

Lucy A Godley

List of Publications by Year in descending order

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Version: 2024-02-01

161
papers

14,597
citations

53794

45
h-index

19749

117
g-index

166
all docs

166
docs citations

166
times ranked

17475
citing authors

#	ARTICLE	IF	CITATIONS
1	Leukemic IDH1 and IDH2 Mutations Result in a Hypermethylation Phenotype, Disrupt TET2 Function, and Impair Hematopoietic Differentiation. <i>Cancer Cell</i> , 2010, 18, 553-567.	16.8	2,328
2	Tet2 Loss Leads to Increased Hematopoietic Stem Cell Self-Renewal and Myeloid Transformation. <i>Cancer Cell</i> , 2011, 20, 11-24.	16.8	1,105
3	Selective chemical labeling reveals the genome-wide distribution of 5-hydroxymethylcytosine. <i>Nature Biotechnology</i> , 2011, 29, 68-72.	17.5	955
4	Dnmt3a is essential for hematopoietic stem cell differentiation. <i>Nature Genetics</i> , 2012, 44, 23-31.	21.4	916
5	TET2 Inactivation Results in Pleiotropic Hematopoietic Abnormalities in Mouse and Is a Recurrent Event during Human Lymphomagenesis. <i>Cancer Cell</i> , 2011, 20, 25-38.	16.8	792
6	5-hmC-mediated epigenetic dynamics during postnatal neurodevelopment and aging. <i>Nature Neuroscience</i> , 2011, 14, 1607-1616.	14.8	746
7	Recurrent somatic TET2 mutations in normal elderly individuals with clonal hematopoiesis. <i>Nature Genetics</i> , 2012, 44, 1179-1181.	21.4	692
8	Inherited and Somatic Defects in DDX41 in Myeloid Neoplasms. <i>Cancer Cell</i> , 2015, 27, 658-670.	16.8	341
9	Germline ETV6 mutations in familial thrombocytopenia and hematologic malignancy. <i>Nature Genetics</i> , 2015, 47, 180-185.	21.4	299
10	Microbial signals drive pre-leukaemic myeloproliferation in a Tet2-deficient host. <i>Nature</i> , 2018, 557, 580-584.	27.8	296
11	Large conserved domains of low DNA methylation maintained by Dnmt3a. <i>Nature Genetics</i> , 2014, 46, 17-23.	21.4	276
12	Therapy-Related Myeloid Leukemia. <i>Seminars in Oncology</i> , 2008, 35, 418-429.	2.2	272
13	Therapy-related myeloid neoplasms: when genetics and environment collide. <i>Nature Reviews Cancer</i> , 2017, 17, 513-527.	28.4	270
14	DNA Hydroxymethylation Profiling Reveals that WT1 Mutations Result in Loss of TET2 Function in Acute Myeloid Leukemia. <i>Cell Reports</i> , 2014, 9, 1841-1855.	6.4	237
15	Fumarate and Succinate Regulate Expression of Hypoxia-inducible Genes via TET Enzymes. <i>Journal of Biological Chemistry</i> , 2016, 291, 4256-4265.	3.4	234
16	Geriatric assessment to predict survival in older allogeneic hematopoietic cell transplantation recipients. <i>Haematologica</i> , 2014, 99, 1373-1379.	3.5	213
17	Genomic analysis of germ line and somatic variants in familial myelodysplasia/acute myeloid leukemia. <i>Blood</i> , 2015, 126, 2484-2490.	1.4	207
18	Mechanism-Based Epigenetic Chemosensitization Therapy of Diffuse Large B-Cell Lymphoma. <i>Cancer Discovery</i> , 2013, 3, 1002-1019.	9.4	180

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19	Novel germ line DDX41 mutations define families with a lower age of MDS/AML onset and lymphoid malignancies. <i>Blood</i> , 2016, 127, 1017-1023.	1.4	179
20	Inhibition of TET2-mediated conversion of 5-methylcytosine to 5-hydroxymethylcytosine disturbs erythroid and granulomonocytic differentiation of human hematopoietic progenitors. <i>Blood</i> , 2011, 118, 2551-2555.	1.4	163
21	Reduced-intensity conditioning with combined haploidentical and cord blood transplantation results in rapid engraftment, low GVHD, and durable remissions. <i>Blood</i> , 2011, 118, 6438-6445.	1.4	158
22	TET1-Mediated Hydroxymethylation Facilitates Hypoxic Gene Induction in Neuroblastoma. <i>Cell Reports</i> , 2014, 7, 1343-1352.	6.4	146
23	Genetic predisposition to hematologic malignancies: management and surveillance. <i>Blood</i> , 2017, 130, 424-432.	1.4	145
24	Inherited mutations in cancer susceptibility genes are common among survivors of breast cancer who develop therapy-related leukemia. <i>Cancer</i> , 2016, 122, 304-311.	4.1	129
25	Genome-Wide Variation of Cytosine Modifications Between European and African Populations and the Implications for Complex Traits. <i>Genetics</i> , 2013, 194, 987-996.	2.9	117
26	Effects of TET2 mutations on DNA methylation in chronic myelomonocytic leukemia. <i>Epigenetics</i> , 2012, 7, 201-207.	2.7	110
27	ClinGen Myeloid Malignancy Variant Curation Expert Panel recommendations for germline RUNX1 variants. <i>Blood Advances</i> , 2019, 3, 2962-2979.	5.2	110
28	RUNX1-mutated families show phenotype heterogeneity and a somatic mutation profile unique to germline predisposed AML. <i>Blood Advances</i> , 2020, 4, 1131-1144.	5.2	102
29	Familial myelodysplastic syndrome/acute leukemia syndromes: a review and utility for translational investigations. <i>Annals of the New York Academy of Sciences</i> , 2014, 1310, 111-118.	3.8	95
30	Hydroxymethylation at Gene Regulatory Regions Directs Stem/Early Progenitor Cell Commitment during Erythropoiesis. <i>Cell Reports</i> , 2014, 6, 231-244.	6.4	93
31	Proposal for the clinical detection and management of patients and their family members with familial myelodysplastic syndrome/acute leukemia predisposition syndromes. <i>Leukemia and Lymphoma</i> , 2013, 54, 28-35.	1.3	88
32	DNA Methylation Dynamics of Germinal Center B Cells Are Mediated by AID. <i>Cell Reports</i> , 2015, 12, 2086-2098.	6.4	87
33	Inherited Predisposition to Acute Myeloid Leukemia. <i>Seminars in Hematology</i> , 2014, 51, 306-321.	3.4	85
34	Prognostic tumor sequencing panels frequently identify germ line variants associated with hereditary hematopoietic malignancies. <i>Blood Advances</i> , 2018, 2, 146-150.	5.2	83
35	TET2 Mutations Affect Non-CpG Island DNA Methylation at Enhancers and Transcription Factor Binding Sites in Chronic Myelomonocytic Leukemia. <i>Cancer Research</i> , 2015, 75, 2833-2843.	0.9	80
36	5-hydroxymethylcytosine in cancer: significance in diagnosis and therapy. <i>Cancer Genetics</i> , 2015, 208, 167-177.	0.4	77

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37	Evaluation of Patients and Families With Concern for Predispositions to Hematologic Malignancies Within the Hereditary Hematologic Malignancy Clinic (HHMC). <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2016, 16, 417-428.e2.	0.4	74
38	Genome-wide hydroxymethylation tested using the HELP-GT assay shows redistribution in cancer. <i>Nucleic Acids Research</i> , 2013, 41, e157-e157.	14.5	69
39	Germline ETV6 mutations and predisposition to hematological malignancies. <i>International Journal of Hematology</i> , 2017, 106, 189-195.	1.6	64
40	Genetic predisposition to leukemia and other hematologic malignancies. <i>Seminars in Oncology</i> , 2016, 43, 598-608.	2.2	58
41	DNMT3B7, a Truncated DNMT3B Isoform Expressed in Human Tumors, Disrupts Embryonic Development and Accelerates Lymphomagenesis. <i>Cancer Research</i> , 2010, 70, 5840-5850.	0.9	56
42	Germline Lysine-Specific Demethylase 1 (<i>LSD1/KDM1A</i>) Mutations Confer Susceptibility to Multiple Myeloma. <i>Cancer Research</i> , 2018, 78, 2747-2759.	0.9	56
43	Recognizing familial myeloid leukemia in adults. <i>Therapeutic Advances in Hematology</i> , 2013, 4, 254-269.	2.5	55
44	A phase I study of selinexor in combination with high-dose cytarabine and mitoxantrone for remission induction in patients with acute myeloid leukemia. <i>Journal of Hematology and Oncology</i> , 2018, 11, 4.	17.0	52
45	Altered hydroxymethylation is seen at regulatory regions in pancreatic cancer and regulates oncogenic pathways. <i>Genome Research</i> , 2017, 27, 1830-1842.	5.5	51
46	TET-catalyzed 5-hydroxymethylcytosine regulates gene expression in differentiating colonocytes and colon cancer. <i>Scientific Reports</i> , 2015, 5, 17568.	3.3	50
47	New themes in the biological functions of 5-â€methylcytosine and 5-â€hydroxymethylcytosine. <i>Immunological Reviews</i> , 2015, 263, 36-49.	6.0	48
48	Alterations of 5-Hydroxymethylcytosine in Human Cancers. <i>Cancers</i> , 2013, 5, 786-814.	3.7	46
49	Dnmt3b is a haploinsufficient tumor suppressor gene in Myc-induced lymphomagenesis. <i>Blood</i> , 2013, 121, 2059-2063.	1.4	44
50	Germline variants drive myelodysplastic syndrome in young adults. <i>Leukemia</i> , 2021, 35, 2439-2444.	7.2	43
51	Identifying Inherited and Acquired Genetic Factors Involved in Poor Stem Cell Mobilization and Donor-Derived Malignancy. <i>Biology of Blood and Marrow Transplantation</i> , 2016, 22, 2100-2103.	2.0	42
52	Cytokine-Regulated Phosphorylation and Activation of TET2 by JAK2 in Hematopoiesis. <i>Cancer Discovery</i> , 2019, 9, 778-795.	9.4	41
53	The Role of Gene Body Cytosine Modifications in <i>MGMT</i> Expression and Sensitivity to Temozolomide. <i>Molecular Cancer Therapeutics</i> , 2014, 13, 1334-1344.	4.1	40
54	Brca1 deficiency causes bone marrow failure and spontaneous hematologic malignancies in mice. <i>Blood</i> , 2016, 127, 310-313.	1.4	39

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55	Pharmacogenomics of chemotherapeutic susceptibility and toxicity. <i>Genome Medicine</i> , 2012, 4, 90.	8.2	38
56	Increased DNA methylation of Dnmt3b targets impairs leukemogenesis. <i>Blood</i> , 2016, 127, 1575-1586.	1.4	38
57	The use of hypomethylating agents in the treatment of hematologic malignancies. <i>Leukemia and Lymphoma</i> , 2007, 48, 1676-1695.	1.3	37
58	Recognition of familial myeloid neoplasia in adults. <i>Seminars in Hematology</i> , 2017, 54, 60-68.	3.4	37
59	Linking the genetic architecture of cytosine modifications with human complex traits. <i>Human Molecular Genetics</i> , 2014, 23, 5893-5905.	2.9	36
60	Truncated DNMT3B Isoform DNMT3B7 Suppresses Growth, Induces Differentiation, and Alters DNA Methylation in Human Neuroblastoma. <i>Cancer Research</i> , 2012, 72, 4714-4723.	0.9	35
61	The chemotherapeutic CX-5461 primarily targets TOP2B and exhibits selective activity in high-risk neuroblastoma. <i>Nature Communications</i> , 2021, 12, 6468.	12.8	35
62	Integrative genomics reveals hypoxia inducible genes that are associated with a poor prognosis in neuroblastoma patients. <i>Oncotarget</i> , 2016, 7, 76816-76826.	1.8	33
63	Targeted gene panels identify a high frequency of pathogenic germline variants in patients diagnosed with a hematological malignancy and at least one other independent cancer. <i>Leukemia</i> , 2021, 35, 3245-3256.	7.2	32
64	A new family with a germline <i>ANKRD26</i> mutation and predisposition to myeloid malignancies. <i>Leukemia and Lymphoma</i> , 2014, 55, 2945-2946.	1.3	30
65	Reduced intensity haplo plus single cord transplant compared to double cord transplant: improved engraftment and graft-versus-host disease-free, relapse-free survival. <i>Haematologica</i> , 2016, 101, 634-643.	3.5	30
66	Identification and molecular characterization of a novel β mutation in <i>RUNX1</i> in a family with familial platelet disorder. <i>Leukemia and Lymphoma</i> , 2010, 51, 1931-1935.	1.3	29
67	The RUNX1 database (RUNX1db): establishment of an expert curated RUNX1 registry and genomics database as a public resource for familial platelet disorder with myeloid malignancy. <i>Haematologica</i> , 2021, 106, 3004-3007.	3.5	29
68	Identifying potential germline variants from sequencing hematopoietic malignancies. <i>Blood</i> , 2020, 136, 2498-2506.	1.4	27
69	An update on the safety and efficacy of decitabine in the treatment of myelodysplastic syndromes. <i>OncoTargets and Therapy</i> , 2010, 3, 1.	2.0	25
70	Correspondence Regarding the Consensus Statement from the Worldwide Network for Blood and Marrow Transplantation Standing Committee on Donor Issues. <i>Biology of Blood and Marrow Transplantation</i> , 2016, 22, 183-184.	2.0	24
71	Inherited Susceptibility to Hematopoietic Malignancies in the Era of Precision Oncology. <i>JCO Precision Oncology</i> , 2021, 5, 107-122.	3.0	24
72	Telomere biology disorder prevalence and phenotypes in adults with familial hematologic and/or pulmonary presentations. <i>Blood Advances</i> , 2020, 4, 4873-4886.	5.2	23

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73	Heterozygous germ line CSF3R variants as risk alleles for development of hematologic malignancies. <i>Blood Advances</i> , 2020, 4, 5269-5284.	5.2	23
74	How I curate: applying American Society of Hematology-Clinical Genome Resource Myeloid Malignancy Variant Curation Expert Panel rules for RUNX1 variant curation for germline predisposition to myeloid malignancies. <i>Haematologica</i> , 2020, 105, 870-887.	3.5	23
75	5-Hydroxymethylcytosine Profiles in Circulating Cell-Free DNA Associate with Disease Burden in Children with Neuroblastoma. <i>Clinical Cancer Research</i> , 2020, 26, 1309-1317.	7.0	22
76	An Integrated Genomic Approach to the Assessment and Treatment of Acute Myeloid Leukemia. <i>Seminars in Oncology</i> , 2011, 38, 215-224.	2.2	21
77	Gene Mutations, Epigenetic Dysregulation, and Personalized Therapy in Myeloid Neoplasia: Are We There Yet?. <i>Seminars in Oncology</i> , 2011, 38, 196-214.	2.2	21
78	Profiles in Leukemia. <i>New England Journal of Medicine</i> , 2012, 366, 1152-1153.	27.0	21
79	2-Hydroxyglutarate in IDH mutant acute myeloid leukemia: predicting patient responses, minimal residual disease and correlations with methylcytosine and hydroxymethylcytosine levels. <i>Leukemia and Lymphoma</i> , 2013, 54, 408-410.	1.3	21
80	High dose cytarabine and mitoxantrone: an effective induction regimen for high-risk Acute Myeloid Leukemia (AML). <i>Leukemia and Lymphoma</i> , 2012, 53, 445-450.	1.3	20
81	Inherited predisposition to haematopoietic malignancies: overcoming barriers and exploring opportunities. <i>British Journal of Haematology</i> , 2021, 194, 663-676.	2.5	20
82	Identifying familial myelodysplastic/acute leukemia predisposition syndromes through hematopoietic stem cell transplantation donors with thrombocytopenia. <i>Blood</i> , 2012, 120, 5247-5249.	1.4	19
83	Germline predisposition to hematopoietic malignancies. <i>Human Molecular Genetics</i> , 2021, 30, R225-R235.	2.9	19
84	Reduced-Intensity Allogeneic Transplant for Acute Myeloid Leukemia and Myelodysplastic Syndrome Using Combined CD34-Selected Haploidentical Graft and a Single Umbilical Cord Unit Compared with Matched Unrelated Donor Stem Cells in Older Adults. <i>Biology of Blood and Marrow Transplantation</i> , 2018, 24, 997-1004.	2.0	18
85	Identifying patients with genetic predisposition to acute myeloid leukemia. <i>Best Practice and Research in Clinical Haematology</i> , 2018, 31, 373-378.	1.7	18
86	Hereditary Myelodysplastic Syndrome and Acute Myeloid Leukemia: Diagnosis, Questions, and Controversies. <i>Current Hematologic Malignancy Reports</i> , 2018, 13, 426-434.	2.3	17
87	Assessment of technical heterogeneity among diagnostic tests to detect germline risk variants for hematopoietic malignancies. <i>Genetics in Medicine</i> , 2021, 23, 211-214.	2.4	17
88	Treatment of therapy-related myeloid neoplasms with high-dose cytarabine/mitoxantrone followed by hematopoietic stem cell transplant. <i>Leukemia and Lymphoma</i> , 2010, 51, 995-1006.	1.3	16
89	Clinical Assessment and Diagnosis of Germline Predisposition to Hematopoietic Malignancies: The University of Chicago Experience. <i>Frontiers in Pediatrics</i> , 2017, 5, 252.	1.9	16
90	Somatic mutation panels: Time to clear their names. <i>Cancer Genetics</i> , 2019, 235-236, 84-92.	0.4	16

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91	Identifying potential germline variants from sequencing hematopoietic malignancies. Hematology American Society of Hematology Education Program, 2020, 2020, 219-227.	2.5	16
92	DNMT3B7 Expression Promotes Tumor Progression to a More Aggressive Phenotype in Breast Cancer Cells. PLoS ONE, 2015, 10, e0117310.	2.5	15
93	HIF-1 directly induces TET3 expression to enhance 5-hmC density and induce erythroid gene expression in hypoxia. Blood Advances, 2020, 4, 3053-3062.	5.2	15
94	Breaking the spatial constraint between neighboring zinc fingers: a new germline mutation in GATA2 deficiency syndrome. Leukemia, 2021, 35, 264-268.	7.2	15
95	Germline mutations in MDS/AML predisposition disorders. Current Opinion in Hematology, 2021, 28, 86-93.	2.5	15
96	Feasibility and limitations of cultured skin fibroblasts for germline genetic testing in hematologic disorders. Human Mutation, 2022, 43, 950-962.	2.5	15
97	The Next Frontier for Stem Cell Transplantation. JAMA - Journal of the American Medical Association, 2010, 303, 1421.	7.4	14
98	Perturbations of 5-Hydroxymethylcytosine Patterning in Hematologic Malignancies. Seminars in Hematology, 2013, 50, 61-69.	3.4	14
99	5-Hydroxymethylcytosine Profiles Are Prognostic of Outcome in Neuroblastoma and Reveal Transcriptional Networks That Correlate With Tumor Phenotype. JCO Precision Oncology, 2019, 3, 1-12.	3.0	14
100	MYC Regulation of D2HGDH and L2HGDH Influences the Epigenome and Epitranscriptome. Cell Chemical Biology, 2020, 27, 538-550.e7.	5.2	14
101	Study of inherited thrombocytopenia resulting from mutations in ETV6 or RUNX1 using a human pluripotent stem cell model. Stem Cell Reports, 2021, 16, 1458-1467.	4.8	14
102	Germline CHEK2 and ATM Variants in Myeloid and Other Hematopoietic Malignancies. Current Hematologic Malignancy Reports, 2022, 17, 94-104.	2.3	14
103	Clinical Predictors of Transplant Related Mortality after Reduced Intensity Allogeneic Stem Cell Transplantation (RIST).. Blood, 2004, 104, 1145-1145.	1.4	11
104	The identification and characterisation of novel <i>KIT</i> transcripts in aggressive mast cell malignancies and normal CD34+ cells. Leukemia and Lymphoma, 2008, 49, 1567-1577.	1.3	10
105	Characterization of CpG sites that escape methylation on the inactive human X-chromosome. Epigenetics, 2015, 10, 810-818.	2.7	9
106	Incidence and predictors of respiratory viral infections by multiplex PCR in allogeneic hematopoietic cell transplant recipients 50 years and older including geriatric assessment. Leukemia and Lymphoma, 2016, 57, 1807-1813.	1.3	9
107	A phase 1 study of azacitidine with high-dose cytarabine and mitoxantrone in high-risk acute myeloid leukemia. Blood Advances, 2020, 4, 599-606.	5.2	9
108	Genetics of Myelodysplastic Syndromes. Cancers, 2021, 13, 3380.	3.7	9

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109	A practical guide to interpreting germline variants that drive hematopoietic malignancies, bone marrow failure, and chronic cytopenias. <i>Genetics in Medicine</i> , 2022, 24, 931-954.	2.4	9
110	The Histone Code and Treatments for Acute Myeloid Leukemia. <i>New England Journal of Medicine</i> , 2012, 366, 960-961.	27.0	8
111	Dnmt3a Is Essential for Hematopoietic Stem Cell Differentiation. <i>Blood</i> , 2011, 118, 386-386.	1.4	7
112	Epigenetic Control of <i>Apolipoprotein E</i> Expression Mediates Gender-Specific Hematopoietic Regulation. <i>Stem Cells</i> , 2015, 33, 3643-3654.	3.2	6
113	Inherited Thrombocytopenia Caused by Germline <i>ANKRD26</i> Mutation Should Be Considered in Young Patients With Suspected Myelodysplastic Syndrome. <i>Journal of Investigative Medicine High Impact Case Reports</i> , 2020, 8, 232470962093894.	0.6	6
114	Clinical features and survival outcomes in patients with chronic myelomonocytic leukemia arising in the context of germline predisposition syndromes. <i>American Journal of Hematology</i> , 2021, 96, E327-E330.	4.1	6
115	Therapy-Related Myeloid Neoplasms in 109 Patients Following Radiation Monotherapy. <i>Blood Advances</i> , 2021, 5, 4140-4148.	5.2	6
116	Modulators of DNA methylation and histone acetylation. <i>Update on Cancer Therapeutics</i> , 2007, 2, 157-169.	0.4	5
117	<i>HMGA2</i> levels in CML: Reflective of miRNA gene regulation in a hematopoietic tumor?. <i>Leukemia and Lymphoma</i> , 2007, 48, 1898-1899.	1.3	5
118	On the Origin of Leukemic Species. <i>Cell Stem Cell</i> , 2014, 14, 421-422.	11.1	4
119	Characterization of cancer comorbidity prior to allogeneic hematopoietic cell transplantation. <i>Leukemia and Lymphoma</i> , 2019, 60, 629-638.	1.3	4
120	When should transplant physicians think about familial blood cancers?. <i>Advances in Cell and Gene Therapy</i> , 2019, 2, e68.	0.9	4
121	Regulation of telomeric function by DNA methylation differs between humans and mice. <i>Human Molecular Genetics</i> , 2020, 29, 3197-3210.	2.9	4
122	Efficacy and tolerability of a modified pediatric-inspired intensive regimen for acute lymphoblastic leukemia in older adults. <i>EJHaem</i> , 2021, 2, 413-420.	1.0	4
123	Deletion of the der(9q) in chronic myeloid leukemia: the controversy continues. <i>Leukemia and Lymphoma</i> , 2009, 50, 871-872.	1.3	3
124	Preference by Exclusion. <i>Science</i> , 2011, 331, 1017-1018.	12.6	3
125	Regulation of 5-Hydroxymethylcytosine Distribution by the TET Enzymes. <i>RNA Technologies</i> , 2019, , 229-263.	0.3	3
126	Inherited predisposition to myeloid malignancies. <i>Blood Advances</i> , 2019, 3, 2688-2688.	5.2	2

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127	Correct application of variant classification guidelines in germline RUNX1 mutated disorders to assist clinical diagnosis. <i>Leukemia and Lymphoma</i> , 2020, 61, 246-247.	1.3	2
128	RBL2 bi-allelic truncating variants cause severe motor and cognitive impairment without evidence for abnormalities in DNA methylation or telomeric function. <i>Journal of Human Genetics</i> , 2021, 66, 1101-1112.	2.3	2
129	Expanding Use of a Modified Pediatric Intensive Regimen for Acute Lymphoblastic Leukemia (ALL) into an Older Adult Population: Feasibility and Efficacy Results. <i>Blood</i> , 2020, 136, 41-42.	1.4	2
130	Identical Novel C-Kit Transcripts in Two Patients with Mast Cell Leukemia.. <i>Blood</i> , 2004, 104, 2001-2001.	1.4	2
131	Expanded Phenotypic and Genetic Heterogeneity in the Clinical Spectrum of FPD-AML: Lymphoid Malignancies and Skin Disorders Are Common Features in Carriers of Germline RUNX1 Mutations. <i>Blood</i> , 2016, 128, 1212-1212.	1.4	2
132	Assessing the Feasibility and Limitations of Cultured Skin Fibroblasts for Germline Genetic Testing in Hematologic Disorders. <i>Blood</i> , 2020, 136, 35-36.	1.4	2
133	Getting to the root of the stem cell in mutated chronic myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2010, 51, 2147-2148.	1.3	1
134	MBD4: guardian of the epigenetic galaxy. <i>Blood</i> , 2018, 132, 1468-1469.	1.4	1
135	Fludarabine Melphalan and Alemtuzumab (Campath) Conditioning for Pts with High Risk Myeloid Malignancies. High Cure Rate for Pts with Low Leukemia Burden.. <i>Blood</i> , 2004, 104, 2321-2321.	1.4	1
136	Leukemic Relapse after Allogeneic Stem Cell Transplantation with a T-Cell Depleted Reduced Intensity Conditioning (RIST) Regimen.. <i>Blood</i> , 2005, 106, 2022-2022.	1.4	1
137	Preliminary Results of Combined Haploidentical-Cord Blood Transplantation for Patients Lacking HLA Identical Donors. <i>Blood</i> , 2008, 112, 3015-3015.	1.4	1
138	Using sequential next-generation sequencing assays to identify germline cancer predisposition variants.. <i>Journal of Clinical Oncology</i> , 2020, 38, 1581-1581.	1.6	1
139	Limited Effect of TET2 Mutations on Promoter DNA Methylation in Chronic Myelomonocytic Leukemia. <i>Blood</i> , 2011, 118, 1365-1365.	1.4	1
140	Reduced Intensity Conditioning with Combined Haploidentical and Cord Blood Transplantation Results in Rapid Engraftment and Durable Remissions in Hematological Malignancies. <i>Blood</i> , 2011, 118, 830-830.	1.4	1
141	Cytokine-Regulated Phosphorylation and Activation of TET2 by JAK2 in Hematopoiesis. <i>Cancer Discovery</i> , 2019, , .	9.4	0
142	BET inhibitors enhance embryonic and fetal globin expression in erythroleukemia cell lines. <i>Haematologica</i> , 2021, 106, 3223-3227.	3.5	0
143	Anticipation in hematopoietic malignancies: biology, bias, or both?. <i>Leukemia and Lymphoma</i> , 2021, 62, 3070-3072.	1.3	0
144	Phase I Study of XK469R (NSC 698215), a Quinoxaline Phenoxypropionic Acid Derivative, in Patients with Refractory Hematological Malignancies.. <i>Blood</i> , 2006, 108, 1952-1952.	1.4	0

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145	New Cytogenetic Abnormalities Are Frequent in AML and MDS Relapsing after Allogeneic Hematopoietic Cell Transplantation (HCT).. Blood, 2006, 108, 3675-3675.	1.4	0
146	Novel C-KIT Transcripts Identified in Mast Cell Leukemia: An Update of the Full Transcript and Its Distribution.. Blood, 2007, 110, 2396-2396.	1.4	0
147	Clofarabine-Melphalan-Alemtuzumab Conditioning for Allogeneic Hematopoietic Cell Transplantation: Final Report of a Phase I-II Study. Blood, 2011, 118, 1948-1948.	1.4	0
148	Myc-Mediated Lymphomagenesis Is Driven by DNA Methylation Changes Induced by DNMT3B7 Expression and Dnmt3b Heterozygosity. Blood, 2011, 118, 225-225.	1.4	0
149	A Phase II Prospective Feasibility Study of Clofarabine Cyto-reduction Prior to Allogeneic Hematopoietic Cell Transplantation (HCT) for Patients with Relapsed or Refractory Acute Leukemias and Advanced Myelodysplastic Syndromes. Blood, 2011, 118, 496-496.	1.4	0
150	Dynamic Regulation of 5-Hydroxymethylcytosine At the β -Globin Promoter During Erythroid Differentiation. Blood, 2012, 120, 824-824.	1.4	0
151	Allogeneic Hematopoietic Cell Transplantation Is Effective In Patients With Advanced Systemic Mastocytosis: A Multicenter Retrospective Analysis. Blood, 2013, 122, 2145-2145.	1.4	0
152	Large Conserved Domains Of Low DNA Methylation Maintained By 5-Hydroxymethylcytosine and Dnmt3a. Blood, 2013, 122, 2406-2406.	1.4	0
153	Hematopoietic Stem Cell Function Is Regulated By Hormonal and Epigenetic Factors. Blood, 2013, 122, 1194-1194.	1.4	0
154	Identifying Inherited and Acquired Genetic Factors Involved in Poor Stem Cell Mobilization and Donor-Derived Malignancy. Blood, 2015, 126, 3163-3163.	1.4	0
155	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
156	Development of a Data Portal for Aggregation and Analysis of Genomics Data in Familial Platelet Disorder with Predisposition to Myeloid Malignancy - the RUNX1.DB. Blood, 2018, 132, 5241-5241.	1.4	0
157	Final Results from a Phase I Trial Combining Selinexor with High-Dose Cytarabine (HiDAC) and Mitoxantrone (Mito) for Remission Induction in Acute Myeloid Leukemia (AML). Blood, 2018, 132, 4073-4073.	1.4	0
158	Feasibility and Outcomes of T-Cell Depleted Hematopoietic Stem Cell Transplantation in Patients with Relapsed or Refractory AML and High Risk MDS. Blood, 2019, 134, 3324-3324.	1.4	0
159	Deleterious Germline Variants Are Present in Patients with Myelodysplastic Syndrome of All Ages Treated with Related Allogeneic Stem Cell Transplantation. Blood, 2021, 138, 320-320.	1.4	0
160	Spacing Constraints of Neighboring Zinc Finger Modules within GATA2. Blood, 2021, 138, 3306-3306.	1.4	0
161	Therapy-Related Myeloid Neoplasms in 108 Patients Following Radiation Therapy Only. Blood, 2020, 136, 25-26.	1.4	0