

Timothy A Graubert

List of Publications by Year in descending order

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Version: 2024-02-01

123
papers

19,311
citations

34105

52
h-index

25787

108
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126
all docs

126
docs citations

126
times ranked

24738
citing authors

#	ARTICLE	IF	CITATIONS
1	A synthetic small molecule stalls pre-mRNA splicing by promoting an early-stage U2AF2-RNA complex. <i>Cell Chemical Biology</i> , 2021, 28, 1145-1157.e6.	5.2	24
2	U2af1 is a haplo-essential gene required for hematopoietic cancer cell survival in mice. <i>Journal of Clinical Investigation</i> , 2021, 131, .	8.2	9
3	Discovery and Pharmacological Characterization of JNJ-64619178, a Novel Small-Molecule Inhibitor of PRMT5 with Potent Antitumor Activity. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 2317-2328.	4.1	48
4	Spliceosome Mutant Myeloid Malignancies Are Preferentially Sensitive to PARP Inhibition. <i>Blood</i> , 2021, 138, 322-322.	1.4	4
5	Inhibition of ATR with AZD6738 (Ceralasertib) for the Treatment of Progressive or Relapsed Myelodysplastic Syndromes and Chronic Myelomonocytic Leukemia: Safety and Preliminary Activity from a Phase Ib/II Study. <i>Blood</i> , 2021, 138, 1521-1521.	1.4	4
6	ATR/CHK1/WEE1 Dependency in SRSF2-Mutated MDS/AML. <i>Blood</i> , 2021, 138, 3661-3661.	1.4	1
7	A phase 1 study of the antibody-drug conjugate brentuximab vedotin with re-induction chemotherapy in patients with CD30-expressing relapsed/refractory acute myeloid leukemia. <i>Cancer</i> , 2020, 126, 1264-1273.	4.1	15
8	Alisertib plus induction chemotherapy in previously untreated patients with high-risk, acute myeloid leukaemia: a single-arm, phase 2 trial. <i>Lancet Haematology</i> , 2020, 7, e122-e133.	4.6	19
9	Long: molecular tracking of CML with bilineal inv(16) myeloid and del(9) lymphoid blast crisis and durable response to CD19-directed CAR-T therapy. <i>Leukemia</i> , 2020, 34, 3050-3054.	7.2	3
10	Clonal hematopoiesis and measurable residual disease assessment in acute myeloid leukemia. <i>Blood</i> , 2020, 135, 1729-1738.	1.4	80
11	<i>SF3B1</i> -mutant MDS as a distinct disease subtype: a proposal from the International Working Group for the Prognosis of MDS. <i>Blood</i> , 2020, 136, 157-170.	1.4	195
12	Phase I Study of Ixazomib Added to Chemotherapy in the Treatment of Acute Lymphoblastic Leukemia in Older Adults. <i>Blood</i> , 2020, 136, 41-42.	1.4	1
13	Isocitrate dehydrogenase 1 and 2 mutations, 2-hydroxyglutarate levels, and response to standard chemotherapy for patients with newly diagnosed acute myeloid leukemia. <i>Cancer</i> , 2019, 125, 541-549.	4.1	23
14	TP53 mutation status divides myelodysplastic syndromes with complex karyotypes into distinct prognostic subgroups. <i>Leukemia</i> , 2019, 33, 1747-1758.	7.2	195
15	Redirecting T-Cells Against AML in a Multidimensional Targeting Space Using T-Cell Engaging Antibody Circuits (TEAC). <i>Blood</i> , 2019, 134, 2653-2653.	1.4	4
16	Targeting R-loop-associated ATR response in myelodysplastic syndrome. <i>Oncotarget</i> , 2019, 10, 2581-2582.	1.8	9
17	Cellular stressors contribute to the expansion of hematopoietic clones of varying leukemic potential. <i>Nature Communications</i> , 2018, 9, 455.	12.8	150
18	Genomics in childhood acute myeloid leukemia comes of age. <i>Nature Medicine</i> , 2018, 24, 7-9.	30.7	14

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19	High NPM1-mutant allele burden at diagnosis predicts unfavorable outcomes in de novo AML. <i>Blood</i> , 2018, 131, 2816-2825.	1.4	64
20	Immune Escape of Relapsed AML Cells after Allogeneic Transplantation. <i>New England Journal of Medicine</i> , 2018, 379, 2330-2341.	27.0	322
21	Mutation Clearance after Transplantation for Myelodysplastic Syndrome. <i>New England Journal of Medicine</i> , 2018, 379, 1028-1041.	27.0	93
22	Spliceosome Mutations Induce R Loop-Associated Sensitivity to ATR Inhibition in Myelodysplastic Syndromes. <i>Cancer Research</i> , 2018, 78, 5363-5374.	0.9	117
23	Pathobiology of Acute Myeloid Leukemia. , 2018, , 913-923.		1
24	Subclones dominate at MDS progression following allogeneic hematopoietic cell transplant. <i>JCI Insight</i> , 2018, 3, .	5.0	48
25	Phase II Clinical Trial of Alisertib, an Aurora a Kinase Inhibitor, in Combination with Induction Chemotherapy in High-Risk, Untreated Patients with Acute Myeloid Leukemia. <i>Blood</i> , 2018, 132, 766-766.	1.4	9
26	Clinical Outcomes Following Frontline Chemotherapy for Patients with Myeloid Malignancies Harboring Splicing Factor Mutations. <i>Blood</i> , 2018, 132, 4364-4364.	1.4	0
27	Single-Cell RNA-Seq Reveals AML Cellular Hierarchies Relevant to Clinical Outcomes and Immunity. <i>Blood</i> , 2018, 132, 542-542.	1.4	0
28	Potential Barriers to Clinical Trials of New Therapeutics for Myelodysplastic Syndromes: Wide Variation in Risk Definitions and Trial Enrollment Criteria. <i>Blood</i> , 2018, 132, 4378-4378.	1.4	0
29	Mutant U2AF1-expressing cells are sensitive to pharmacological modulation of the spliceosome. <i>Nature Communications</i> , 2017, 8, 14060.	12.8	99
30	Functions of Replication Protein A as a Sensor of R Loops and a Regulator of RNaseH1. <i>Molecular Cell</i> , 2017, 65, 832-847.e4.	9.7	205
31	Phase I study of the aurora A kinase inhibitor alisertib with induction chemotherapy in patients with acute myeloid leukemia. <i>Haematologica</i> , 2017, 102, 719-727.	3.5	33
32	Splicing factor gene mutations in hematologic malignancies. <i>Blood</i> , 2017, 129, 1260-1269.	1.4	99
33	Combined Targeted Therapy for BRAF-Mutant, Treatment-Related Acute Myeloid Leukemia. <i>JCO Precision Oncology</i> , 2017, 1, 1-7.	3.0	3
34	Prevalence and complications associated with off-label use of lenalidomide in older patients with myelodysplastic syndromes (MDS).. <i>Journal of Clinical Oncology</i> , 2017, 35, 7054-7054.	1.6	0
35	Case 37-2016. <i>New England Journal of Medicine</i> , 2016, 375, 2273-2282.	27.0	3
36	The Public Repository of Xenografts Enables Discovery and Randomized Phase II-like Trials in Mice. <i>Cancer Cell</i> , 2016, 29, 574-586.	16.8	227

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37	Rapid expansion of preexisting nonleukemic hematopoietic clones frequently follows induction therapy for de novo AML. <i>Blood</i> , 2016, 127, 893-897.	1.4	94
38	<i>TP53</i> and Decitabine in Acute Myeloid Leukemia and Myelodysplastic Syndromes. <i>New England Journal of Medicine</i> , 2016, 375, 2023-2036.	27.0	663
39	Comprehensive genomic analysis reveals FLT3 activation and a therapeutic strategy for a patient with relapsed adult B-lymphoblastic leukemia. <i>Experimental Hematology</i> , 2016, 44, 603-613.	0.4	44
40	Genomic analysis of germ line and somatic variants in familial myelodysplasia/acute myeloid leukemia. <i>Blood</i> , 2015, 126, 2484-2490.	1.4	207
41	Health care utilization and end-of-life care for older patients with acute myeloid leukemia. <i>Cancer</i> , 2015, 121, 2840-2848.	4.1	113
42	Patterns and functional implications of rare germline variants across 12 cancer types. <i>Nature Communications</i> , 2015, 6, 10086.	12.8	243
43	Quality of life and mood of patients and family caregivers during hospitalization for hematopoietic stem cell transplantation. <i>Cancer</i> , 2015, 121, 951-959.	4.1	157
44	Mutant U2AF1 Expression Alters Hematopoiesis and Pre-mRNA Splicing In Vivo. <i>Cancer Cell</i> , 2015, 27, 631-643.	16.8	259
45	Functional analysis of a chromosomal deletion associated with myelodysplastic syndromes using isogenic human induced pluripotent stem cells. <i>Nature Biotechnology</i> , 2015, 33, 646-655.	17.5	130
46	Detection of Dual IDH1 and IDH2 Mutations by Targeted Next-Generation Sequencing in Acute Myeloid Leukemia and Myelodysplastic Syndromes. <i>Journal of Molecular Diagnostics</i> , 2015, 17, 661-668.	2.8	31
47	Association Between Mutation Clearance After Induction Therapy and Outcomes in Acute Myeloid Leukemia. <i>JAMA - Journal of the American Medical Association</i> , 2015, 314, 811.	7.4	302
48	Role of TP53 mutations in the origin and evolution of therapy-related acute myeloid leukaemia. <i>Nature</i> , 2015, 518, 552-555.	27.8	685
49	New Molecular Abnormalities and Clonal Architecture in AML: From Reciprocal Translocations to Whole-Genome Sequencing. <i>American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting</i> , 2014, , e334-e340.	3.8	10
50	SciClone: Inferring Clonal Architecture and Tracking the Spatial and Temporal Patterns of Tumor Evolution. <i>PLoS Computational Biology</i> , 2014, 10, e1003665.	3.2	400
51	Clonal Architecture of Secondary Acute Myeloid Leukemia Defined by Single-Cell Sequencing. <i>PLoS Genetics</i> , 2014, 10, e1004462.	3.5	115
52	A Call to Action for Acute Lymphoblastic Leukemia. <i>New England Journal of Medicine</i> , 2014, 371, 1064-1066.	27.0	11
53	Functional Heterogeneity of Genetically Defined Subclones in Acute Myeloid Leukemia. <i>Cancer Cell</i> , 2014, 25, 379-392.	16.8	330
54	Integrated analysis of germline and somatic variants in ovarian cancer. <i>Nature Communications</i> , 2014, 5, 3156.	12.8	253

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55	Pathogenic Variants for Mendelian and Complex Traits in Exomes of 6,517 European and African Americans: Implications for the Return of Incidental Results. <i>American Journal of Human Genetics</i> , 2014, 95, 183-193.	6.2	78
56	AML Genomics for the Clinician. <i>Seminars in Hematology</i> , 2014, 51, 322-329.	3.4	6
57	AML Genomics: Introduction. <i>Seminars in Hematology</i> , 2014, 51, 249.	3.4	1
58	Caspase-9 is required for normal hematopoietic development and protection from alkylator-induced DNA damage in mice. <i>Blood</i> , 2014, 124, 3887-3895.	1.4	20
59	Prognostic understanding, quality of life (QOL), and mood in patients undergoing hematopoietic stem cell transplantation (HCT).. <i>Journal of Clinical Oncology</i> , 2014, 32, 219-219.	1.6	4
60	Allele-Specific Effects Of U2AF1 Mutations On Alternative Splicing. <i>Blood</i> , 2013, 122, 2748-2748.	1.4	0
61	Recurrent mutations in the U2AF1 splicing factor in myelodysplastic syndromes. <i>Nature Genetics</i> , 2012, 44, 53-57.	21.4	513
62	Imputation of Exome Sequence Variants into Population- Based Samples and Blood-Cell-Trait-Associated Loci in African Americans: NHLBI GO Exome Sequencing Project. <i>American Journal of Human Genetics</i> , 2012, 91, 794-808.	6.2	123
63	Clonal Architecture of Secondary Acute Myeloid Leukemia. <i>New England Journal of Medicine</i> , 2012, 366, 1090-1098.	27.0	688
64	Clonal evolution in relapsed acute myeloid leukaemia revealed by whole-genome sequencing. <i>Nature</i> , 2012, 481, 506-510.	27.8	1,795
65	The Origin and Evolution of Mutations in Acute Myeloid Leukemia. <i>Cell</i> , 2012, 150, 264-278.	28.9	1,365
66	Mutant U2AF1(S34F) Expression Alters Hematopoiesis in Mice. <i>Blood</i> , 2012, 120, 553-553.	1.4	0
67	Genetics of Myelodysplastic Syndromes: New Insights. <i>Hematology American Society of Hematology Education Program</i> , 2011, 2011, 543-549.	2.5	49
68	Genomics of Acute Myeloid Leukemia. <i>Cancer Journal (Sudbury, Mass)</i> , 2011, 17, 487-491.	2.0	20
69	Use of Whole-Genome Sequencing to Diagnose a Cryptic Fusion Oncogene. <i>JAMA - Journal of the American Medical Association</i> , 2011, 305, 1577.	7.4	233
70	Identification of a Novel <i>TP53</i> ; Cancer Susceptibility Mutation Through Whole-Genome Sequencing of a Patient With Therapy-Related AML. <i>JAMA - Journal of the American Medical Association</i> , 2011, 305, 1568.	7.4	146
71	Sequencing a mouse acute promyelocytic leukemia genome reveals genetic events relevant for disease progression. <i>Journal of Clinical Investigation</i> , 2011, 121, 1445-1455.	8.2	91
72	Complete Sequencing and Comparison of 12 Normal Karyotype M1 AML Genomes with 12 t(15;17) Positive M3-APL Genomes. <i>Blood</i> , 2011, 118, 404-404.	1.4	1

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73	Integrated genomics of susceptibility to alkylator-induced leukemia in mice. BMC Genomics, 2010, 11, 638.	2.8	5
74	Therapy-Related Myelodysplastic Syndrome: Models and Genetics. Biology of Blood and Marrow Transplantation, 2010, 16, S45-S47.	2.0	11
75	<i>DNMT3A</i> Mutations in Acute Myeloid Leukemia. New England Journal of Medicine, 2010, 363, 2424-2433.	27.0	1,777
76	Molecular basis of hematology. , 2010, , 1-26.		0
77	DNA Sequence of the Cancer Genome of a Patient with Therapy-Related Acute Myeloid Leukemia. Blood, 2010, 116, 580-580.	1.4	0
78	Mutations In the DNA Methyltransferase Gene DNMT3A Are Highly Recurrent In Patients with Intermediate Risk Acute Myeloid Leukemia, and Predict Poor Outcomes. Blood, 2010, 116, 99-99.	1.4	9
79	Recurrent DNMT3A Mutations In Patients with Myelodysplastic Syndrome. Blood, 2010, 116, 608-608.	1.4	0
80	Detection of Novel Mutations In MDS/AML by Whole Genome Sequencing. Blood, 2010, 116, 299-299.	1.4	0
81	Murine Models of Human Acute Myeloid Leukemia. Cancer Treatment and Research, 2009, 145, 183-196.	0.5	16
82	Therapy related acute myeloid leukemia in breast cancer survivors, a population-based study. Breast Cancer Research and Treatment, 2009, 118, 593-598.	2.5	41
83	The impact of copy number variation on local gene expression in mouse hematopoietic stem and progenitor cells. Nature Genetics, 2009, 41, 430-437.	21.4	112
84	Recurring Mutations Found by Sequencing an Acute Myeloid Leukemia Genome. New England Journal of Medicine, 2009, 361, 1058-1066.	27.0	2,009
85	Next-generation sequencing of cancer genomes: back to the future. Personalized Medicine, 2009, 6, 653-662.	1.5	26
86	Acquired copy number alterations in adult acute myeloid leukemia genomes. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 12950-12955.	7.1	231
87	Genome-wide association study to identify novel loci associated with therapy-related myeloid leukemia susceptibility. Blood, 2009, 113, 5575-5582.	1.4	93
88	BRCA1 and BRCA2 Nucleotide Variants in Young Women with Therapy Related Acute Myeloid Leukemia.. Blood, 2009, 114, 1102-1102.	1.4	5
89	Integrated Genomic Analysis Implicates Haploinsufficiency of Multiple Chromosome 5q31.2 Genes in De Novo Myelodysplastic Syndromes Pathogenesis. PLoS ONE, 2009, 4, e4583.	2.5	48
90	POU4F1 Is Associated with t(8;21) AML and Contributes Directly to Its Unique Transcriptional Signature.. Blood, 2009, 114, 2623-2623.	1.4	6

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91	DNA sequencing of a cytogenetically normal acute myeloid leukaemia genome. <i>Nature</i> , 2008, 456, 66-72.	27.8	1,275
92	Identification of somatic JAK1 mutations in patients with acute myeloid leukemia. <i>Blood</i> , 2008, 111, 4809-4812.	1.4	84
93	wuHMM: a robust algorithm to detect DNA copy number variation using long oligonucleotide microarray data. <i>Nucleic Acids Research</i> , 2008, 36, e41.	14.5	25
94	Somatic mutations and germline sequence variants in the expressed tyrosine kinase genes of patients with de novo acute myeloid leukemia. <i>Blood</i> , 2008, 111, 4797-4808.	1.4	198
95	Quantitative trait loci associated with susceptibility to therapy-related acute murine promyelocytic leukemia in hCG-PML/RARA transgenic mice. <i>Blood</i> , 2008, 112, 1434-1442.	1.4	11
96	AML1 and Evi1: coconspirators in MDS/AML?. <i>Blood</i> , 2008, 111, 3916-3917.	1.4	0
97	Bcl2, a Candidate Murine Therapy-Related Acute Myeloid Leukemia Susceptibility Factor, Exhibits Strain-Dependent and Alkylator-Responsive Expression.. <i>Blood</i> , 2008, 112, 1499-1499.	1.4	0
98	A High-Resolution Map of Segmental DNA Copy Number Variation in the Mouse Genome. <i>PLoS Genetics</i> , 2007, 3, e3.	3.5	196
99	A Randomized Double-Blind Trial of Hydroxychloroquine for the Prevention of Chronic Graft-versus-Host Disease after Allogeneic Peripheral Blood Stem Cell Transplantation. <i>Biology of Blood and Marrow Transplantation</i> , 2007, 13, 1201-1206.	2.0	24
100	Distinct patterns of mutations occurring in de novo AML versus AML arising in the setting of severe congenital neutropenia. <i>Blood</i> , 2007, 110, 1648-1655.	1.4	88
101	Comprehensive Genomic Copy Number and Sequence Analysis of 28 Chromosome 5q31.2 Candidate Genes in De Novo MDS.. <i>Blood</i> , 2007, 110, 117-117.	1.4	1
102	Identification of Polymorphisms Associated with Susceptibility to Therapy-Related MDS and AML.. <i>Blood</i> , 2007, 110, 15-15.	1.4	5
103	A Phase II Study of Intravenous Azacitidine Alone in Patients with Myelodysplastic Syndromes NCT00384956.. <i>Blood</i> , 2007, 110, 1451-1451.	1.4	1
104	Pharmacogenetics of alkylator-associated acute myeloid leukemia. <i>Pharmacogenomics</i> , 2006, 7, 719-729.	1.3	10
105	Identification of Candidate Alkylator-Induced Cancer Susceptibility Genes by Whole Genome Scanning in Mice. <i>Cancer Research</i> , 2006, 66, 5029-5038.	0.9	44
106	Phase II Trial of the Tyrosine Kinase Inhibitor PKC412 in Advanced Systemic Mastocytosis: Preliminary Results.. <i>Blood</i> , 2006, 108, 3609-3609.	1.4	2
107	Roles of Sca-1 in hematopoietic stem/progenitor cell function. <i>Experimental Hematology</i> , 2005, 33, 836-843.	0.4	108
108	Dominant Negative Effects of the AML1/ETO Fusion Oncoprotein. <i>Cell Cycle</i> , 2005, 4, 33-36.	2.6	4

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109	Sca-1 negatively regulates proliferation and differentiation of muscle cells. <i>Developmental Biology</i> , 2005, 283, 240-252.	2.0	96
110	A Mouse Model of Alkylator-Induced Myelodysplastic Syndrome.. <i>Blood</i> , 2005, 106, 368-368.	1.4	17
111	A high resolution map of segmental DNA copy number variation in the mouse genome. <i>PLoS Genetics</i> , 2005, preprint, e3.	3.5	0
112	A Randomized, Double Blind Trial, of Hydroxychloroquine for the Prevention of Graft-Versus-Host Disease after Allogeneic Peripheral Blood Stem Cell Transplantation.. <i>Blood</i> , 2005, 106, 1800-1800.	1.4	0
113	Stem cell expression of the AML1/ETO fusion protein induces a myeloproliferative disorder in mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2004, 101, 15184-15189.	7.1	81
114	Recruitment of Bone Marrow-Derived Endothelial Cells to Sites of Pancreatic β -Cell Injury. <i>Diabetes</i> , 2004, 53, 91-98.	0.6	172
115	Enhanced green fluorescent protein targeted to the Sca-1 (Ly-6A) locus in transgenic mice results in efficient marking of hematopoietic stem cells in vivo. <i>Experimental Hematology</i> , 2003, 31, 159-167.	0.4	47
116	Long-term outcomes of allogeneic stem cell transplant recipients after calcineurin inhibitor-induced neurotoxicity. <i>British Journal of Haematology</i> , 2003, 123, 110-113.	2.5	28
117	A pilot study of high-throughput, sequence-based mutational profiling of primary human acute myeloid leukemia cell genomes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2003, 100, 14275-14280.	7.1	55
118	A mouse-based strategy for cyclophosphamide pharmacogenomic discovery. <i>Journal of Applied Physiology</i> , 2003, 95, 1352-1360.	2.5	27
119	Sca-1pos Cells in the Mouse Mammary Gland Represent an Enriched Progenitor Cell Population. <i>Developmental Biology</i> , 2002, 245, 42-56.	2.0	491
120	Characterization of Ly-6M, a novel member of the Ly-6 family of hematopoietic proteins. <i>Blood</i> , 2000, 95, 3125-3132.	1.4	19
121	Granzyme A Initiates an Alternative Pathway for Granule-Mediated Apoptosis. <i>Immunity</i> , 1999, 10, 595-605.	14.3	140
122	How do cytotoxic lymphocytes kill their targets?. <i>Current Opinion in Immunology</i> , 1998, 10, 581-587.	5.5	353
123	Recombinant retroviral systems for the analysis of drug resistant HIV. <i>Nucleic Acids Research</i> , 1993, 21, 4836-4842.	14.5	17