## Tathiane M Malta

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	The Immune Landscape of Cancer. Immunity, 2018, 48, 812-830.e14.	6.6	3,706
2	TCGAbiolinks: an R/Bioconductor package for integrative analysis of TCGA data. Nucleic Acids Research, 2016, 44, e71-e71.	6.5	2,519
3	Cell-of-Origin Patterns Dominate the Molecular Classification of 10,000 Tumors from 33 Types of Cancer. Cell, 2018, 173, 291-304.e6.	13.5	1,718
4	Molecular Profiling Reveals Biologically Discrete Subsets and Pathways of Progression in Diffuse Glioma. Cell, 2016, 164, 550-563.	13.5	1,695
5	Machine Learning Identifies Stemness Features Associated with Oncogenic Dedifferentiation. Cell, 2018, 173, 338-354.e15.	13.5	1,417
6	Longitudinal molecular trajectories of diffuse glioma in adults. Nature, 2019, 576, 112-120.	13.7	320
7	Glioma CpG island methylator phenotype (G-CIMP): biological and clinical implications. Neuro-Oncology, 2018, 20, 608-620.	0.6	194
8	DNA methylation profiling to predict recurrence risk in meningioma: development and validation of a nomogram to optimize clinical management. Neuro-Oncology, 2019, 21, 901-910.	0.6	184
9	A global metagenomic map of urban microbiomes and antimicrobial resistance. Cell, 2021, 184, 3376-3393.e17.	13.5	164
10	A Distinct DNA Methylation Shift in a Subset of Glioma CpG Island Methylator Phenotypes during Tumor Recurrence. Cell Reports, 2018, 23, 637-651.	2.9	137
11	Patient-derived organoids and orthotopic xenografts of primary and recurrent gliomas represent relevant patient avatars for precision oncology. Acta Neuropathologica, 2020, 140, 919-949.	3.9	72
12	A serum-based DNA methylation assay provides accurate detection of glioma. Neuro-Oncology, 2021, 23, 1494-1508.	0.6	53
13	Mutated CEACAMs Disrupt Transforming Growth Factor Beta Signaling and Alter the Intestinal Microbiome to Promote Colorectal Carcinogenesis. Gastroenterology, 2020, 158, 238-252.	0.6	46
14	Cultured Human Adipose Tissue Pericytes and Mesenchymal Stromal Cells Display a Very Similar Gene Expression Profile. Stem Cells and Development, 2015, 24, 2822-2840.	1.1	44
15	HTLV-1/2 seroprevalence and coinfection rate in Brazilian first-time blood donors: an 11-year follow-up. Revista Do Instituto De Medicina Tropical De Sao Paulo, 2012, 54, 123-130.	0.5	35
16	Metabolic reprogramming associated with aggressiveness occurs in the G-CIMP-high molecular subtypes of IDH1mut lower grade gliomas. Neuro-Oncology, 2020, 22, 480-492.	0.6	31
17	Predicting master transcription factors from pan-cancer expression data. Science Advances, 2021, 7, eabf6123.	4.7	30
18	Transcriptomic comparisons between cultured human adipose tissue-derived pericytes and mesenchymal stromal cells. Genomics Data, 2016, 7, 20-25.	1.3	25

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19	Identification of subsets of IDH-mutant glioblastomas with distinct epigenetic and copy number alterations and stratified clinical risks. Neuro-Oncology Advances, 2019, 1, vdz015.	0.4	22
20	Generation of induced pluripotent stem cells from large domestic animals. Stem Cell Research and Therapy, 2020, 11, 247.	2.4	21
21	Molecular landscape of IDH-mutant primary astrocytoma Grade IV/glioblastomas. Modern Pathology, 2021, 34, 1245-1260.	2.9	21
22	Genes Related to Antiviral Activity, Cell Migration, and Lysis Are Differentially Expressed in CD4+T Cells in Human T Cell Leukemia Virus Type 1-Associated Myelopathy/Tropical Spastic Paraparesis Patients. AIDS Research and Human Retroviruses, 2014, 30, 610-622.	0.5	20
23	The gene expression profile of non-cultured, highly purified human adipose tissue pericytes: Transcriptomic evidence that pericytes are stem cells in human adipose tissue. Experimental Cell Research, 2016, 349, 239-254.	1.2	19
24	Targeting the E3 Ubiquitin Ligase PJA1 Enhances Tumor-Suppressing TGFÎ <sup>2</sup> Signaling. Cancer Research, 2020, 80, 1819-1832.	0.4	17
25	TCGAbiolinksGUI: A graphical user interface to analyze cancer molecular and clinical data. F1000Research, 0, 7, 439.	0.8	14
26	Short Communication Forced expression of OCT4 influences the expression of pluripotent genes in human mesenchymal stem cells and fibroblasts. Genetics and Molecular Research, 2013, 12, 1054-1060.	0.3	13
27	Leukotrienes Are Upregulated and Associated with Human T-Lymphotropic Virus Type 1 (HTLV-1)-Associated Neuroinflammatory Disease. PLoS ONE, 2012, 7, e51873.	1.1	10
28	Detection of tumor-specific DNA methylation markers in the blood of patients with pituitary neuroendocrine tumors. Neuro-Oncology, 2022, 24, 1126-1139.	0.6	9
29	Molecular landscape of <scp><i>IDH</i></scp> â€wild type, <scp>p<i>TERT</i></scp> â€wild type adult glioblastomas. Brain Pathology, 2022, 32, .	2.1	9
30	Functional and bioinformatics analyses reveal conservation of <i>cis</i> â€regulatory elements between sciaridae and drosophilidae. Genesis, 2008, 46, 43-51.	0.8	6
31	T cell receptor signaling pathway is overexpressed in CD4+ T cells from HAM/TSP individuals. Brazilian Journal of Infectious Diseases, 2015, 19, 578-584.	0.3	6
32	DNA methylation-based signatures classify sporadic pituitary tumors according to clinicopathological features. Neuro-Oncology, 2021, 23, 1292-1303.	0.6	6
33	Altered Expression of Degranulation-Related Genes in CD8+T Cells in Human T Lymphotropic Virus Type I Infection. AIDS Research and Human Retroviruses, 2013, 29, 826-836.	0.5	4
34	Genes related to antiviral activity are differentially expressed in CD4+ T cell in HAM/TSP patients. Retrovirology, 2014, 11, .	0.9	1
35	GENE-61. METHYLATION-BASED LIQUID BIOPSY OF MENINGIOMA PRIMARY AND RECURRENT SAMPLES. Neuro-Oncology, 2019, 21, vi111-vi111.	0.6	1

Abstract LB-373: Comprehensive analysis of cancer stemness. , 2018, , .

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37	Distribution of QPY and RAH haplotypes of granzyme B gene in distinct Brazilian populations. Revista Da Sociedade Brasileira De Medicina Tropical, 2012, 45, 496-499.	0.4	0
38	Novel polymorphisms in the promoter region of the perforin gene among distinct <scp>B</scp> razilian populations and their functional impact. International Journal of Immunogenetics, 2014, 41, 198-205.	0.8	0
39	EPIG-14EPIGENOMIC (DNA METHYLATION AND EXPRESSION) SIGNATURES DEFINE SUBSETS OF BOTH IDHmut AND IDHwt GLIOMA WITH DISTINCT CLINICAL OUTCOMES. Neuro-Oncology, 2015, 17, v89.2-v89.	0.6	0
40	EPIG-16MOST DIFFERENTIAL DNA METHYLATION CHANGES OCCUR AT CANDIDATE ENHANCER ELEMENTS FOR RECURRENT LOWER GRADE GLIOMA AND GLIOBLASTOMA. Neuro-Oncology, 2015, 17, v89.4-v90.	0.6	0
41	GENO-06A PAN-GLIOMA CHARACTERIZATION OF GENOMIC, EPIGENOMIC AND TRANSCRIPTOMIC ACTIVITIES REVEALS NOVEL RELATIONSHIPS BETWEEN HISTOLOGICAL SUBTYPES AND MOLECULAR SIGNATURES. Neuro-Oncology, 2015, 17, v92.2-v92.	0.6	0
42	GENT-33. EPIGENETIC ALTERATIONS AT INTERGENIC REGIONS ASSOCIATED WITH PROGRESSION IN AÂSUBSET OF IDH MUTANT GLIOMAS. Neuro-Oncology, 2016, 18, vi81-vi81.	0.6	0
43	GENT-34. EPIGENOMIC STEMNESS SIGNATURE ASSOCIATED WITH GLIOMA MOLECULAR SUBTYPES. Neuro-Oncology, 2016, 18, vi81-vi81.	0.6	0
44	IMMU-60. PD-L1 EXPRESSION ASSOCIATED WITH GLIOMA MOLECULAR SUBTYPES: CLINICAL IMPLICATIONS. Neuro-Oncology, 2017, 19, vi126-vi126.	0.6	0
45	GENE-52. EPIGENOMIC GLIOMA SUBTYPE EVALUATION ACROSS 31 TUMOR TYPES. Neuro-Oncology, 2017, 19, vi103-vi104.	0.6	0
46	GENE-09. FUNCTIONAL GENOMIC ELEMENTS DEFINED BY DNA METHYLATION CAN DISTINGUISH MENINGIOMA SUBGROUPS. Neuro-Oncology, 2018, 20, vi104-vi104.	0.6	0
47	GENE-07. GENOMIC ENHANCER METHYLATION IS ASSOCIATED WITH BIOLOGICAL AND CLINICAL FEATURES IN PITUITARY TUMORS. Neuro-Oncology, 2018, 20, vi104-vi104.	0.6	0
48	CBMT-42. LOSS OF PROMOTER METHYLATION IN GLYCOLYTIC GENES IS ASSOCIATED WITH AGGRESSIVENESS IN IDH1-MUTANT LOWER GRADE GLIOMAS. Neuro-Oncology, 2018, 20, vi41-vi42.	0.6	0
49	Combined epigenetic signature and gene copy number variations in IDH-mutant glioblastomas showed varied risk stratification. Neuro-Oncology, 2019, 21, iv2-iv2.	0.6	0
50	Detection of Glioma and Prognostic Subtypes by Noninvasive Circulating Cell-Free DNA Methylation Markers. Neurosurgery, 2019, 66, 310-630.	0.6	0
51	Candidate Regulatory Elements in Coding and Noncoding Regions are Associated With Invasive Behavior in Pituitary Tumors. Neurosurgery, 2019, 66, .	0.6	0
52	GENE-24. DNA METHYLATION SIGNATURES DETECTED IN A SERUM-BASED LIQUID BIOPSY DISTINGUISH FUNCTIONAL AND INVASIVENESS FEATURES IN PITUITARY ADENOMAS. Neuro-Oncology, 2019, 21, vi102-vi102.	0.6	0
53	GENE-58. ACTIVE TRANSCRIPTION START SITES OF MENINGIOMA SAMPLES ASSOCIATED WITH RISK OF RECURRENCE. Neuro-Oncology, 2019, 21, vi110-vi110.	0.6	0

54 Abstract LB-004: Molecular hallmarks of cancer: Stemness. , 2017, , .

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55	Distinct Epigenetic Shift in a Subset of Glioma CpG Island Methylator Phenotype (G-CIMP) During Tumor Recurrence. SSRN Electronic Journal, 0, , .	0.4	0
56	IDO1 expression in glioma molecular subtypes Journal of Clinical Oncology, 2018, 36, e14029-e14029.	0.8	0
57	OR24-5 Aberrant DNA Methylation in Regulatory Genomic Elements Are Associated with Invasive Behavior of Pituitary Macroadenomas: An Integrative Analysis of Epigenome-Wide Studies. Journal of the Endocrine Society, 2019, 3, .	0.1	0
58	Abstract 1939: Non-invasive prediction of glioma tumor stemness using multimodal MRI. , 2019, , .		0
59	Abstract 1640: Deep learning classification of neuro-oncology medical documents. , 2019, , .		0
60	Abstract LB-234: Pre-surgical identification of diagnostic, prognostic and predictive DNA methylation-based markers in serum (liquid biopsy) of patients harboring gliomas. , 2019, , .		0
61	Abstract A11: Serum cell-free DNA methylome-based signatures distinguish pituitary tumor from other neoplasias and by clinicopathologic features. , 2020, , .		0
62	Abstract A10: Glioma cell-free DNA methylation marker for diagnosis and monitoring. , 2020, , .		0
63	Abstract 841: Meningioma subgroups associated with functional genomic elements defined by DNA methylation. , 2019, , .		0