Jaime I Davila

List of Publications by Year in descending order

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IAIME I DAVILA

#	Article	IF	CITATIONS
1	Inherited Mutations in 17 Breast Cancer Susceptibility Genes Among a Large Triple-Negative Breast Cancer Cohort Unselected for Family History of Breast Cancer. Journal of Clinical Oncology, 2015, 33, 304-311.	1.6	521
2	MAP-RSeq: Mayo Analysis Pipeline for RNA sequencing. BMC Bioinformatics, 2014, 15, 224.	2.6	284
3	Double-strand break repair processes drive evolution of the mitochondrial genome in Arabidopsis. BMC Biology, 2011, 9, 64.	3.8	209
4	Diversity of the Arabidopsis Mitochondrial Genome Occurs via Nuclear-Controlled Recombination Activity. Genetics, 2009, 183, 1261-1268.	2.9	161
5	MutS HOMOLOG1 Is a Nucleoid Protein That Alters Mitochondrial and Plastid Properties and Plant Response to High Light Â. Plant Cell, 2011, 23, 3428-3441.	6.6	125
6	ILâ€33 facilitates oncogeneâ€induced cholangiocarcinoma in mice by an interleukinâ€6â€sensitive mechanism. Hepatology, 2015, 61, 1627-1642.	7.3	115
7	Integrated mate-pair and RNA sequencing identifies novel, targetable gene fusions in peripheral T-cell lymphoma. Blood, 2016, 128, 1234-1245.	1.4	105
8	RNA Toxicity and Missplicing in the Common Eye Disease Fuchs Endothelial Corneal Dystrophy. Journal of Biological Chemistry, 2015, 290, 5979-5990.	3.4	104
9	Gastroblastoma harbors a recurrent somatic MALAT1–GLI1 fusion gene. Modern Pathology, 2017, 30, 1443-1452.	5.5	93
10	Fast and Practical Algorithms for Planted (I, d) Motif Search. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2007, 4, 544-552.	3.0	92
11	Extensive Rearrangement of the Arabidopsis Mitochondrial Genome Elicits Cellular Conditions for Thermotolerance. Plant Physiology, 2010, 152, 1960-1970.	4.8	77
12	Recurrent STAT3-JAK2 fusions in indolent T-cell lymphoproliferative disorder of the gastrointestinal tract. Blood, 2018, 131, 2262-2266.	1.4	77
13	Contraction of T cell richness in lung cancer brain metastases. Scientific Reports, 2018, 8, 2171.	3.3	74
14	Development and characterization of human-induced pluripotent stem cell-derived cholangiocytes. Laboratory Investigation, 2015, 95, 684-696.	3.7	66
15	Spindle cell rhabdomyosarcoma of bone with <i><scp>FUS</scp>–<scp>TFCP</scp>2</i> fusion: confirmation of a very recently described rhabdomyosarcoma subtype. Histopathology, 2018, 73, 514-520.	2.9	63
16	Cytoplasmic Male Sterility-Associated Chimeric Open Reading Frames Identified by Mitochondrial Genome Sequencing of Four Cajanus Genotypes. DNA Research, 2013, 20, 485-495.	3.4	58
17	ST3GAL1 is a target of the SOX2-GL11 transcriptional complex and promotes melanoma metastasis through AXL. Nature Communications, 2020, 11, 5865.	12.8	54
18	Novel <i>TRAF1â€ALK</i> fusion identified by deep RNA sequencing of anaplastic large cell lymphoma. Genes Chromosomes and Cancer, 2013, 52, 1097-1102.	2.8	51

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19	Proteomic Detection of Immunoglobulin Light Chain Variable Region Peptides from Amyloidosis Patient Biopsies. Journal of Proteome Research, 2015, 14, 1957-1967.	3.7	50
20	<i>TP53</i> mutations, tetraploidy and homologous recombination repair defects in early stage high-grade serous ovarian cancer. Nucleic Acids Research, 2015, 43, 6945-6958.	14.5	46
21	Uterine inflammatory myofibroblastic tumors in pregnant women with and without involvement of the placenta: a study of 6 cases with identification of a novel TIMP3-RET fusion. Human Pathology, 2020, 97, 29-39.	2.0	43
22	Composite hemangioendothelioma with neuroendocrine marker expression: an aggressive variant. Modern Pathology, 2017, 30, 1589-1602.	5.5	38
23	qPMS7: A Fast Algorithm for Finding (â"", d)-Motifs in DNA and Protein Sequences. PLoS ONE, 2012, 7, e41425.	2.5	38
24	Bioinformatics and DNA-extraction strategies to reliably detect genetic variants from FFPE breast tissue samples. BMC Genomics, 2019, 20, 689.	2.8	37
25	Murine Leukemia Virus Uses NXF1 for Nuclear Export of Spliced and Unspliced Viral Transcripts. Journal of Virology, 2014, 88, 4069-4082.	3.4	36
26	Development and Verification of an RNA Sequencing (RNA-Seq) Assay for the Detection of Gene Fusions in Tumors. Journal of Molecular Diagnostics, 2018, 20, 495-511.	2.8	36
27	RVboost: RNA-seq variants prioritization using a boosting method. Bioinformatics, 2014, 30, 3414-3416.	4.1	34
28	Impact of RNA degradation on fusion detection by RNA-seq. BMC Genomics, 2016, 17, 814.	2.8	34
29	Gene expression differences between matched pairs of ovarian cancer patient tumors and patient-derived xenografts. Scientific Reports, 2019, 9, 6314.	3.3	33
30	RNA sequencing identifies a novel <i>USP9Xâ€USP6</i> promoter swap gene fusion in a primary aneurysmal bone cyst. Genes Chromosomes and Cancer, 2019, 58, 589-594.	2.8	27
31	Space and Time Efficient Algorithms for Planted Motif Search. Lecture Notes in Computer Science, 2006, , 822-829.	1.3	25
32	Desmoplastic Infantile Ganglioglioma: A MAPK Pathway-Driven and Microglia/Macrophage-Rich Neuroepithelial Tumor. Journal of Neuropathology and Experimental Neurology, 2019, 78, 1011-1021.	1.7	21
33	RNA-Seq Reveals Differences in Expressed Tumor Mutation Burden in Colorectal and Endometrial Cancers with and without Defective DNA-Mismatch Repair. Journal of Molecular Diagnostics, 2021, 23, 555-564.	2.8	16
34	Molecular and Immunohistochemical Analysis of Mucinous Cystic Neoplasm of the Liver. American Journal of Clinical Pathology, 2020, 154, 837-847.	0.7	14
35	Xanthogranulomatous epithelial tumor: report of 6 cases of a novel, potentially deceptive lesion with a predilection for young women. Modern Pathology, 2020, 33, 1889-1895.	5.5	13
36	Polymorphous Low-Grade Neuroepithelial Tumor of the Young (PLNTY): Molecular Profiling Confirms Frequent MAPK Pathway Activation. Journal of Neuropathology and Experimental Neurology, 2021, 80, 821-829.	1.7	13

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37	Leiomyoma with KAT6B-KANSL1 fusion: case report of a rapidly enlarging uterine mass in a postmenopausal woman. Diagnostic Pathology, 2019, 14, 32.	2.0	11
38	Identification and Development of a Lung Adenocarcinoma PDX Model With STRN-ALK Fusion. Clinical Lung Cancer, 2019, 20, e142-e147.	2.6	11
39	Frequent POLE-driven hypermutation in ovarian endometrioid cancer revealed by mutational signatures in RNA sequencing. BMC Medical Genomics, 2021, 14, 165.	1.5	10
40	Transcriptomic and Proteomic Analysis of Steatohepatitic Hepatocellular Carcinoma Reveals Novel Distinct Biologic Features. American Journal of Clinical Pathology, 2021, 155, 87-96.	0.7	9
41	Comprehensive Genomic Profiling of a Rare Thyroid Follicular Dendritic Cell Sarcoma. Rare Tumors, 2017, 9, 50-53.	0.6	8
42	Malignant Peritoneal Mesothelioma Arising in Young Adults With Long-standing Indwelling Intra-abdominal Shunt Catheters. American Journal of Surgical Pathology, 2021, 45, 255-262.	3.7	6
43	Molecular Genetic Landscape of Sclerosing Pneumocytomas. American Journal of Clinical Pathology, 2021, 155, 397-404.	0.7	5
44	Determining mutational burden and signature using RNA-seq from tumor-only samples. BMC Medical Genomics, 2021, 14, 65.	1.5	5
45	Semi-Supervised Topological Analysis for Elucidating Hidden Structures in High-Dimensional Transcriptome Datasets. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2021, 18, 1620-1631.	3.0	4
46	Gene Expression Profiling Identifies Distinct Signatures for Dysplastic and Proliferative Chronic Myelomonocytic Leukemia. Blood, 2016, 128, 110-110.	1.4	4
47	Novel Phenotypic and Genetic Analysis of T-Cell Prolymphocytic Leukemia (T-PLL). Blood, 2014, 124, 1682-1682.	1.4	3
48	Stage-Specific Non-Coding RNA Expression Patterns during In Vitro Human B Cell Differentiation into Antibody Secreting Plasma Cells. Non-coding RNA, 2022, 8, 15.	2.6	3
49	RANDOMIZED SORTING ON THE POPS NETWORK. International Journal of Foundations of Computer Science, 2005, 16, 105-116.	1.1	2
50	Multiple isodicentric Y chromosomes in myeloid malignancies: a unique cytogenetic entity and potential therapeutic target. Leukemia and Lymphoma, 2019, 60, 821-824.	1.3	2
51	Network-directed cis-mediator analysis of normal prostate tissue expression profiles reveals downstream regulatory associations of prostate cancer susceptibility loci. Oncotarget, 2017, 8, 85896-85908.	1.8	2
52	Packet routing and selection on the POPS network. Journal of Parallel and Distributed Computing, 2005, 65, 927-933.	4.1	1
53	Extending Pattern Branching to Handle Challenging Instances. , 2006, , .		1

54 Sorting- and FFT-Based Techniques in the Discovery of Biopatterns. , 2007, , 93-115.

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#	Article	IF	CITATIONS
55	Fast Algorithms for Selecting Specific siRNA in Complete mRNA Data. Lecture Notes in Computer Science, 2007, , 302-309.	1.3	1
56	CLL Mesenchymal Stromal Cells Have Decreased Replicative Potential and Senescent Phenotype: Clinical and Biologic Implications. Blood, 2014, 124, 3282-3282.	1.4	1
57	Developing a FHIR-based Framework for Phenome Wide Association Studies: A Case Study with A Pan-Cancer Cohort. AMIA Summits on Translational Science Proceedings, 2020, 2020, 750-759.	0.4	1
58	Data-driven Sublanguage Analysis for Cancer Genomics Knowledge Modeling: Applications in Mining Oncological Genetics Information from Patients' Genetic Reports. AMIA Summits on Translational Science Proceedings, 2020, 2020, 720-729.	0.4	1
59	Challenges of using RNA-seq in the clinical setting. , 2017, , .		0
60	Randomized Packet Routing, Selection, and Sorting on the POPS Network. Chapman & Hall/CRC Computer and Information Science Series, 2007, , 13-1-13-14.	0.4	0
61	A tool to predict post-transcriptional instability related to the dysregulation of the SETD2 histone methyltransferase in renal cell carcinoma (RCC) Journal of Clinical Oncology, 2014, 32, 11072-11072.	1.6	Ο
62	Abstract 1894: Extensive genomic profiling of a rare extranodal-follicular dendritic cell sarcoma: Implications for future individualized therapy. , 2014, , .		0
63	Abstract 3462: Comparative RNA-Seq analysis of MIS signaling: Potential relevance as therapeutic strategy in ovarian cancer treatment. , 2014, , .		Ο
64	Abstract 5585: Integrated DNA/RNA sequencing for discovery and orthogonal validation of expressed fusion genes in peripheral T-cell lymphomas. , 2014, , .		0
65	RNA-Seq Based Immunoglobulin Repertoire Analysis of Normal Plasma Cells Generated in an in Vitro B Cell Differentiation System. Blood, 2019, 134, 1051-1051.	1.4	0