Pedro Alexandre Favoretto Galante

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Monoallelic deleterious <scp> <i>MUTYH </i> </scp> germline variants as a driver for tumorigenesis. Journal of Pathology, 2022, 256, 214-222.	2.1	12
2	Identification of NID1 as a novel candidate susceptibility gene for familial non-medullary thyroid carcinoma using whole-exome sequencing. Endocrine Connections, 2022, 11, .	0.8	3
3	Whole-genome sequencing of 1,171 elderly admixed individuals from Brazil. Nature Communications, 2022, 13, 1004.	5.8	35
4	The RNA-Binding Protein Musashi1 Regulates a Network of Cell Cycle Genes in Group 4 Medulloblastoma. Cells, 2022, 11, 56.	1.8	3
5	Severe Acute Respiratory Syndrome Coronavirus 2 Variants of Concern: A Perspective for Emerging More Transmissible and Vaccine-Resistant Strains. Viruses, 2022, 14, 827.	1.5	14
6	Association between tumor mutational burden (TMB) and mutational profile and its effect on overall survival: A post hoc analysis of patients with TMB-high and TMB-low metastatic cancer treated with immune checkpoint inhibitors (ICI) Journal of Clinical Oncology, 2022, 40, 2632-2632.	0.8	0
7	sideRETRO: a pipeline for identifying somatic and polymorphic insertions of processed pseudogenes or retrocopies. Bioinformatics, 2021, 37, 419-421.	1.8	7
8	Synergism of Proneurogenic miRNAs Provides a More Effective Strategy to Target Glioma Stem Cells. Cancers, 2021, 13, 289.	1.7	7
9	Deciphering the Role of Intestinal Crypt Cell Populations in Resistance to Chemotherapy. Cancer Research, 2021, 81, 2730-2744.	0.4	4
10	Reboot: a straightforward approach to identify genes and splicing isoforms associated with cancer patient prognosis. NAR Cancer, 2021, 3, zcab024.	1.6	8
11	Evidence of Cooperation between Hippo Pathway and RAS Mutation in Thyroid Carcinomas. Cancers, 2021, 13, 2306.	1.7	4
12	Molecular Signature Expands the Landscape of Driver Negative Thyroid Cancers. Cancers, 2021, 13, 5184.	1.7	0
13	Chemogenetic modulation of sensory neurons reveals their regulating role in melanoma progression. Acta Neuropathologica Communications, 2021, 9, 183.	2.4	21
14	The RNA-binding protein SERBP1 functions as a novel oncogenic factor in glioblastoma by bridging cancer metabolism and epigenetic regulation. Genome Biology, 2020, 21, 195.	3.8	55
15	Depletion of Ric-8B leads to reduced mTORC2 activity. PLoS Genetics, 2020, 16, e1008255.	1.5	3
16	Gene expression changes associated with trajectories of psychopathology in a longitudinal cohort of children and adolescents. Translational Psychiatry, 2020, 10, 99.	2.4	3
17	Genomic Biomarkers and Underlying Mechanism of Benefit from BCG Immunotherapy in Non-Muscle Invasive Bladder Cancer. Bladder Cancer, 2020, 6, 171-186.	0.2	4
18	Proneural and mesenchymal glioma stem cells display major differences in splicing and lncRNA profiles. Npj Genomic Medicine, 2020, 5, 2.	1.7	29

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19	Comprehensive germline mutation analysis and clinical profile in a large cohort of Brazilian xeroderma pigmentosum patients. Journal of the European Academy of Dermatology and Venereology, 2020, 34, 2392-2401.	1.3	17
20	Depletion of Ric-8B leads to reduced mTORC2 activity. , 2020, 16, e1008255.		0
21	Depletion of Ric-8B leads to reduced mTORC2 activity. , 2020, 16, e1008255.		0
22	Depletion of Ric-8B leads to reduced mTORC2 activity. , 2020, 16, e1008255.		0
23	Depletion of Ric-8B leads to reduced mTORC2 activity. , 2020, 16, e1008255.		0
24	Uncovering the mouse olfactory long non-coding transcriptome with a novel machine-learning model. DNA Research, 2019, 26, 365-378.	1,5	8
25	Retroposed copies of RET gene: a somatically acquired event in medullary thyroid carcinoma. BMC Medical Genomics, 2019, 12, 104.	0.7	10
26	The Effects of Neoadjuvant Chemoradiation in Locally Advanced Rectal Cancer—The Impact in Intratumoral Heterogeneity. Frontiers in Oncology, 2019, 9, 974.	1.3	20
27	Antagonism between the RNA-binding protein Musashi1 and miR-137 and its potential impact on neurogenesis and glioblastoma development. Rna, 2019, 25, 768-782.	1.6	25
28	Transposon insertion profiling by sequencing (TIPseq) for mapping LINE-1 insertions in the human genome. Mobile DNA, 2019, 10, 8.	1.3	22
29	Tumor mutational burden (TMB) and BCG responsiveness in high-risk non-muscle invasive bladder cancer (NMIBC) Journal of Clinical Oncology, 2019, 37, 442-442.	0.8	2
30	Identification and Validation of Potential Differential miRNA Regulation via Alternative Polyadenylation. Methods in Molecular Biology, 2018, 1733, 87-92.	0.4	1
31	Patient-derived conditionally reprogrammed cells maintain intra-tumor genetic heterogeneity. Scientific Reports, 2018, 8, 4097.	1.6	34
32	<i>MRPL53</i> , a New Candidate Gene for Orofacial Clefting, Identified Using an eQTL Approach. Journal of Dental Research, 2018, 97, 33-40.	2.5	8
33	TSC2 rare germline variants in non-tuberous sclerosis patients with neuroendocrine neoplasias. Endocrine-Related Cancer, 2018, 25, L1-L5.	1.6	6
34	miRNA-based signatures in cerebrospinal fluid as potential diagnostic tools for early stage Parkinson's disease. Oncotarget, 2018, 9, 17455-17465.	0.8	94
35	Histopathological Characterization and Whole Exome Sequencing of Ectopic Thyroid: Fetal Architecture in a Functional Ectopic Gland from Adult Patient. International Journal of Endocrinology, 2018, 2018, 1-10.	0.6	7
36	Tumor mutational burden (TMB), intratumoral genetic heterogeneity (ITGH) and BCG responsiveness in high-risk non-muscle invasive bladder cancer (NMIBC) Journal of Clinical Oncology, 2018, 36, e16516-e16516.	0.8	1

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37	Intratumoral Genetic Heterogeneity in Rectal Cancer. Annals of Surgery, 2017, 265, e4-e6.	2.1	56
38	A straightforward assay to evaluate DNA integrity and optimize next-generation sequencing for clinical diagnosis in oncology. Experimental and Molecular Pathology, 2017, 103, 294-299.	0.9	18
39	A genetic cluster of patients with variant xeroderma pigmentosum with two different founder mutations. British Journal of Dermatology, 2017, 176, 1270-1278.	1.4	23
40	High IL-1R8 expression in breast tumors promotes tumor growth and contributes to impaired antitumor immunity. Oncotarget, 2017, 8, 49470-49483.	0.8	24
41	Unveiling the Impact of the Genomic Architecture on the Evolution of Vertebrate microRNAs. Frontiers in Genetics, 2017, 8, 34.	1.1	14
42	MiRIAD update: using alternative polyadenylation, protein interaction network analysis and additional species to enhance exploration of the role of intragenic miRNAs and their host genes. Database: the Journal of Biological Databases and Curation, 2017, 2017, .	1.4	10
43	Measuring plasma levels of three microRNAs can improve the accuracy for identification of malignant breast lesions in women with BI-RADS 4 mammography. Oncotarget, 2017, 8, 83940-83948.	0.8	7
44	Functional genomics analyses of RNA-binding proteins reveal the splicing regulator SNRPB as an oncogenic candidate in glioblastoma. Genome Biology, 2016, 17, 125.	3.8	83
45	Host gene constraints and genomic context impact the expression and evolution of human microRNAs. Nature Communications, 2016, 7, 11438.	5.8	47
46	MicroRNA Expression Signature Is Altered in the Cardiac Remodeling Induced by High Fat Diets. Journal of Cellular Physiology, 2016, 231, 1771-1783.	2.0	27
47	Signaling transcript profile of the asexual intraerythrocytic development cycle of Plasmodium falciparum induced by melatonin and cAMP. Genes and Cancer, 2016, 7, 323-339.	0.6	16
48	A Mouse Model of Targeted Musashi1 Expression in Whole Intestinal Epithelium Suggests Regulatory Roles in Cell Cycle and Stemness. Stem Cells, 2015, 33, 3621-3634.	1.4	25
49	Comprehensive cancer-gene panels can be used to estimate mutational load and predict clinical benefit to PD-1 blockade in clinical practice. Oncotarget, 2015, 6, 34221-34227.	0.8	198
50	Alternative Polyadenylation Allows Differential Negative Feedback of Human miRNA miR-579 on Its Host Gene ZFR. PLoS ONE, 2015, 10, e0121507.	1.1	24
51	The use of personalized biomarkers and liquid biopsies to monitor treatment response and disease recurrence in locally advanced rectal cancer after neoadjuvant chemoradiation. Oncotarget, 2015, 6, 38360-38371.	0.8	52
52	Identification of agonists for a group of human odorant receptors. Frontiers in Pharmacology, 2015, 6, 35.	1.6	37
53	Comprehensive evaluation of the effectiveness of gene expression signatures to predict complete response to neoadjuvant chemoradiotherapy and guide surgical intervention in rectal cancer. Cancer Genetics, 2015, 208, 319-326.	0.2	45
54	A Genome-Wide Landscape of Retrocopies in Primate Genomes. Genome Biology and Evolution, 2015, 7, 2265-2275.	1.1	46

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55	ICRmax: An optimized approach to detect tumor-specific interchromosomal rearrangements for clinical application. Genomics, 2015, 105, 265-272.	1.3	4
56	Recurrent somatic mutation in DROSHA induces microRNA profile changes in Wilms tumour. Nature Communications, 2014, 5, 4039.	5.8	159
57	Overexpression of miR-21-5p as a predictive marker for complete tumor regression to neoadjuvant chemoradiotherapy in rectal cancer patients. BMC Medical Genomics, 2014, 7, 68.	0.7	58
58	miRIAD—integrating microRNA inter- and intragenic data. Database: the Journal of Biological Databases and Curation, 2014, 2014, .	1.4	85
59	Mutational analysis of genes coding for cell surface proteins in colorectal cancer cell lines reveal novel altered pathways, druggable mutations and mutated epitopes for targeted therapy. Oncotarget, 2014, 5, 9199-9213.	0.8	31
60	Setting Up an Intronic miRNA Database. Methods in Molecular Biology, 2013, 936, 69-76.	0.4	6
61	The Human Cell Surfaceome of Breast Tumors. BioMed Research International, 2013, 2013, 1-11.	0.9	6
62	Gene Copy-Number Polymorphism Caused by Retrotransposition in Humans. PLoS Genetics, 2013, 9, e1003242.	1.5	88
63	RCPedia: a database of retrocopied genes. Bioinformatics, 2013, 29, 1235-1237.	1.8	32
64	SPLOOCE. RNA Biology, 2012, 9, 1339-1343.	1.5	7
65	SurfaceomeDB: a cancer-orientated database for genes encoding cell surface proteins. Cancer Immunity, 2012, 12, 15.	3.2	7
66	HPV-58 Molecular Variants Exhibit Different Transcriptional Activity. Intervirology, 2011, 54, 146-150.	1.2	8
67	Temporal blastemal cell gene expression analysis in the kidney reveals new Wnt and related signaling pathway genes to be essential for Wilms' tumor onset. Cell Death and Disease, 2011, 2, e224-e224.	2.7	23
68	PVALB, a New Hürthle Adenoma Diagnostic Marker Identified through Gene Expression. Journal of Clinical Endocrinology and Metabolism, 2011, 96, E151-E160.	1.8	22
69	Analysis of allelic differential expression in the human genome using allele-specific serial analysis of gene expression tags. Genome, 2011, 54, 120-127.	0.9	5
70	Alternative splicing and genetic diversity: silencers are more frequently modified by SNVs associated with alternative exon/intron borders. Nucleic Acids Research, 2011, 39, 4942-4948.	6.5	11
71	Distinct patterns of somatic alterations in a lymphoblastoid and a tumor genome derived from the same individual. Nucleic Acids Research, 2011, 39, 6056-6068.	6.5	19
72	Common Promoter Elements in Odorant and Vomeronasal Receptor Genes. PLoS ONE, 2011, 6, e29065.	1.1	32

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73	A potential role for intragenic miRNAs on their hosts' interactome. BMC Genomics, 2010, 11, 533.	1.2	142
74	Alternative splicing enriched cDNA libraries identify breast cancer-associated transcripts. BMC Genomics, 2010, 11, S4.	1.2	10
75	DSGeo: Software tools for cross-platform analysis of gene expression data in GEO. Journal of Biomedical Informatics, 2010, 43, 709-715.	2.5	13
76	Systematic detection of putative tumor suppressor genes through the combined use of exome and transcriptome sequencing. Genome Biology, 2010, 11, R114.	13.9	35
77	Application of Bioinformatics in Cancer Research. , 2010, , 211-233.		7
78	Transcriptome-guided characterization of genomic rearrangements in a breast cancer cell line. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 1886-1891.	3.3	104
79	A comprehensive in silico expression analysis of RNA binding proteins in normal and tumor tissue; identification of potential players in tumor formation. RNA Biology, 2009, 6, 426-433.	1.5	51
80	Bioinformatics construction of the human cell surfaceome. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 16752-16757.	3.3	119
81	A score system for quality evaluation of RNA sequence tags: an improvement for gene expression profiling. BMC Bioinformatics, 2009, 10, 170.	1.2	2
82	Evaluation of a large-scale biomedical data annotation initiative. BMC Bioinformatics, 2009, 10, S10.	1.2	16
83	Towards large-scale sample annotation in gene expression repositories. BMC Bioinformatics, 2009, 10, S9.	1.2	7
84	Identification of FAM46D as a novel cancer/testis antigen using EST data and serological analysis. Genomics, 2009, 94, 153-160.	1.3	23
85	Different Evolutionary Strategies for the Origin of Caspase-1 Inhibitors. Journal of Molecular Evolution, 2008, 66, 591-597.	0.8	14
86	Differential gene expression analysis of iodide-treated rat thyroid follicular cell line PCCl3. Genomics, 2008, 91, 356-366.	1.3	29
87	Genome-Wide Detection of Serpentine Receptor-Like Proteins in Malaria Parasites. PLoS ONE, 2008, 3, e1889.	1.1	43
88	Automatic correspondence of tags and genes (ACTG): a tool for the analysis of SAGE, MPSS and SBS data. Bioinformatics, 2007, 23, 903-905.	1.8	10
89	Sense-antisense pairs in mammals: functional and evolutionary considerations. Genome Biology, 2007, 8, R40.	13.9	55
90	Alternative splicing: a bioinformatics perspective. Molecular BioSystems, 2007, 3, 473.	2.9	13

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91	Identification of potential regulatory motifs in odorant receptor genes by analysis of promoter sequences. Genome Research, 2006, 16, 1091-1098.	2.4	59
92	The impact of SNPs on the interpretation of SAGE and MPSS experimental data. Nucleic Acids Research, 2004, 32, 6104-6110.	6.5	34
93	Detection and evaluation of intron retention events in the human transcriptome. Rna, 2004, 10, 757-765.	1.6	193
94	Splicing factors are differentially expressed in tumors. Genetics and Molecular Research, 2004, 3, 512-20.	0.3	21
95	A novel human G protein-coupled receptor is over-expressed in prostate cancer. Genetics and Molecular Research, 2004, 3, 521-31.	0.3	6
96	ORESTES are enriched in rare exon usage variants affecting the encoded proteins. Comptes Rendus - Biologies, 2003, 326, 979-985.	0.1	13
97	The generation and utilization of a cancer-oriented representation of the human transcriptome by using expressed sequence tags. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 13418-13423.	3.3	105