Catrina C Fronick

List of Publications by Year in descending order

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48 5,594 papers citations

279798 276875 41
h-index g-index

50 50 all docs citations

50 times ranked 12056 citing authors

#	Article	IF	CITATIONS
1	Genome remodelling in a basal-like breast cancer metastasis and xenograft. Nature, 2010, 464, 999-1005.	27.8	1,077
2	<i>TP53</i> and Decitabine in Acute Myeloid Leukemia and Myelodysplastic Syndromes. New England Journal of Medicine, 2016, 375, 2023-2036.	27.0	663
3	Signatures of Adaptation to Obligate Biotrophy in the <i>Hyaloperonospora arabidopsidis</i> Genome. Science, 2010, 330, 1549-1551.	12.6	492
4	Genome of <i>Rhodnius prolixus</i> , an insect vector of Chagas disease, reveals unique adaptations to hematophagy and parasite infection. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 14936-14941.	7.1	329
5	Gibbon genome and the fast karyotype evolution of small apes. Nature, 2014, 513, 195-201.	27.8	320
6	TREM2 Modulation Remodels the Tumor Myeloid Landscape Enhancing Anti-PD-1 Immunotherapy. Cell, 2020, 182, 886-900.e17.	28.9	309
7	High-Dimensional Analysis Delineates Myeloid and Lymphoid Compartment Remodeling during Successful Immune-Checkpoint Cancer Therapy. Cell, 2018, 175, 1014-1030.e19.	28.9	292
8	An "off-the-shelf―fratricide-resistant CAR-T for the treatment of T cell hematologic malignancies. Leukemia, 2018, 32, 1970-1983.	7.2	282
9	Heterogeneity of meningeal B cells reveals a lymphopoietic niche at the CNS borders. Science, 2021, 373,	12.6	218
10	Rapid and Extraction-Free Detection of SARS-CoV-2 from Saliva by Colorimetric Reverse-Transcription Loop-Mediated Isothermal Amplification. Clinical Chemistry, 2021, 67, 415-424.	3.2	192
11	Optimizing Cancer Genome Sequencing and Analysis. Cell Systems, 2015, 1, 210-223.	6.2	174
12	Recurrent somatic mutations affecting B-cell receptor signaling pathway genes in follicular lymphoma. Blood, 2017, 129, 473-483.	1.4	147
13	A general approach for detecting expressed mutations in AML cells using single cell RNA-sequencing. Nature Communications, 2019, 10, 3660.	12.8	147
14	Subsets of ILC3â°'ILC1-like cells generate a diversity spectrum of innate lymphoid cells in human mucosal tissues. Nature Immunology, 2019, 20, 980-991.	14.5	141
15	TYK2 Protein-Coding Variants Protect against Rheumatoid Arthritis and Autoimmunity, with No Evidence of Major Pleiotropic Effects on Non-Autoimmune Complex Traits. PLoS ONE, 2015, 10, e0122271.	2.5	120
16	The prognostic effects of somatic mutations in ER-positive breast cancer. Nature Communications, 2018, 9, 3476.	12.8	89
17	The <i>Physarum polycephalum</i> Genome Reveals Extensive Use of Prokaryotic Two-Component and Metazoan-Type Tyrosine Kinase Signaling. Genome Biology and Evolution, 2016, 8, 109-125.	2.5	87
18	Haploinsufficiency for DNA methyltransferase 3A predisposes hematopoietic cells to myeloid malignancies. Journal of Clinical Investigation, 2017, 127, 3657-3674.	8.2	80

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19	Harnessing Expressed Single Nucleotide Variation and Single Cell RNA Sequencing To Define Immune Cell Chimerism in the Rejecting Kidney Transplant. Journal of the American Society of Nephrology: JASN, 2020, 31, 1977-1986.	6.1	71
20	Subtelomeric CTCF and cohesin binding site organization using improved subtelomere assemblies and a novel annotation pipeline. Genome Research, 2014, 24, 1039-1050.	5.5	64
21	Subclones dominate at MDS progression following allogeneic hematopoietic cell transplant. JCI Insight, 2018, 3, .	5.0	48
22	A multiple myeloma-specific capture sequencing platform discovers novel translocations and frequent, risk-associated point mutations in IGLL5. Blood Cancer Journal, 2018, 8, 35.	6.2	41
23	The clonal evolution of metastatic colorectal cancer. Science Advances, 2020, 6, eaay9691.	10.3	41
24	Sequence analysis in <i>Bos taurus</i> reveals pervasiveness of X–Y arms races in mammalian lineages. Genome Research, 2020, 30, 1716-1726.	5.5	29
25	Remethylation of <i>Dnmt3a</i> ^{â°'/â°'} hematopoietic cells is associated with partial correction of gene dysregulation and reduced myeloid skewing. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 3123-3134.	7.1	27
26	Immunosuppression and outcomes in adult patients with de novo acute myeloid leukemia with normal karyotypes. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118,	7.1	24
27	Genetic and Transcriptional Contributions to Relapse in Normal Karyotype Acute Myeloid Leukemia. Blood Cancer Discovery, 2022, 3, 32-49.	5.0	14
28	A domestic cat whole exome sequencing resource for trait discovery. Scientific Reports, 2021, 11, 7159.	3.3	13
29	Genetic Heterogeneity of Induced Pluripotent Stem Cells: Results from 24 Clones Derived from a Single C57BL/6 Mouse. PLoS ONE, 2015, 10, e0120585.	2.5	12
30	Rare Pre-Existing MDS Subclones Contribute to Secondary AML Progression. Blood, 2016, 128, 959-959.	1.4	12
31	Convergent Clonal Evolution of Signaling Gene Mutations Is a Hallmark of Myelodysplastic Syndrome Progression. Blood Cancer Discovery, 2022, 3, 330-345.	5.0	10
32	Brief Report: The Role of Rare Proteinâ€Coding Variants in Anti–Tumor Necrosis Factor Treatment Response in Rheumatoid Arthritis. Arthritis and Rheumatology, 2017, 69, 735-741.	5.6	8
33	Whole-Genome Bisulfite Sequencing of Primary AML Cells with the DNMT3A R882H Mutation Identifies Regions of Focal Hypomethylation That Are Associated with Open Chromatin. Blood, 2014, 124, 608-608.	1.4	3
34	Dynamic Changes in the Clonal Structure of MDS and AML in Response to Epigenetic Therapy. Blood, 2015, 126, 610-610.	1.4	3
35	Recurrent switch 2 domain <i>RAC2</i> mutations in intravascular large B-cell lymphoma. Blood Advances, 2022, 6, 6051-6055.	5.2	3
36	Recurrent Somatic Genomic Alterations in Follicular NHL (FL) Revealed By Exome and Custom-Capture Next Generation Sequencing. Blood, 2015, 126, 574-574.	1.4	2

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37	An Off-the-Shelfâ,,¢ Fratricide-Resistant CAR-T for the Treatment of T Cell Hematologic Malignancies. Blood, 2017, 130, 844-844.	1.4	2
38	Draft Genome Sequences of Two Polycyclic Tetramate Macrolactam Producers, Streptomyces sp. Strains JV180 and SP18CM02. Microbiology Resource Announcements, 2020, 9, .	0.6	1
39	Single-Cell Transcriptomic and Proteomic Diversity in Multiple Myeloma. Blood, 2019, 134, 5531-5531.	1.4	1
40	Specific Patterns of DNA Remethylation in the Bone Marrow Cells of Dnmt3a Deficient Mice after Induced Expression of Wild Type Human DNMT3A. Blood, 2015, 126, 433-433.	1.4	0
41	Dynamic Changes in MDS Clonal Architecture Following Allogeneic Stem Cell Transplant. Blood, 2016, 128, 5506-5506.	1.4	O
42	Evidence for Complete Mutation Clearance in Normal Karyotype AML Patients with Very Long (> 5) Tj ETQq0 0 (orgBT/Ov	erlock 10 Tf 5
43	Improving Risk Assessment of AML with a Precision Genomic Strategy to Assess Mutation Clearance. Blood, 2018, 132, 5277-5277.	1.4	O
44	Direct Detection of Expressed Mutations in AML Cells Using Single Cell RNA-Sequencing, and Its Impact on Defining Sources of Expression Heterogeneity. Blood, 2018, 132, 1314-1314.	1.4	0
45	The Molecular Basis of Long First Remissions in Normal Karyotype AML Patients. Blood, 2019, 134, 3827-3827.	1.4	О
46	Whole Genome Bisulfite Sequencing of 63 Primary AML Samples Identifies a Unique DNA Hypermethylation Signature for Mutant IDH1/2 Cases That Is Different from That of TET2 Mutant AML. Blood, 2019, 134, 3755-3755.	1.4	0
47	Identification of predicted neoantigen vaccine candidates in follicular lymphoma patients Journal of Clinical Oncology, 2020, 38, 8054-8054.	1.6	0
48	Signaling Gene Mutations Are Characterized By Diverse Patterns of Expansion and Contraction during Progression from MDS to Secondary AML. Blood, 2020, 136, 2-3.	1.4	0