

Tessy Thomas Maliekal

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/10958480/publications.pdf>

Version: 2024-02-01

11
papers

140
citations

1163117

8
h-index

1281871

11
g-index

12
all docs

12
docs citations

12
times ranked

149
citing authors

#	ARTICLE	IF	CITATIONS
1	Histone Chaperone Nucleophosmin Regulates Transcription of Key Genes Involved in Oral Tumorigenesis. <i>Molecular and Cellular Biology</i> , 2022, 42, MCB0066920.	2.3	9
2	Reporters of Cancer Stem Cells as a Tool for Drug Discovery. <i>Frontiers in Oncology</i> , 2021, 11, 669250.	2.8	19
3	Markers and Reporters to Reveal the Hierarchy in Heterogeneous Cancer Stem Cells. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 668851.	3.7	16
4	SSTP1, a Host Defense Peptide, Exploits the Immunomodulatory IL6 Pathway to Induce Apoptosis in Cancer Cells. <i>Frontiers in Immunology</i> , 2021, 12, 740620.	4.8	2
5	̢-Tubulin Isotype, TUBB4B, Regulates The Maintenance of Cancer Stem Cells. <i>Frontiers in Oncology</i> , 2021, 11, 788024.	2.8	5
6	B1CTcu5: A frog-derived brevinin-1 peptide with anti-tuberculosis activity. <i>Peptides</i> , 2020, 132, 170373.	2.4	12
7	A novel reporter construct for screening small molecule inhibitors that specifically target self-renewing cancer cells. <i>Experimental Cell Research</i> , 2019, 383, 111551.	2.6	7
8	Single cell biology beyond the era of antibodies: relevance, challenges, and promises in biomedical research. <i>Cellular and Molecular Life Sciences</i> , 2017, 74, 1177-1189.	5.4	10
9	CD66 and CD49f expressing cells are associated with distinct neoplastic phenotypes and progression in human cervical cancer. <i>European Journal of Cancer</i> , 2016, 60, 166-178.	2.8	16
10	TM1-IR680 peptide for assessment of surgical margin and lymph node metastasis in murine orthotopic model of oral cancer. <i>Scientific Reports</i> , 2016, 6, 36726.	3.3	11
11	Notch Signaling in CD66+ Cells Drives the Progression of Human Cervical Cancers. <i>Cancer Research</i> , 2011, 71, 4888-4897.	0.9	33