitamin D-related gene variants, and risk of non-Hodgkin lymphoma. Cancer Causes and Control, 2012, 23, 1017-1029.	0.8	34
139	RVboost: RNA-seq variants prioritization using a boosting method. Bioinformatics, 2014, 30, 3414-3416.	1.8	34
140	HLA Class I and II Diversity Contributes to the Etiologic Heterogeneity of Non-Hodgkin Lymphoma Subtypes. Cancer Research, 2018, 78, 4086-4096.	0.4	34
141	Familial chronic lymphocytic leukemia. Current Opinion in Hematology, 2010, 17, 350-355.	1.2	33
142	Associations Between Intake of Folate and Related Micronutrients with Molecularly Defined Colorectal Cancer Risks in the Iowa Women's Health Study. Nutrition and Cancer, 2012, 64, 899-910.	0.9	33
143	Incidence of chronic lymphocytic leukemia and highâ€count monoclonal Bâ€cell lymphocytosis using the 2008 guidelines. Cancer, 2014, 120, 2000-2005.	2.0	33
144	Pharmacovigilance during ibrutinib therapy for chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL) in routine clinical practice. Leukemia and Lymphoma, 2017, 58, 1376-1383.	0.6	33

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145	Cigarette Smoking and Colorectal Cancer Risk by KRAS Mutation Status Among Older Women. American Journal of Gastroenterology, 2012, 107, 782-789.	0.2	32
146	Medical History, Lifestyle, Family History, and Occupational Risk Factors for Lymphoplasmacytic Lymphoma/Waldenstrom's Macroglobulinemia: The InterLymph Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 87-97.	0.9	32
147	Medical History, Lifestyle, Family History, and Occupational Risk Factors for Sporadic Burkitt Lymphoma/Leukemia: The Interlymph Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 106-114.	0.9	32
148	Atrial fibrillation in patients with chronic lymphocytic leukemia (CLL) treated with ibrutinib: risk prediction, management, and clinical outcomes. Annals of Hematology, 2021, 100, 143-155.	0.8	32
149	Genetic Susceptibility Variants for Chronic Lymphocytic Leukemia. Cancer Epidemiology Biomarkers and Prevention, 2010, 19, 1098-1102.	1.1	31
150	Medical History, Lifestyle, and Occupational Risk Factors for Hairy Cell Leukemia: The InterLymph Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 115-124.	0.9	31
151	Medical History, Lifestyle, Family History, and Occupational Risk Factors for Mantle Cell Lymphoma: The InterLymph Non-Hodgkin Lymphoma Subtypes Project. Journal of the National Cancer Institute Monographs, 2014, 2014, 76-86.	0.9	31
152	Identification of independent association signals and putative functional variants for breast cancer risk through fine-scale mapping of the 12p11 locus. Breast Cancer Research, 2016, 18, 64.	2.2	31
153	Novel pedigree analysis implicates DNA repair and chromatin remodeling in multiple myeloma risk. PLoS Genetics, 2018, 14, e1007111.	1.5	30
154	CXCR5 polymorphisms in non-Hodgkin lymphoma risk and prognosis. Cancer Immunology, Immunotherapy, 2013, 62, 1475-1484.	2.0	28
155	Impact of post-alignment processing in variant discovery from whole exome data. BMC Bioinformatics, 2016, 17, 403.	1.2	28
156	Genetic overlap between autoimmune diseases and nonâ€Hodgkin lymphoma subtypes. Genetic Epidemiology, 2019, 43, 844-863.	0.6	28
157	Targeting of inflammatory pathways with R2CHOP in high-risk DLBCL. Leukemia, 2021, 35, 522-533.	3.3	28
158	Lack of intrafollicular memory CD4â€‰+â€‰T cells is predictive of early clinical failure in newly diagnosed follicular lymphoma. Blood Cancer Journal, 2021, 11, 130.	2.8	27
159	Susceptibility genes and Bâ€chronic lymphocytic leukaemia. British Journal of Haematology, 2007, 139, 762-771.	1.2	26
160	Genetic Susceptibility to Chronic Lymphocytic Leukemia. Seminars in Hematology, 2013, 50, 296-302.	1.8	26
161	Genetic polymorphisms in oxidative stressâ€related genes are associated with outcomes following treatment for aggressive Bâ€cell nonâ€Hodgkin lymphoma. American Journal of Hematology, 2014, 89, 639-645.	2.0	26
162	History of autoimmune conditions and lymphoma prognosis. Blood Cancer Journal, 2018, 8, 73.	2.8	26

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164	Genome-wide Association Study Identifies HLA-DPB1 as a Significant Risk Factor for Severe Aplastic Anemia. American Journal of Human Genetics, 2020, 106, 264-271.	2.6	25
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