## Catrina C Fronick

## List of Publications by Year in descending order

Source: https:/|exaly.com/author-pdf/3720395/publications.pdf
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| 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 â€œoff-the-shelfâ $€ \cdot f r a t r i c i d e-r e s i s t a n t ~ C A R-T ~ f o r ~ t h e ~ t r e a t m e n t ~ o f ~ T ~ c e l l ~ h e m a t o l o g i c ~ m a l i g n a n c i e s . ~$ 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 |

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.
Genetic Heterogeneity of Induced Pluripotent Stem Cells: Results from 24 Clones Derived from a
29 Single C57BL/6 Mouse. PLoS ONE, 2015, 10, e0120585.
2.5 ..... 12
Rare Pre-Existing MDS Subclones Contribute to Secondary AML Progression. Blood, 2016, 128, 959-959.1.4121.4
31 Convergent Clonal Evolution of Signaling Gene Mutations Is a Hallmark of Myelodysplastic Syndrome
Progression. Blood Cancer Discovery, 2022, 3, 330-345.5.010Brief Report: The Role of Rare Proteinâ€Coding Variants in Antiâ€"Tumor Necrosis Factor TreatmentResponse in Rheumatoid Arthritis. Arthritis and Rheumatology, 2017, 69, 735-741.
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Whole-Genome Bisulfite Sequencing of Primary AML Cells with the DNMT3A R882H Mutation Identifies
33 Regions of Focal Hypomethylation That Are Associated with Open Chromatin. Blood, 2014, 124, 608-608.
1.43

Dynamic Changes in the Clonal Structure of MDS and AML in Response to Epigenetic Therapy. Blood, 2015, 126, 610-610.
37 An Off-the-Shelfâ," $¢$ Fratricide-Resistant CAR-T for the Treatment of T Cell Hematologic Malignancies.
Blood, 2017, 130, $844-844$.
1.4

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Draft Genome Sequences of Two Polycyclic Tetramate Macrolactam Producers, Streptomyces sp. Strains JV180 and SP18CM02. Microbiology Resource Announcements, 2020, 9, .
39 Single-Cell Transcriptomic and Proteomic Diversity in Multiple Myeloma. Blood, 2019, 134,5531-5531. 1.4 .1

$40 \quad$| Specific Patterns of DNA Remethylation in the Bone Marrow Cells of Dnmt3a Deficient Mice after |
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| Induced Expression of Wild Type Human DNMT3A. Blood, 2015, 126, 433-433. |

Dynamic Changes in MDS Clonal Architecture Following Allogeneic Stem Cell Transplant. Blood, 2016,
$128,5506-5506$.

Evidence for Complete Mutation Clearance in Normal Karyotype AML Patients with Very Long (>5) Tj ETQq0 00 rgBT /Overlock 10 Tf 50

| 43 | Improving Risk Assessment of AML with a Precision Genomic Strategy to Assess Mutation Clearance. Blood, 2018, 132, 5277-5277. | 1.4 | 0 |
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| 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 | 0 |

Whole Genome Bisulfite Sequencing of 63 Primary AML Samples Identifies a Unique DNA

Identification of predicted neoantigen vaccine candidates in follicular lymphoma patients.. Journal of Clinical Oncology, 2020, 38, 8054-8054.

Signaling Gene Mutations Are Characterized By Diverse Patterns of Expansion and Contraction during Progression from MDS to Secondary AML. Blood, 2020, 136, 2-3.

