

# Stanley B Pounds

## List of Publications by Year in descending order

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218  
papers

18,890  
citations

23567

58  
h-index

12946

131  
g-index

222  
all docs

222  
docs citations

222  
times ranked

23838  
citing authors

#	ARTICLE	IF	CITATIONS
1	Polygenic Ara-C Response Score Identifies Pediatric Patients With Acute Myeloid Leukemia in Need of Chemotherapy Augmentation. <i>Journal of Clinical Oncology</i> , 2022, 40, 772-783.	1.6	7
2	Proteomics: a new era in pediatric acute myeloid leukemia research. <i>Haematologica</i> , 2022, , .	3.5	0
3	Phase Separation Mediates NUP98 Fusion Oncoprotein Leukemic Transformation. <i>Cancer Discovery</i> , 2022, 12, 1152-1169.	9.4	68
4	Integrated Genomic Analysis Identifies <i>UBTF</i> Tandem Duplications as a Recurrent Lesion in Pediatric Acute Myeloid Leukemia. <i>Blood Cancer Discovery</i> , 2022, 3, 194-207.	5.0	38
5	Changes in body mass index, weight, and height in children with acute myeloid leukemia and the associations with outcome. <i>Blood Advances</i> , 2022, 6, 2824-2834.	5.2	3
6	Preclinical and Pilot Study of Type I FLT3 Tyrosine Kinase Inhibitor, Crenolanib, with Sorafenib in Acute Myeloid Leukemia and <i>FLT3</i> -Internal Tandem Duplication. <i>Clinical Cancer Research</i> , 2022, 28, 2536-2546.	7.0	3
7	DNA Methylation-Based Epigenetic Repression of <i>SLC22A4</i> Promotes Resistance to Cytarabine in Acute Myeloid Leukemia. <i>Clinical and Translational Science</i> , 2021, 14, 137-142.	3.1	16
8	SequencErr: measuring and suppressing sequencer errors in next-generation sequencing data. <i>Genome Biology</i> , 2021, 22, 37.	8.8	15
9	Network-based systems pharmacology reveals heterogeneity in LCK and BCL2 signaling and therapeutic sensitivity of T-cell acute lymphoblastic leukemia. <i>Nature Cancer</i> , 2021, 2, 284-299.	13.2	70
10	The Common Germline <i>TP53-R337H</i> Mutation Is Hypomorphic and Confers Incomplete Penetrance and Late Tumor Onset in a Mouse Model. <i>Cancer Research</i> , 2021, 81, 2442-2456.	0.9	9
11	The acquisition of molecular drivers in pediatric therapy-related myeloid neoplasms. <i>Nature Communications</i> , 2021, 12, 985.	12.8	31
12	Gene-set distance analysis (GSDA): a powerful tool for gene-set association analysis. <i>BMC Bioinformatics</i> , 2021, 22, 207.	2.6	0
13	Molecular classification improves risk assessment in adult <i>BCR-ABL1</i> negative B-ALL. <i>Blood</i> , 2021, 138, 948-958.	1.4	59
14	Global Proteomic Profiling of Pediatric AML: A Pilot Study. <i>Cancers</i> , 2021, 13, 3161.	3.7	6
15	Cellular Metabolomics Profiles Associated With Drug Chemosensitivity in AML. <i>Frontiers in Oncology</i> , 2021, 11, 678008.	2.8	8
16	Integrative Genomic Analysis of Pediatric Myeloid-Related Acute Leukemias Identifies Novel Subtypes and Prognostic Indicators. <i>Blood Cancer Discovery</i> , 2021, 2, 586-599.	5.0	21
17	Clinical Features and Cytoreduction Therapy in Children with Newly Diagnosed Acute Myeloid Leukemia and Hyperleukocytosis. <i>Blood</i> , 2021, 138, 2295-2295.	1.4	0
18	Integrated Genomic Analysis Identifies <i>UBTF</i> Tandem Duplications As a Subtype-Defining Lesion in Pediatric Acute Myeloid Leukemia. <i>Blood</i> , 2021, 138, LBA-4-LBA-4.	1.4	0

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19	A six-gene leukemic stem cell score identifies high risk pediatric acute myeloid leukemia. <i>Leukemia</i> , 2020, 34, 735-745.	7.2	56
20	DNA Methylation Clusters and Their Relation to Cytogenetic Features in Pediatric AML. <i>Cancers</i> , 2020, 12, 3024.	3.7	5
21	Venetoclax in combination with cytarabine with or without idarubicin in children with relapsed or refractory acute myeloid leukaemia: a phase 1, dose-escalation study. <i>Lancet Oncology</i> , The, 2020, 21, 551-560.	10.7	92
22	Integrative genomic analyses reveal mechanisms of glucocorticoid resistance in acute lymphoblastic leukemia. <i>Nature Cancer</i> , 2020, 1, 329-344.	13.2	44
23	Mutational Landscape and Patterns of Clonal Evolution in Relapsed Pediatric Acute Lymphoblastic Leukemia. <i>Blood Cancer Discovery</i> , 2020, 1, 96-111.	5.0	93
24	Pharmacogenomics of intracellular methotrexate polyglutamates in patients' leukemia cells in vivo. <i>Journal of Clinical Investigation</i> , 2020, 130, 6600-6615.	8.2	18
25	The St. Jude STEM Clubs: An Afterschool STEM Club for Upper Elementary School Students in Memphis, TN. <i>Journal of STEM Outreach</i> , 2020, 3, .	0.5	0
26	Uncovering the Genomic Landscape in Newly Diagnosed and Relapsed Pediatric Cytogenetically Normal FLT3-ITD AML. <i>Clinical and Translational Science</i> , 2019, 12, 641-647.	3.1	12
27	Clofarabine Can Replace Anthracyclines and Etoposide in Remission Induction Therapy for Childhood Acute Myeloid Leukemia: The AML08 Multicenter, Randomized Phase III Trial. <i>Journal of Clinical Oncology</i> , 2019, 37, 2072-2081.	1.6	34
28	Sorafenib Population Pharmacokinetics and Skin Toxicities in Children and Adolescents with Refractory/Relapsed Leukemia or Solid Tumor Malignancies. <i>Clinical Cancer Research</i> , 2019, 25, 7320-7330.	7.0	14
29	Bacterial Factors Required for Transmission of <i>Streptococcus pneumoniae</i> in Mammalian Hosts. <i>Cell Host and Microbe</i> , 2019, 25, 884-891.e6.	11.0	48
30	MicroRNAs Mediated Regulation of Expression of Nucleoside Analog Pathway Genes in Acute Myeloid Leukemia. <i>Genes</i> , 2019, 10, 319.	2.4	6
31	A phase II clinical trial of adoptive transfer of haploidentical natural killer cells for consolidation therapy of pediatric acute myeloid leukemia. , 2019, 7, 81.		74
32	Genomic subtyping and therapeutic targeting of acute erythroleukemia. <i>Nature Genetics</i> , 2019, 51, 694-704.	21.4	97
33	H3.3 K27M depletion increases differentiation and extends latency of diffuse intrinsic pontine glioma growth in vivo. <i>Acta Neuropathologica</i> , 2019, 137, 637-655.	7.7	85
34	Forty-five patient-derived xenografts capture the clinical and biological heterogeneity of Wilms tumor. <i>Nature Communications</i> , 2019, 10, 5806.	12.8	27
35	DNA Methylation Profiling Reveals Prognostically Significant Groups in Pediatric Adrenocortical Tumors: A Report From the International Pediatric Adrenocortical Tumor Registry. <i>JCO Precision Oncology</i> , 2019, 3, 1-21.	3.0	6
36	Histone H3.3 K27M Accelerates Spontaneous Brainstem Glioma and Drives Restricted Changes in Bivalent Gene Expression. <i>Cancer Cell</i> , 2019, 35, 140-155.e7.	16.8	194

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37	PAX5-driven subtypes of B-progenitor acute lymphoblastic leukemia. <i>Nature Genetics</i> , 2019, 51, 296-307.	21.4	384
38	Integrative Analysis of Pediatric Acute Leukemia Identifies Immature Subtypes That Span a T Lineage and Myeloid Continuum with Distinct Prognoses. <i>Blood</i> , 2019, 134, 918-918.	1.4	1
39	Integrated Transcriptomic and Genomic Sequencing Identifies Prognostic Constellations of Driver Mutations in Acute Myeloid Leukemia and Myelodysplastic Syndromes. <i>Blood</i> , 2019, 134, LBA-4-LBA-4.	1.4	20
40	OR02-1 DNA Methylation Profiling in Pediatric Adrenocortical Tumors Reveals Distinct Methylation Signatures with Prognostic Significance: A Report from the International Pediatric Adrenocortical Tumor Registry. <i>Journal of the Endocrine Society</i> , 2019, 3, .	0.2	0
41	SAT-LB058 Effect of a Genetic Modifier of Cancer Risk in TP53 Mutation Carriers. <i>Journal of the Endocrine Society</i> , 2019, 3, .	0.2	0
42	Venetoclax in Combination with High-Dose Chemotherapy Is Active and Well-Tolerated in Children with Relapsed or Refractory Acute Myeloid Leukemia. <i>Blood</i> , 2019, 134, 178-178.	1.4	0
43	The Genomic Landscape of Childhood Acute Lymphoblastic Leukemia. <i>Blood</i> , 2019, 134, 649-649.	1.4	5
44	A 5-Gene Ara-C, Daunorubicin and Etoposide (ADE) Drug Response Score As a Prognostic Tool to Predict AML Treatment Outcome. <i>Blood</i> , 2019, 134, 1429-1429.	1.4	1
45	Gut Microbiome Composition Predicts Infection Risk During Chemotherapy in Children With Acute Lymphoblastic Leukemia. <i>Clinical Infectious Diseases</i> , 2018, 67, 541-548.	5.8	122
46	Pan-cancer genome and transcriptome analyses of 1,699 paediatric leukaemias and solid tumours. <i>Nature</i> , 2018, 555, 371-376.	27.8	649
47	Malignant rhabdoid tumors originating within and outside the central nervous system are clinically and molecularly heterogeneous. <i>Acta Neuropathologica</i> , 2018, 136, 315-326.	7.7	26
48	Bithalamic gliomas may be molecularly distinct from their unilateral high-grade counterparts. <i>Brain Pathology</i> , 2018, 28, 112-120.	4.1	26
49	The genetic basis and cell of origin of mixed phenotype acute leukaemia. <i>Nature</i> , 2018, 562, 373-379.	27.8	236
50	POST: A framework for set-based association analysis in high-dimensional data. <i>Methods</i> , 2018, 145, 76-81.	3.8	1
51	Statistical selection of biological models for genome-wide association analyses. <i>Methods</i> , 2018, 145, 67-75.	3.8	5
52	Comprehensive Ara-C SNP score predicts leukemic cell intracellular ara-CTP levels in pediatric acute myeloid leukemia patients. <i>Pharmacogenomics</i> , 2018, 19, 1101-1110.	1.3	7
53	Molecular heterogeneity and CXorf67 alterations in posterior fossa group A (PFA) ependymomas. <i>Acta Neuropathologica</i> , 2018, 136, 211-226.	7.7	199
54	Metabolomics Profiling Reveals Markers for Chemosensitivity and Clinical Outcomes in Pediatric AML Patients. <i>Blood</i> , 2018, 132, 1536-1536.	1.4	5

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55	Integrated epigenetic and genetic analysis identifies markers of prognostic significance in pediatric acute myeloid leukemia. <i>Oncotarget</i> , 2018, 9, 26711-26723.	1.8	26
56	Genome-wide association analysis identifies SNPs predictive of <i>in vitro</i> leukemic cell sensitivity to cytarabine in pediatric AML. <i>Oncotarget</i> , 2018, 9, 34859-34875.	1.8	12
57	Pediatric LSC3 (pLSC3) Score Derived from DNMT3B-CD34-GPR56 As a Prognostic Tool to Predict AML Patient Outcome: Results from Two Independent Pediatric AML Cohorts. <i>Blood</i> , 2018, 132, 290-290.	1.4	1
58	Characterization of Novel Subtypes in B Progenitor Acute Lymphoblastic Leukemia. <i>Blood</i> , 2018, 132, 565-565.	1.4	14
59	Integrated Genome Wide Association Study (GWAS) Identifies SNPs Associated with Outcome in Pediatric AML. <i>Blood</i> , 2018, 132, 2758-2758.	1.4	2
60	Transcriptome profiling of patient derived xenograft models established from pediatric acute myeloid leukemia patients confirm maintenance of FLT3-ITD mutation. <i>Leukemia and Lymphoma</i> , 2017, 58, 247-250.	1.3	5
61	A Robust and Powerful Set-Valued Approach to Rare Variant Association Analyses of Secondary Traits in Case-Control Sequencing Studies. <i>Genetics</i> , 2017, 205, 1049-1062.	2.9	4
62	OCTN1 Is a High-Affinity Carrier of Nucleoside Analogues. <i>Cancer Research</i> , 2017, 77, 2102-2111.	0.9	41
63	MLF1 is a proapoptotic antagonist of HOP complex-mediated survival. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2017, 1864, 719-727.	4.1	5
64	High Frequency and Poor Outcome of Philadelphia Chromosome-Like Acute Lymphoblastic Leukemia in Adults. <i>Journal of Clinical Oncology</i> , 2017, 35, 394-401.	1.6	326
65	Decreased relapsed rate and treatment-related mortality contribute to improved outcomes for pediatric acute myeloid leukemia in successive clinical trials. <i>Cancer</i> , 2017, 123, 3791-3798.	4.1	34
66	Contribution of the TP53 R337H mutation to the cancer burden in southern Brazil: Insights from the study of 55 families of children with adrenocortical tumors. <i>Cancer</i> , 2017, 123, 3150-3158.	4.1	26
67	The genomic landscape of pediatric and young adult T-lineage acute lymphoblastic leukemia. <i>Nature Genetics</i> , 2017, 49, 1211-1218.	21.4	693
68	Statistical selection of biological models for genome-wide association analyses. , 2017, , .		0
69	POST: A framework for set-based association analysis in high-dimensional data. , 2017, , .		0
70	Identification of Clinical and Biologic Correlates Associated With Outcome in Children With Adrenocortical Tumors Without Germline TP53 Mutations: A St Jude Adrenocortical Tumor Registry and Children's Oncology Group Study. <i>Journal of Clinical Oncology</i> , 2017, 35, 3956-3963.	1.6	33
71	Hypoxia-induced upregulation of BMX kinase mediates therapeutic resistance in acute myeloid leukemia. <i>Journal of Clinical Investigation</i> , 2017, 128, 369-380.	8.2	39
72	Higher-order oligomerization promotes localization of SPOP to liquid nuclear speckles. <i>EMBO Journal</i> , 2016, 35, 1254-1275.	7.8	172

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73	Quantitative Assessment of Commutability for Clinical Viral Load Testing Using a Digital PCR-Based Reference Standard. <i>Journal of Clinical Microbiology</i> , 2016, 54, 1616-1623.	3.9	16
74	Comparative Evaluation of Four Real-Time PCR Methods for the Quantitative Detection of Epstein-Barr Virus from Whole Blood Specimens. <i>Journal of Molecular Diagnostics</i> , 2016, 18, 527-534.	2.8	15
75	Comparative Performance of Reagents and Platforms for Quantitation of Cytomegalovirus DNA by Digital PCR. <i>Journal of Clinical Microbiology</i> , 2016, 54, 2602-2608.	3.9	25
76	Multi-organ Mapping of Cancer Risk. <i>Cell</i> , 2016, 166, 1132-1146.e7.	28.9	128
77	TERT promoter mutations and prognosis in solitary fibrous tumor. <i>Modern Pathology</i> , 2016, 29, 1511-1522.	5.5	88
78	Rapid Antimicrobial Susceptibility Testing Using Forward Laser Light Scatter Technology. <i>Journal of Clinical Microbiology</i> , 2016, 54, 2701-2706.	3.9	36
79	The genomic landscape of core-binding factor acute myeloid leukemias. <i>Nature Genetics</i> , 2016, 48, 1551-1556.	21.4	215
80	CC-PROMISE effectively integrates two forms of molecular data with multiple biologically related endpoints. <i>BMC Bioinformatics</i> , 2016, 17, 382.	2.6	5
81	Prognostic Significance of Major Histocompatibility Complex Class II Expression in Pediatric Adrenocortical Tumors: A St. Jude and Children's Oncology Group Study. <i>Clinical Cancer Research</i> , 2016, 22, 6247-6255.	7.0	22
82	Inherited variation in OATP1B1 is associated with treatment outcome in acute myeloid leukemia. <i>Clinical Pharmacology and Therapeutics</i> , 2016, 99, 651-660.	4.7	27
83	Comparative evaluation of whole blood versus plasma for quantitative detection of cytomegalovirus using an automated system. <i>Diagnostic Microbiology and Infectious Disease</i> , 2016, 85, 23-25.	1.8	6
84	Gliomatosis cerebri in children shares molecular characteristics with other pediatric gliomas. <i>Acta Neuropathologica</i> , 2016, 131, 299-307.	7.7	38
85	Clinical significance of <i>in vivo</i> cytarabine-induced gene expression signature in AML. <i>Leukemia and Lymphoma</i> , 2016, 57, 909-920.	1.3	7
86	The Genomic Landscape of Childhood and Adult Acute Erythroid Leukemia. <i>Blood</i> , 2016, 128, 39-39.	1.4	2
87	Genomic Landscape of Pediatric Mixed Phenotype Acute Leukemia. <i>Blood</i> , 2016, 128, 454-454.	1.4	4
88	Linking Subclonal Genetic Diversity with Functional Heterogeneity Identifies Diagnosis Subclones Destined to Relapse. <i>Blood</i> , 2016, 128, 605-605.	1.4	0
89	Genomic Profiling Identifies Novel Mutations and Fusion Genes in Newly Diagnosed and Relapsed Pediatric FLT3-ITD-Positive AML. <i>Blood</i> , 2016, 128, 2838-2838.	1.4	0
90	MPTH-26 MOLECULAR REFINEMENT OF PEDIATRIC POSTERIOR FOSSA EPENDYMOMA. <i>Neuro-Oncology</i> , 2015, 17, v144.1-v144.	1.2	0

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91	SVSI: Fast and Powerful Set-Valued System Identification Approach to Identifying Rare Variants in Sequencing Studies for Ordered Categorical Traits. <i>Annals of Human Genetics</i> , 2015, 79, 294-309.	0.8	9
92	The Genomic Landscape of Childhood and Adolescent Melanoma. <i>Journal of Investigative Dermatology</i> , 2015, 135, 816-823.	0.7	148
93	Comparative Evaluation of Three Commercial Quantitative Cytomegalovirus Standards by Use of Digital and Real-Time PCR. <i>Journal of Clinical Microbiology</i> , 2015, 53, 1500-1505.	3.9	37
94	Genomic landscape of paediatric adrenocortical tumours. <i>Nature Communications</i> , 2015, 6, 6302.	12.8	166
95	The landscape of somatic mutations in infant MLL-rearranged acute lymphoblastic leukemias. <i>Nature Genetics</i> , 2015, 47, 330-337.	21.4	405
96	CONSERTING: integrating copy-number analysis with structural-variation detection. <i>Nature Methods</i> , 2015, 12, 527-530.	19.0	68
97	Commutability of the First World Health Organization International Standard for Human Cytomegalovirus. <i>Journal of Clinical Microbiology</i> , 2015, 53, 3325-3333.	3.9	76
98	MicroRNA-mRNA Pairs Associated with Outcome in AML: From In Vitro Cell-Based Studies to AML Patients. <i>Frontiers in Pharmacology</i> , 2015, 6, 324.	3.5	19
99	The methylome of pediatric acute myeloid leukemia.. <i>Journal of Clinical Oncology</i> , 2015, 33, 10027-10027.	1.6	1
100	Abstract 5464: Host variation in OATP1B1 is associated with treatment outcome in pediatric AML. , 2015, , .		0
101	Methylation of DNMT3B Strongly Associates with the Methylome, Cytogenetic Risk Groups, and Prognosis of Pediatric Acute Myeloid Leukemia. <i>Blood</i> , 2015, 126, 2434-2434.	1.4	0
102	Identification and Characterization of Novel Fusion Proteins in Pediatric Acute Megakaryoblastic Leukemia. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2014, 14, S123-S124.	0.4	0
103	Feasibility, efficacy, and adverse effects of outpatient antibacterial prophylaxis in children with acute myeloid leukemia. <i>Cancer</i> , 2014, 120, 1985-1992.	4.1	53
104	Definition of cure in childhood acute myeloid leukemia. <i>Cancer</i> , 2014, 120, 2490-2496.	4.1	12
105	A therapeutic trial of decitabine and vorinostat in combination with chemotherapy for relapsed/refractory acute lymphoblastic leukemia. <i>American Journal of Hematology</i> , 2014, 89, 889-895.	4.1	82
106	The most informative spacing test effectively discovers biologically relevant outliers or multiple modes in expression. <i>Bioinformatics</i> , 2014, 30, 1400-1408.	4.1	10
107	An R package that automatically collects and archives details for reproducible computing. <i>BMC Bioinformatics</i> , 2014, 15, 138.	2.6	14
108	The genomic landscape of diffuse intrinsic pontine glioma and pediatric non-brainstem high-grade glioma. <i>Nature Genetics</i> , 2014, 46, 444-450.	21.4	871

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109	A New System Identification Approach to Identify Genetic Variants in Sequencing Studies for a Binary Phenotype. <i>Human Heredity</i> , 2014, 78, 104-116.	0.8	19
110	A Conditional Approach for Regression Analysis of Longitudinal Data with Informative Observation Time and Non-negligible Observation Duration. <i>Communications in Statistics - Theory and Methods</i> , 2014, 43, 4998-5011.	1.0	1
111	RB1 gene inactivation by chromothripsis in human retinoblastoma. <i>Oncotarget</i> , 2014, 5, 438-450.	1.8	104
112	Abstract LB-63: Cell ground state dictates cancer susceptibility across organs. , 2014, , .		0
113	Abstract PR03: The genomic landscape of diffuse intrinsic pontine glioma and pediatric non-brainstem high-grade glioma. , 2014, , .		2
114	High-Throughput, High-Content siRNA/Drug Modifier Screen for Validation of Transcriptional Profiles Predictive of Cytarabine Response in AML. <i>Blood</i> , 2014, 124, 3615-3615.	1.4	0
115	Gemtuzumab ozogamicin can reduce minimal residual disease in patients with childhood acute myeloid leukemia. <i>Cancer</i> , 2013, 119, 4036-4043.	4.1	41
116	Comparison of Droplet Digital PCR to Real-Time PCR for Quantitative Detection of Cytomegalovirus. <i>Journal of Clinical Microbiology</i> , 2013, 51, 540-546.	3.9	280
117	Joint analysis of longitudinal data and recurrent episodes data with application to medical cost analysis. <i>Biometrical Journal</i> , 2013, 55, 5-16.	1.0	5
118	Prognostic features in acute megakaryoblastic leukemia in children without Down syndrome: a report from the AML02 multicenter trial and the Children's Oncology Group Study POG 9421. <i>Leukemia</i> , 2013, 27, 731-734.	7.2	41
119	Comparison of two multiplexed PCR assays for the detection of HSV-1, HSV-2, and VZV with extracted and unextracted cutaneous and mucosal specimens. <i>Journal of Clinical Virology</i> , 2013, 58, 84-88.	3.1	17
120	A genomic random interval model for statistical analysis of genomic lesion data. <i>Bioinformatics</i> , 2013, 29, 2088-2095.	4.1	17
121	PAIR: paired allelic log-intensity-ratio-based normalization method for SNP-CGH arrays. <i>Bioinformatics</i> , 2013, 29, 299-307.	4.1	6
122	Clinical Significance of CD33 Nonsynonymous Single-Nucleotide Polymorphisms in Pediatric Patients with Acute Myeloid Leukemia Treated with Gemtuzumab-Ozogamicin-Containing Chemotherapy. <i>Clinical Cancer Research</i> , 2013, 19, 1620-1627.	7.0	58
123	<i>RRM1</i> and <i>RRM2</i> pharmacogenetics: association with phenotypes in HapMap cell lines and acute myeloid leukemia patients. <i>Pharmacogenomics</i> , 2013, 14, 1449-1466.	1.3	27
124	An empirical Bayes approach for analysis of diverse periodic trends in time-course gene expression data. <i>Bioinformatics</i> , 2013, 29, 182-188.	4.1	4
125	Comprehensive genetic analysis of cytarabine sensitivity in a cell-based model identifies polymorphisms associated with outcome in AML patients. <i>Blood</i> , 2013, 121, 4366-4376.	1.4	42
126	Cross-species genomic and epigenomic landscape of retinoblastoma. <i>Oncotarget</i> , 2013, 4, 844-859.	1.8	37



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127	Statistical Methods for Overdispersion in mRNA-Seq Count Data. <i>Open Bioinformatics Journal</i> , 2013, 7, 34-40.	1.0	8
128	Empirical Bayesian Selection of Hypothesis Testing Procedures for Analysis of Sequence Count Expression Data. <i>Statistical Applications in Genetics and Molecular Biology</i> , 2012, 11, .	0.6	12
129	High-resolution genomic profiling of adult and pediatric core-binding factor acute myeloid leukemia reveals new recurrent genomic alterations. <i>Blood</i> , 2012, 119, e67-e75.	1.4	66
130	An Inv(16)(p13.3q24.3)-Encoded CBFA2T3-GLIS2 Fusion Protein Defines an Aggressive Subtype of Pediatric Acute Megakaryoblastic Leukemia. <i>Cancer Cell</i> , 2012, 22, 683-697.	16.8	213
131	Comparative Analysis of Different Approaches to Measure Treatment Response in Acute Myeloid Leukemia. <i>Journal of Clinical Oncology</i> , 2012, 30, 3625-3632.	1.6	188
132	High-resolution genomic profiling of chronic lymphocytic leukemia reveals new recurrent genomic alterations. <i>Blood</i> , 2012, 120, 4783-4794.	1.4	179
133	The genetic basis of early T-cell precursor acute lymphoblastic leukaemia. <i>Nature</i> , 2012, 481, 157-163.	27.8	1,430
134	A novel retinoblastoma therapy from genomic and epigenetic analyses. <i>Nature</i> , 2012, 481, 329-334.	27.8	442
135	Novel mutations target distinct subgroups of medulloblastoma. <i>Nature</i> , 2012, 488, 43-48.	27.8	742
136	Effect of body mass index on the outcome of children with acute myeloid leukemia. <i>Cancer</i> , 2012, 118, 5989-5996.	4.1	56
137	Treatment outcome in older patients with childhood acute myeloid leukemia. <i>Cancer</i> , 2012, 118, 6253-6259.	4.1	32
138	A Mouse Model of the Most Aggressive Subgroup of Human Medulloblastoma. <i>Cancer Cell</i> , 2012, 21, 168-180.	16.8	250
139	Analysis of MDM2 and MDM4 Single Nucleotide Polymorphisms, mRNA Splicing and Protein Expression in Retinoblastoma. <i>PLoS ONE</i> , 2012, 7, e42739.	2.5	68
140	Abstract 1434: A mouse model of the most aggressive subgroup of human medulloblastoma. , 2012, , .		0
141	Abstract 2487: CONSERTING: an accurate method for detecting focal and gross somatic copy number alterations in cancer genome by next generation sequencing. , 2012, , .		0
142	Abstract 4867: Identification of an inv(16)-encoded CBFA2T3-GLIS2 fusion protein in 34% of non-infant acute megakaryoblastic leukemias: A report from the Pediatric Cancer Genome Project. , 2012, , .		0
143	Abstract LB-228: Association of XRCC1 SNPs with clinical response in AML patients. , 2012, , .		0
144	Cytarabine-Induced Gene Expression Signatures in AML Patients and Its Association with Clinical Outcome.. <i>Blood</i> , 2012, 120, 2470-2470.	1.4	0

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145	Integrated analysis of pharmacologic, clinical and SNP microarray data using Projection Onto the Most Interesting Statistical Evidence with Adaptive Permutation Testing. <i>International Journal of Data Mining and Bioinformatics</i> , 2011, 5, 143.	0.1	11
146	Impact of genetic variation in FKBP5 on clinical response in pediatric acute myeloid leukemia patients: a pilot study. <i>Leukemia</i> , 2011, 25, 1354-1356.	7.2	19
147	Regression analysis of longitudinal data with informative observation times and application to medical cost data. <i>Statistics in Medicine</i> , 2011, 30, 1429-1440.	1.6	8
148	Randomized trial of 2 dosages of prophylactic granulocyte "colony" stimulating factor after induction chemotherapy in pediatric acute myeloid leukemia. <i>Cancer</i> , 2011, 117, 1313-1320.	4.1	13
149	Identification of predictive markers of cytarabine response in AML by integrative analysis of gene-expression profiles with multiple phenotypes. <i>Pharmacogenomics</i> , 2011, 12, 327-339.	1.3	27
150	IDH1 and IDH2 mutations in pediatric acute leukemia. <i>Leukemia</i> , 2011, 25, 1570-1577.	7.2	80
151	Genetic Variants in Cytosolic 5' Nucleotidase II Are Associated with Its Expression and Cytarabine Sensitivity in HapMap Cell Lines and in Patients with Acute Myeloid Leukemia. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2011, 339, 9-23.	2.5	50
152	Phase I Pharmacokinetic and Pharmacodynamic Study of the Multikinase Inhibitor Sorafenib in Combination With Clofarabine and Cytarabine in Pediatric Relapsed/Refractory Leukemia. <i>Journal of Clinical Oncology</i> , 2011, 29, 3293-3300.	1.6	142
153	Activity of the Multikinase Inhibitor Sorafenib in Combination With Cytarabine in Acute Myeloid Leukemia. <i>Journal of the National Cancer Institute</i> , 2011, 103, 893-905.	6.3	50
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