

# Vidyalakshmi Sethunath

## List of Publications by Year in descending order

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Version: 2024-02-01

22  
papers

650  
citations

840776

11  
h-index

996975

15  
g-index

23  
all docs

23  
docs citations

23  
times ranked

1251  
citing authors

#	ARTICLE	IF	CITATIONS
1	Integrative clinical and molecular characterization of translocation renal cell carcinoma. Cell Reports, 2022, 38, 110190.	6.4	40
2	A genome-scale CRISPR screen reveals PRMT1 as a critical regulator of androgen receptor signaling in prostate cancer. Cell Reports, 2022, 38, 110417.	6.4	17
3	Abstract PD3-09:HER2 L755Smutation is acquired upon resistance to lapatinib and neratinib and confers cross-resistance to tucatinib and trastuzumab in HER2-positive breast cancer cell models. , 2021, , .		2
4	Abstract PS5-29: Insights into the molecular underpinnings of the mevalonate pathway-YAP/TAZ-driven anti-HER2 therapy resistance in HER2+ breast cancer (BC). , 2021, , .		0
5	Activation of the IFN Signaling Pathway is Associated with Resistance to CDK4/6 Inhibitors and Immune Checkpoint Activation in ER-Positive Breast Cancer. Clinical Cancer Research, 2021, 27, 4870-4882.	7.0	49
6	Neratinib plus trastuzumab is superior to pertuzumab plus trastuzumab in HER2-positive breast cancer xenograft models. Npj Breast Cancer, 2021, 7, 63.	5.2	4
7	Evaluation of the Predictive Role of Tumor Immune Infiltrate in Patients with HER2-Positive Breast Cancer Treated with Neoadjuvant Anti-HER2 Therapy without Chemotherapy. Clinical Cancer Research, 2020, 26, 738-745.	7.0	31
8	Towards personalized treatment for early stage HER2-positive breast cancer. Nature Reviews Clinical Oncology, 2020, 17, 233-250.	27.6	166
9	Abstract 1911: HER2 L755S mutation is associated with acquired resistance to lapatinib and neratinib, and confers cross-resistance to tucatinib in HER2-positive breast cancer models. Cancer Research, 2020, 80, 1911-1911.	0.9	8
10	Abstract GS2-01: High levels of interferon-response gene signatures are associated withde novoand acquired resistance to CDK4/6 inhibitors in ER+ breast cancer. , 2020, , .		2
11	Abstract PD2-02: Activation of the EGFR/RAS/p42,44 MAPK axis as a convergent mechanism of resistance to CDK4/6 inhibitors in ER+ breast cancer. , 2020, , .		0
12	Abstract P3-06-07: ADGRF1 overexpression inhibits tumor growthin vivoby inducing cell cycle arrest in HER2+ breast cancer. , 2020, , .		0
13	Abstract P6-04-02: Integrative cistromic/transcriptomic profiling identifies a high FOXA1/ER-activated pro-metastatic secretome in endocrine-resistant breast cancer. , 2020, , .		0
14	Targeting the Mevalonate Pathway to Overcome Acquired Anti-HER2 Treatment Resistance in Breast Cancer. Molecular Cancer Research, 2019, 17, 2318-2330.	3.4	41
15	FOXA1 upregulation promotes enhancer and transcriptional reprogramming in endocrine-resistant breast cancer. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 26823-26834.	7.1	103
16	Abstract 4827: The therapeutic superiority of neratinib in combination with trastuzumab compared to pertuzumab plus trastuzumab in HER2-positivein vivobreast cancer models. , 2019, , .		0
17	Abstract 4757: Targeting the mevalonate pathway in HER2+breast cancer to overcome resistance and enhance anti-HER2 therapy efficacy. , 2019, , .		0
18	GPCRs profiling and identification of GPR110 as a potential new target in HER2+ breast cancer. Breast Cancer Research and Treatment, 2018, 170, 279-292.	2.5	22

#	ARTICLE	IF	CITATIONS
19	HER2 Reactivation through Acquisition of the HER2 L755S Mutation as a Mechanism of Acquired Resistance to HER2-targeted Therapy in HER2+ Breast Cancer. <i>Clinical Cancer Research</i> , 2017, 23, 5123-5134.	7.0	85
20	Drug-sensitive FGFR3 mutations in lung adenocarcinoma. <i>Annals of Oncology</i> , 2017, 28, 597-603.	1.2	36
21	Notch pathway activation is essential for maintenance of stem-like cells in early tongue cancer. <i>Oncotarget</i> , 2016, 7, 50437-50449.	1.8	40
22	Abstract 4811: Pro-oncogenic role of NOTCH1 in early tongue squamous cell carcinoma. <i>Cancer Research</i> , 2015, 75, 4811-4811.	0.9	1