

Preethi Srinivasan

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

19
papers

2,644
citations

10
h-index

22
g-index

22
ext. papers

3,602
ext. citations

13.8
avg, IF

3.54
L-index

#	Paper	IF	Citations
19	Mutational landscape of metastatic cancer revealed from prospective clinical sequencing of 10,000 patients. <i>Nature Medicine</i> , 2017 , 23, 703-713	50.5	1638
18	Analysis of the Prevalence of Microsatellite Instability in Prostate Cancer and Response to Immune Checkpoint Blockade. <i>JAMA Oncology</i> , 2019 , 5, 471-478	13.4	257
17	Microsatellite Instability Is Associated With the Presence of Lynch Syndrome Pan-Cancer. <i>Journal of Clinical Oncology</i> , 2019 , 37, 286-295	2.2	203
16	Tumour lineage shapes BRCA-mediated phenotypes. <i>Nature</i> , 2019 , 571, 576-579	50.4	170
15	Clinical relevance of KRAS-mutated subclones detected with picodroplet digital PCR in advanced colorectal cancer treated with anti-EGFR therapy. <i>Clinical Cancer Research</i> , 2015 , 21, 1087-97	12.9	122
14	Multiplex picoliter-droplet digital PCR for quantitative assessment of DNA integrity in clinical samples. <i>Clinical Chemistry</i> , 2013 , 59, 815-23	5.5	78
13	Clinical Utility of Prospective Molecular Characterization in Advanced Endometrial Cancer. <i>Clinical Cancer Research</i> , 2018 , 24, 5939-5947	12.9	60
12	Genomic Landscape of Uterine Sarcomas Defined Through Prospective Clinical Sequencing. <i>Clinical Cancer Research</i> , 2020 , 26, 3881-3888	12.9	16
11	A Comprehensive Comparison of Early-Onset and Average-Onset Colorectal Cancers. <i>Journal of the National Cancer Institute</i> , 2021 ,	9.7	12
10	Enhanced specificity of clinical high-sensitivity tumor mutation profiling in cell-free DNA via paired normal sequencing using MSK-ACCESS. <i>Nature Communications</i> , 2021 , 12, 3770	17.4	10
9	Tumor fraction-guided cell-free DNA profiling in metastatic solid tumor patients. <i>Genome Medicine</i> , 2021 , 13, 96	14.4	8
8	Ampullary cancer: Evaluation of somatic and germline genetic alterations and association with clinical outcomes. <i>Cancer</i> , 2019 , 125, 1441-1448	6.4	8
7	Propagation of uncertainty in Bayesian diagnostic test interpretation. <i>Southern Medical Journal</i> , 2012 , 105, 452-9	0.6	7
6	Protein-altering germline mutations implicate novel genes related to lung cancer development. <i>Nature Communications</i> , 2020 , 11, 2220	17.4	6
5	The context-specific role of germline pathogenicity in tumorigenesis. <i>Nature Genetics</i> , 2021 , 53, 1577-1585	35.3	6
4	Prevalence and Characterization of Biallelic and Monoallelic and Variant Carriers From a Pan-Cancer Patient Population. <i>JCO Precision Oncology</i> , 2021 , 5,	3.6	3
3	Enhanced specificity of high sensitivity somatic variant profiling in cell-free DNA via paired normal sequencing: design, validation, and clinical experience of the MSK-ACCESS liquid biopsy assay		2

- 2 Understanding inherited risk in unselected newly diagnosed patients with endometrial cancer. *JCO Precision Oncology*, **2019**, 3, 3.6 2
- 1 Germline Contributions to Clonal Hematopoiesis in Solid Cancer Patients. *Blood*, **2020**, 136, 30-31 2.2