## **Daniel Rico**

List of Publications by Year in descending order

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DANIEL RICO

| #  | Article                                                                                                                                                                          | IF   | CITATIONS |
|----|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------|-----------|
| 1  | Mutational mechanisms shaping the coding and noncoding genome of germinal center derived B-cell<br>lymphomas. Leukemia, 2021, 35, 2002-2016.                                     | 3.3  | 34        |
| 2  | Resolution of R-loops by INO80 promotes DNA replication and maintains cancer cell proliferation and viability. Nature Communications, 2020, 11, 4534.                            | 5.8  | 63        |
| 3  | Intronic CNVs and gene expression variation in human populations. PLoS Genetics, 2019, 15, e1007902.                                                                             | 1.5  | 61        |
| 4  | Protection against gamma-radiation injury by protein tyrosine phosphatase 1B. Redox Biology, 2018, 17, 213-223.                                                                  | 3.9  | 9         |
| 5  | Dynamics of Transcription Regulation in Human Bone Marrow Myeloid Differentiation to Mature<br>Blood Neutrophils. Cell Reports, 2018, 24, 2784-2794.                             | 2.9  | 104       |
| 6  | Genome-wide analysis of differential transcriptional and epigenetic variability across human immune cell types. Genome Biology, 2017, 18, 18.                                    | 3.8  | 97        |
| 7  | Automatic identification of informative regions with epigenomic changes associated to hematopoiesis. Nucleic Acids Research, 2017, 45, 9244-9259.                                | 6.5  | 19        |
| 8  | Integrating epigenomic data and 3D genomic structure with a new measure of chromatin assortativity.<br>Genome Biology, 2016, 17, 152.                                            | 3.8  | 46        |
| 9  | The BLUEPRINT Data Analysis Portal. Cell Systems, 2016, 3, 491-495.e5.                                                                                                           | 2.9  | 123       |
| 10 | The International Human Epigenome Consortium: A Blueprint for Scientific Collaboration and Discovery. Cell, 2016, 167, 1145-1149.                                                | 13.5 | 404       |
| 11 | Epigenomic Co-localization and Co-evolution Reveal a Key Role for 5hmC as a Communication Hub in the Chromatin Network of ESCs. Cell Reports, 2016, 14, 1246-1257.               | 2.9  | 38        |
| 12 | Whole-genome fingerprint of the DNA methylome during human B cell differentiation. Nature<br>Genetics, 2015, 47, 746-756.                                                        | 9.4  | 278       |
| 13 | Higher gene expression variability in the more aggressive subtype of chronic lymphocytic leukemia.<br>Genome Medicine, 2015, 7, 8.                                               | 3.6  | 57        |
| 14 | The UBC-40 Urothelial Bladder Cancer cell line index: a genomic resource for functional studies. BMC<br>Genomics, 2015, 16, 403.                                                 | 1.2  | 86        |
| 15 | Transcriptome characterization by RNA sequencing identifies a major molecular and clinical subdivision in chronic lymphocytic leukemia. Genome Research, 2014, 24, 212-226.      | 2.4  | 175       |
| 16 | Transcriptional dissection of pancreatic tumors engrafted in mice. Genome Medicine, 2014, 6, 27.                                                                                 | 3.6  | 41        |
| 17 | Subfunctionalization via Adaptive Evolution Influenced by Genomic Context: The Case of Histone Chaperones ASF1a and ASF1b. Molecular Biology and Evolution, 2013, 30, 1853-1866. | 3.5  | 60        |
| 18 | Recurrent inactivation of STAG2 in bladder cancer is not associated with aneuploidy. Nature Genetics, 2013, 45, 1464-1469.                                                       | 9.4  | 224       |

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| #  | Article                                                                                                                                                                                                 | IF  | CITATIONS |
|----|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 19 | Late-replicating CNVs as a source of new genes. Biology Open, 2013, 2, 1402-1411.                                                                                                                       | 0.6 | 9         |
| 20 | Epigenomic analysis detects widespread gene-body DNA hypomethylation in chronic lymphocytic leukemia. Nature Genetics, 2012, 44, 1236-1242.                                                             | 9.4 | 525       |
| 21 | Assessment of copy number variation using the Illumina Infinium 1M SNP-array: a comparison of methodological approaches in the Spanish Bladder Cancer/EPICURO study. Human Mutation, 2011, 32, 240-248. | 1.1 | 57        |
| 22 | Mosaic Uniparental Disomies and Aneuploidies as Large Structural Variants of the Human Genome.<br>American Journal of Human Genetics, 2010, 87, 129-138.                                                | 2.6 | 111       |
| 23 | waviCGH: a web application for the analysis and visualization of genomic copy number alterations.<br>Nucleic Acids Research, 2010, 38, W182-W187.                                                       | 6.5 | 18        |
| 24 | Substrate Fate in Activated Macrophages: A Comparison between Innate, Classic, and Alternative<br>Activation. Journal of Immunology, 2010, 185, 605-614.                                                | 0.4 | 820       |
| 25 | Tetrahymena Metallothioneins Fall into Two Discrete Subfamilies. PLoS ONE, 2007, 2, e291.                                                                                                               | 1.1 | 76        |
| 26 | Identification of conserved domains in the promoter regions of nitric oxide synthase 2: implications for the species-specific transcription and evolutionary differences. BMC Genomics, 2007, 8, 271.   | 1.2 | 17        |