

Scott A Armstrong

List of Publications by Year in descending order

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133
papers

14,868
citations

34105

52
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19749

117
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137
all docs

137
docs citations

137
times ranked

17533
citing authors

#	ARTICLE	IF	CITATIONS
1	MLL translocations specify a distinct gene expression profile that distinguishes a unique leukemia. <i>Nature Genetics</i> , 2002, 30, 41-47.	21.4	1,720
2	Transformation from committed progenitor to leukaemia stem cell initiated by MLL- Δ AF9. <i>Nature</i> , 2006, 442, 818-822.	27.8	1,317
3	MLL translocations, histone modifications and leukaemia stem-cell development. <i>Nature Reviews Cancer</i> , 2007, 7, 823-833.	28.4	1,039
4	Selective Killing of Mixed Lineage Leukemia Cells by a Potent Small-Molecule DOT1L Inhibitor. <i>Cancer Cell</i> , 2011, 20, 53-65.	16.8	842
5	MLL-Rearranged Leukemia Is Dependent on Aberrant H3K79 Methylation by DOT1L. <i>Cancer Cell</i> , 2011, 20, 66-78.	16.8	791
6	H3K79 Methylation Profiles Define Murine and Human MLL-AF4 Leukemias. <i>Cancer Cell</i> , 2008, 14, 355-368.	16.8	494
7	Inhibition of FLT3 in MLL. <i>Cancer Cell</i> , 2003, 3, 173-183.	16.8	389
8	The DOT1L inhibitor pinometostat reduces H3K79 methylation and has modest clinical activity in adult acute leukemia. <i>Blood</i> , 2018, 131, 2661-2669.	1.4	313
9	Acute Myelogenous Leukemia-Induced Sympathetic Neuropathy Promotes Malignancy in an Altered Hematopoietic Stem Cell Niche. <i>Cell Stem Cell</i> , 2014, 15, 365-375.	11.1	308
10	Mediator kinase inhibition further activates super-enhancer-associated genes in AML. <i>Nature</i> , 2015, 526, 273-276.	27.8	307
11	Modulation of splicing catalysis for therapeutic targeting of leukemia with mutations in genes encoding spliceosomal proteins. <i>Nature Medicine</i> , 2016, 22, 672-678.	30.7	301
12	HOXA9 is required for survival in human MLL-rearranged acute leukemias. <i>Blood</i> , 2009, 113, 2375-2385.	1.4	292
13	A dominant-negative effect drives selection of TP53 missense mutations in myeloid malignancies. <i>Science</i> , 2019, 365, 599-604.	12.6	265
14	FLT3 mutations in childhood acute lymphoblastic leukemia. <i>Blood</i> , 2004, 103, 3544-3546.	1.4	235
15	A Menin-MLL Inhibitor Induces Specific Chromatin Changes and Eradicates Disease in Models of MLL-Rearranged Leukemia. <i>Cancer Cell</i> , 2019, 36, 660-673.e11.	16.8	231
16	PI3K pathway regulates ER-dependent transcription in breast cancer through the epigenetic regulator KMT2D. <i>Science</i> , 2017, 355, 1324-1330.	12.6	217
17	Genetic and Pharmacologic Inhibition of β -Catenin Targets Imatinib-Resistant Leukemia Stem Cells in CML. <i>Cell Stem Cell</i> , 2012, 10, 412-424.	11.1	209
18	Degradation of the BAF Complex Factor BRD9 by Heterobifunctional Ligands. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 5738-5743.	13.8	207

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19	ENL links histone acetylation to oncogenic gene expression in acute myeloid leukaemia. <i>Nature</i> , 2017, 543, 265-269.	27.8	203
20	DOT1L inhibits SIRT1-mediated epigenetic silencing to maintain leukemic gene expression in MLL-rearranged leukemia. <i>Nature Medicine</i> , 2015, 21, 335-343.	30.7	200
21	DNMT3A mutations promote anthracycline resistance in acute myeloid leukemia via impaired nucleosome remodeling. <i>Nature Medicine</i> , 2016, 22, 1488-1495.	30.7	195
22	DNA-damage-induced differentiation of leukaemic cells as an anti-cancer barrier. <i>Nature</i> , 2014, 514, 107-111.	27.8	174
23	A UTX-MLL4-p300 Transcriptional Regulatory Network Coordinately Shapes Active Enhancer Landscapes for Eliciting Transcription. <i>Molecular Cell</i> , 2017, 67, 308-321.e6.	9.7	172
24	Mutations in epigenetic regulators including SETD2 are gained during relapse in paediatric acute lymphoblastic leukaemia. <i>Nature Communications</i> , 2014, 5, 3469.	12.8	171
25	Targeting Chromatin Regulators Inhibits Leukemogenic Gene Expression in <i>NPM1</i> Mutant Leukemia. <i>Cancer Discovery</i> , 2016, 6, 1166-1181.	9.4	171
26	CDK9-mediated transcription elongation is required for MYC addiction in hepatocellular carcinoma. <i>Genes and Development</i> , 2014, 28, 1800-1814.	5.9	167
27	Myeloid progenitor cluster formation drives emergency and leukaemic myelopoiesis. <i>Nature</i> , 2017, 544, 53-58.	27.8	155
28	AF10 Regulates Progressive H3K79 Methylation and HOX Gene Expression in Diverse AML Subtypes. <i>Cancer Cell</i> , 2014, 26, 896-908.	16.8	153
29	Leukemic transformation by the MLL-AF6 fusion oncogene requires the H3K79 methyltransferase Dot1l. <i>Blood</i> , 2013, 121, 2533-2541.	1.4	149
30	Therapeutic targeting of preleukemia cells in a mouse model of <i>NPM1</i> mutant acute myeloid leukemia. <i>Science</i> , 2020, 367, 586-590.	12.6	145
31	Requirement for CDK6 in MLL-rearranged acute myeloid leukemia. <i>Blood</i> , 2014, 124, 13-23.	1.4	139
32	Targeted degradation of BRD9 reverses oncogenic gene expression in synovial sarcoma. <i>ELife</i> , 2018, 7, .	6.0	125
33	Exploiting the Epigenome to Control Cancer-Promoting Gene-Expression Programs. <i>Cancer Cell</i> , 2016, 29, 464-476.	16.8	122
34	Conditional MLL-CBP targets GMP and models therapy-related myeloproliferative disease. <i>EMBO Journal</i> , 2005, 24, 368-381.	7.8	111
35	NUP98 Fusion Proteins Interact with the NSL and MLL1 Complexes to Drive Leukemogenesis. <i>Cancer Cell</i> , 2016, 30, 863-878.	16.8	111
36	SETD2 alterations impair DNA damage recognition and lead to resistance to chemotherapy in leukemia. <i>Blood</i> , 2017, 130, 2631-2641.	1.4	102

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37	Targeting DOT1L and HOX gene expression in MLL-rearranged leukemia and beyond. <i>Experimental Hematology</i> , 2015, 43, 673-684.	0.4	97
38	Hematopoietic Differentiation Is Required for Initiation of Acute Myeloid Leukemia. <i>Cell Stem Cell</i> , 2015, 17, 611-623.	11.1	97
39	A Non-catalytic Function of SETD1A Regulates Cyclin K and the DNA Damage Response. <i>Cell</i> , 2018, 172, 1007-1021.e17.	28.9	97
40	HOXA9 Reprograms the Enhancer Landscape to Promote Leukemogenesis. <i>Cancer Cell</i> , 2018, 34, 643-658.e5.	16.8	94
41	LSD1 inhibition exerts its antileukemic effect by recommissioning PU.1- and C/EBP β -dependent enhancers in AML. <i>Blood</i> , 2018, 131, 1730-1742.	1.4	92
42	The PZP Domain of AF10 Senses Unmodified H3K27 to Regulate DOT1L-Mediated Methylation of H3K79. <i>Molecular Cell</i> , 2015, 60, 319-327.	9.7	78
43	JMJD1C is required for the survival of acute myeloid leukemia by functioning as a coactivator for key transcription factors. <i>Genes and Development</i> , 2015, 29, 2123-2139.	5.9	76
44	Functional screen of MSI2 interactors identifies an essential role for SYNCRIP in myeloid leukemia stem cells. <i>Nature Genetics</i> , 2017, 49, 866-875.	21.4	75
45	Telomerase Inhibition Effectively Targets Mouse and Human AML Stem Cells and Delays Relapse following Chemotherapy. <i>Cell Stem Cell</i> , 2014, 15, 775-790.	11.1	74
46	PGBD5 promotes site-specific oncogenic mutations in human tumors. <i>Nature Genetics</i> , 2017, 49, 1005-1014.	21.4	69
47	MLL-AF9 and HOXA9-mediated acute myeloid leukemia stem cell self-renewal requires JMJD1C. <i>Journal of Clinical Investigation</i> , 2016, 126, 997-1011.	8.2	69
48	Peptidomimetic blockade of MYB in acute myeloid leukemia. <i>Nature Communications</i> , 2018, 9, 110.	12.8	68
49	IKZF2 Drives Leukemia Stem Cell Self-Renewal and Inhibits Myeloid Differentiation. <i>Cell Stem Cell</i> , 2019, 24, 153-165.e7.	11.1	66
50	Ezh2 Controls an Early Hematopoietic Program and Growth and Survival Signaling in Early T Cell Precursor Acute Lymphoblastic Leukemia. <i>Cell Reports</i> , 2016, 14, 1953-1965.	6.4	65
51	Targeting the kinase activities of ATR and ATM exhibits antitumoral activity in mouse models of MLL-rearranged AML. <i>Science Signaling</i> , 2016, 9, ra91.	3.6	63
52	MLL-rearranged leukemias: insights from gene expression profiling. <i>Seminars in Hematology</i> , 2003, 40, 268-273.	3.4	61
53	MEF2C Phosphorylation Is Required for Chemotherapy Resistance in Acute Myeloid Leukemia. <i>Cancer Discovery</i> , 2018, 8, 478-497.	9.4	59
54	Synergistic targeting of FLT3 mutations in AML via combined menin-MLL and FLT3 inhibition. <i>Blood</i> , 2020, 136, 2442-2456.	1.4	59

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55	Selective Inhibition of HDAC1 and HDAC2 as a Potential Therapeutic Option for B-ALL. <i>Clinical Cancer Research</i> , 2015, 21, 2348-2358.	7.0	57
56	Targeting chromatin complexes in fusion protein-driven malignancies. <i>Nature Reviews Cancer</i> , 2019, 19, 255-269.	28.4	55
57	Chromatin modifications as therapeutic targets in MLL-rearranged leukemia. <i>Trends in Immunology</i> , 2012, 33, 563-570.	6.8	52
58	The cell fate determinant <i>Lgl1</i> influences HSC fitness and prognosis in AML. <i>Journal of Experimental Medicine</i> , 2013, 210, 15-22.	8.5	47
59	Drugging Chromatin in Cancer: Recent Advances and Novel Approaches. <i>Molecular Cell</i> , 2015, 60, 561-570.	9.7	47
60	Mixed-Lineage Leukemia Fusions and Chromatin in Leukemia. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2017, 7, a026658.	6.2	46
61	MLL partial tandem duplication leukemia cells are sensitive to small molecule DOT1L inhibition. <i>Haematologica</i> , 2015, 100, e190-e193.	3.5	45
62	<i>miR-99</i> regulates normal and malignant hematopoietic stem cell self-renewal. <i>Journal of Experimental Medicine</i> , 2017, 214, 2453-2470.	8.5	44
63	The menin-MLL1 interaction is a molecular dependency in <i>NUP98</i> -rearranged AML. <i>Blood</i> , 2022, 139, 894-906.	1.4	42
64	The role of DOT1L in the maintenance of leukemia gene expression. <i>Current Opinion in Genetics and Development</i> , 2016, 36, 68-72.	3.3	41
65	Histone Acetyltransferase Activity of MOF Is Required for <i>MLL-AF9</i> Leukemogenesis. <i>Cancer Research</i> , 2017, 77, 1753-1762.	0.9	38
66	Myeloid Leukemia Cells With MLL partial Tandem Duplication Are Sensitive To Pharmacological Inhibition Of The H3K79 Methyltransferase DOT1L. <i>Blood</i> , 2013, 122, 1256-1256.	1.4	35
67	MLL1 and DOT1L cooperate with meningioma-1 to induce acute myeloid leukemia. <i>Journal of Clinical Investigation</i> , 2016, 126, 1438-1450.	8.2	33
68	A chromatin-independent role of Polycomb-like 1 to stabilize p53 and promote cellular quiescence. <i>Genes and Development</i> , 2015, 29, 2231-2243.	5.9	32
69	Antigen presentation safeguards the integrity of the hematopoietic stem cell pool. <i>Cell Stem Cell</i> , 2022, 29, 760-775.e10.	11.1	29
70	Menin Inhibitors in Acute Myeloid Leukemia—What Does the Future Hold?. <i>Cancer Journal (Sudbury, MA)</i> 2019, 25(10):1000-1008. TJ ETQq0 0 0 rgBT /Overlock 10 T	2.0	28
71	Resistance Mechanisms to SYK Inhibition in Acute Myeloid Leukemia. <i>Cancer Discovery</i> , 2020, 10, 214-231.	9.4	27
72	Evolutionarily Conserved Signaling Pathways: Acting in the Shadows of Acute Myelogenous Leukemia's Genetic Diversity. <i>Clinical Cancer Research</i> , 2015, 21, 240-248.	7.0	25

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73	Novel inhibitors of the histone methyltransferase DOT1L show potent antileukemic activity in patient-derived xenografts. <i>Blood</i> , 2020, 136, 1983-1988.	1.4	25
74	Inhibition Of LSD1 As a Therapeutic Strategy For The Treatment Of Acute Myeloid Leukemia. <i>Blood</i> , 2013, 122, 3964-3964.	1.4	25
75	Oncogenic Gene-Expression Programs in Leiomyosarcoma and Characterization of Conventional, Inflammatory, and Uterogenic Subtypes. <i>Molecular Cancer Research</i> , 2020, 18, 1302-1314.	3.4	24
76	Leukemia Cell of Origin Influences Apoptotic Priming and Sensitivity to LSD1 Inhibition. <i>Cancer Discovery</i> , 2020, 10, 1500-1513.	9.4	24
77	High-resolution characterization of gene function using single-cell CRISPR tiling screen. <i>Nature Communications</i> , 2021, 12, 4063.	12.8	23
78	Designed to Kill: Novel Menin-MLL Inhibitors Target MLL -Rearranged Leukemia. <i>Cancer Cell</i> , 2015, 27, 431-433.	16.8	22
79	MLL::AF9 degradation induces rapid changes in transcriptional elongation and subsequent loss of an active chromatin landscape. <i>Molecular Cell</i> , 2022, 82, 1140-1155.e11.	9.7	21
80	Structure-Guided DOT1L Probe Optimization by Label-Free Ligand Displacement. <i>ACS Chemical Biology</i> , 2015, 10, 667-674.	3.4	20
81	Inactivation of Eed impedes MLL-AF9-mediated leukemogenesis through Cdkn2a-dependent and Cdkn2a-independent mechanisms in a murine model. <i>Experimental Hematology</i> , 2015, 43, 930-935.e6.	0.4	20
82	Gastrointestinal stromal tumor enhancers support a transcription factor network predictive of clinical outcome. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E5746-E5755.	7.1	20
83	Paediatric Strategy Forum for medicinal product development of epigenetic modifiers for children. <i>European Journal of Cancer</i> , 2020, 139, 135-148.	2.8	20
84	YBX1 mediates translation of oncogenic transcripts to control cell competition in AML. <i>Leukemia</i> , 2022, 36, 426-437.	7.2	18
85	Transmembrane Inhibitor of RICTOR/mTORC2 in Hematopoietic Progenitors. <i>Stem Cell Reports</i> , 2014, 3, 832-840.	4.8	17
86	Rationale for targeting BCL6 in MLL-rearranged acute lymphoblastic leukemia. <i>Genes and Development</i> , 2019, 33, 1265-1279.	5.9	17
87	Enhancer Domains in Gastrointestinal Stromal Tumor Regulate KIT Expression and Are Targetable by BET Bromodomain Inhibition. <i>Cancer Research</i> , 2019, 79, 994-1009.	0.9	17
88	IKAROS and MENIN coordinate therapeutically actionable leukemogenic gene expression in MLL-r acute myeloid leukemia. <i>Nature Cancer</i> , 2022, 3, 595-613.	13.2	16
89	Acute myeloid leukemia driven by the CALM-AF10 fusion gene is dependent on BMI1. <i>Experimental Hematology</i> , 2019, 74, 42-51.e3.	0.4	15
90	Degradation of the BAF Complex Factor BRD9 by Heterobifunctional Ligands. <i>Angewandte Chemie</i> , 2017, 129, 5832-5837.	2.0	14

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91	HAND1 and BARX1 Act as Transcriptional and Anatomic Determinants of Malignancy in Gastrointestinal Stromal Tumor. <i>Clinical Cancer Research</i> , 2021, 27, 1706-1719.	7.0	14
92	SETD2 Haploinsufficiency Enhances Germinal Center-Associated AICDA Somatic Hypermutation to Drive B-cell Lymphomagenesis. <i>Cancer Discovery</i> , 2022, 12, 1782-1803.	9.4	14
93	Pathprinting: An integrative approach to understand the functional basis of disease. <i>Genome Medicine</i> , 2013, 5, 68.	8.2	13
94	Forward genetic screen of human transposase genomic rearrangements. <i>BMC Genomics</i> , 2016, 17, 548.	2.8	13
95	Chromatin Complexes Maintain Self-Renewal of Myeloid Progenitors in AML: Opportunities for Therapeutic Intervention. <i>Stem Cell Reports</i> , 2020, 15, 6-12.	4.8	13
96	Targeting Chromatin Complexes in Myeloid Malignancies and Beyond: From Basic Mechanisms to Clinical Innovation. <i>Cells</i> , 2020, 9, 2721.	4.1	13
97	Inhibition of MEK and ATR is effective in a B-cell acute lymphoblastic leukemia model driven by Mll-Af4 and activated Ras. <i>Blood Advances</i> , 2018, 2, 2478-2490.	5.2	12
98	MOZ and Menin-MLL Complexes Are Complementary Regulators of Chromatin Association and Transcriptional Output in Gastrointestinal Stromal Tumor. <i>Cancer Discovery</i> , 2022, 12, 1804-1823.	9.4	10
99	Genomic approaches to the pathogenesis and treatment of acute lymphoblastic leukemias. <i>Current Opinion in Hematology</i> , 2002, 9, 339-344.	2.5	9
100	A JAK/STAT-mediated inflammatory signaling cascade drives oncogenesis in AF10-rearranged AML. <i>Blood</i> , 2021, 137, 3403-3415.	1.4	8
101	Genome-Wide Identification of Prednisolone-Responsive Genes in Primary Acute Lymphoblastic Leukemia Cells. <i>Blood</i> , 2005, 106, 103-103.	1.4	8
102	It's Not What You Say But How You Say It: Targeting RNA Methylation in AML. <i>Molecular Cell</i> , 2020, 78, 996-998.	9.7	6
103	Loss of H3K36 Methyltransferase SETD2 Impairs V(D)J Recombination during Lymphoid Development. <i>IScience</i> , 2020, 23, 100941.	4.1	6
104	Location, Location, Location: Mutant NPM1c Cytoplasmic Localization Is Required to Maintain Stem Cell Genes in AML. <i>Cancer Cell</i> , 2018, 34, 355-357.	16.8	5
105	Genomic Dark Matter Sheds Light on EVI1-Driven Leukemia. <i>Cancer Cell</i> , 2014, 25, 407-408.	16.8	4
106	Histone PTM Crosstalk Stimulates Dot1 Methyltransferase Activity. <i>Trends in Biochemical Sciences</i> , 2021, 46, 522-524.	7.5	4
107	Genome-Wide RNAi Screen Identifies The Mechanistic Role For DOT1L In MLL-Rearranged Leukemia. <i>Blood</i> , 2013, 122, 598-598.	1.4	4
108	Bone Marrow Surveillance of Pediatric Cancer Survivors Identifies Clones that Predict Therapy-Related Leukemia. <i>Clinical Cancer Research</i> , 2022, 28, 1614-1627.	7.0	4

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109	Haploinsufficiency of Dnmt1 Impairs Leukemia Stem Cell Function Through Derepression of Bivalent Chromatin Domains. Blood, 2011, 118, 3459-3459.	1.4	3
110	Taspase1 orchestrates fetal liver hematopoietic stem cell and vertebrae fates through cleaving TFIIA. JCI Insight, 2021, 6, .	5.0	2
111	Patient Derived Xenograft (PDX) Models Faithfully Recapitulate The Genetic Composition Of Primary AML. Blood, 2013, 122, 1328-1328.	1.4	2
112	Identification Of Actionable Genomic Alterations In Hematologic Malignancies By a Clinical Next Generation Sequencing-Based Assay. Blood, 2013, 122, 230-230.	1.4	2
113	Regulation Of Normal and Malignant Hoxa Gene Expression Through Higher H3K79 Methylated States. Blood, 2013, 122, 2492-2492.	1.4	2
114	HoxA9 Knockdown Inhibits Proliferation and Induces Cell Death in Human MLL-Rearranged Leukemias.. Blood, 2006, 108, 734-734.	1.4	2
115	Potent Ikaros Degradation By the Cereblon E3 Ligase Modulator CC-92480 Is Effective in Combination with Menin-MLL1 Inhibition in <i>MLL1</i>-Rearranged and <i>NPM1</i>-Mutant AML. Blood, 2021, 138, 208-208.	1.4	2
116	Selective Killing of Mixed Lineage Leukemia Cells by a Potent Small-Molecule DOT1L Inhibitor. Blood, 2010, 116, 780-780.	1.4	1
117	Î²-Catenin Suppression Targets Imatinib Resistant Leukemia Stem Cells In Mice with BCR-ABL Induced Myeloproliferative Disease. Blood, 2010, 116, 93-93.	1.4	1
118	Bmi-1 Is Dispensable for the Development of Acute Myeloid Leukemia Mediated by MLL-AF9. Blood, 2010, 116, 63-63.	1.4	1
119	MLL Rearranged Infant Acute Lymphoblastic Leukemia Is Characterized by Silencing of the Putative Tumor Suppressor Gene FHIT.. Blood, 2004, 104, 525-525.	1.4	0
120	MLL-AF9 and FLT3 Cooperation during Myeloid Leukemogenesis: Development of a Model for Rapid Testing of New Therapeutics.. Blood, 2005, 106, 1605-1605.	1.4	0
121	HOXA9 Represses Bim Expression in MLL Rearranged Leukemia: Implications for Drug Therapy.. Blood, 2007, 110, 57-57.	1.4	0
122	Hoxa9+Meis1a Efficiently Transform Hematopoietic Stem Cells but Not Committed Progenitors.. Blood, 2007, 110, 3375-3375.	1.4	0
123	A Pilot Trial of Rapamycin with Glucocorticoids In Children and Adults with Relapsed ALL. Blood, 2010, 116, 3244-3244.	1.4	0
124	Demonstration of a Role for Dot1l In MLL-Rearranged Leukemia Using a Conditional Loss of Function Model. Blood, 2010, 116, 62-62.	1.4	0
125	Targeting BCL6-Mediated Drug-Resistance in High-Risk Childhood ALL. Blood, 2012, 120, 776-776.	1.4	0
126	The Interaction Between DOT1L and AF10 Is Required for H3K79 Dimethylation and MLL-AF9 Leukemia. Blood, 2012, 120, 401-401.	1.4	0

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127	Inhibition Of Telomerase Is a Novel and Effective Therapy In MLL-Rearranged Acute Myeloid Leukemia (AML). Blood, 2013, 122, 2887-2887.	1.4	0
128	Acute Myeloid Leukemia Alters The Mesenchymal Stem Cell Potential Of The HSC Niche: Evidence For Modulation By β^2 -Adrenergic Signals. Blood, 2013, 122, 342-342.	1.4	0
129	Identification Of BCL6 As a Therapeutic Target In MLL-Rearranged ALL. Blood, 2013, 122, 72-72.	1.4	0
130	Peptidomimetic Blockade of MYB in Acute Myeloid Leukemia. Blood, 2016, 128, 3945-3945.	1.4	0
131	Aberrant Phosphorylation of MEF2C Is Dispensable for Hematopoiesis, and Induces Chemotherapy Resistance and Susceptibility to MARK Kinase Inhibition Therapy in Acute Myeloid Leukemia. Blood, 2016, 128, 436-436.	1.4	0
132	Oncogenic Feedback Activation Between BCL6 and MLL Promotes Malignant Transformation in MLL-Rearranged Acute Lymphoblastic Leukemia. Blood, 2016, 128, 907-907.	1.4	0
133	Comparison of human genomics and genetic models of cancer to identify novel therapeutic targets. Cell Cycle, 2003, 2, 408-9.	2.6	0