

# Manfred JÄ¼cker

## List of Publications by Year in descending order

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Version: 2024-02-01

42  
papers

1,678  
citations

304602

22  
h-index

302012

39  
g-index

42  
all docs

42  
docs citations

42  
times ranked

2788  
citing authors

#	ARTICLE	IF	CITATIONS
1	Distinct functions of AKT isoforms in breast cancer: a comprehensive review. <i>Cell Communication and Signaling</i> , 2019, 17, 154.	2.7	192
2	PI3K/AKT/mTOR signaling as a molecular target in head and neck cancer. <i>Biochemical Pharmacology</i> , 2020, 172, 113729.	2.0	174
3	The Role of mTOR Signaling as a Therapeutic Target in Cancer. <i>International Journal of Molecular Sciences</i> , 2021, 22, 1743.	1.8	128
4	Combined targeting of AKT and mTOR synergistically inhibits proliferation of hepatocellular carcinoma cells. <i>Molecular Cancer</i> , 2012, 11, 85.	7.9	97
5	COSMC knockdown mediated aberrant O-glycosylation promotes oncogenic properties in pancreatic cancer. <i>Molecular Cancer</i> , 2015, 14, 109.	7.9	89
6	Expression of Hedgehog Pathway Mediator <i>GLI1</i> Represents a Negative Prognostic Marker in Human Acute Myeloid Leukemia and Its Inhibition Exerts Antileukemic Effects. <i>Clinical Cancer Research</i> , 2015, 21, 2388-2398.	3.2	88
7	Characterization of circulating breast cancer cells with tumorigenic and metastatic capacity. <i>EMBO Molecular Medicine</i> , 2020, 12, e11908.	3.3	77
8	Distinct functional roles of Akt isoforms for proliferation, survival, migration and EGF-mediated signalling in lung cancer derived disseminated tumor cells. <i>Cellular Signalling</i> , 2011, 23, 1952-1960.	1.7	76
9	Combined targeting of AKT and mTOR using MK2206 and RAD001 is synergistic in the treatment of cholangiocarcinoma. <i>International Journal of Cancer</i> , 2013, 133, 2065-2076.	2.3	71
10	The Functional Role of Extracellular Matrix Proteins in Cancer. <i>Cancers</i> , 2022, 14, 238.	1.7	65
11	Downregulation of AKT3 Increases Migration and Metastasis in Triple Negative Breast Cancer Cells by Upregulating S100A4. <i>PLoS ONE</i> , 2016, 11, e0146370.	1.1	61
12	Dual Inhibition of PI3K-AKT-mTOR- and RAF-MEK-ERK-signaling is synergistic in cholangiocarcinoma and reverses acquired resistance to MEK-inhibitors. <i>Investigational New Drugs</i> , 2014, 32, 1144-1154.	1.2	50
13	Suppression of Early Hematogenous Dissemination of Human Breast Cancer Cells to Bone Marrow by Retinoic Acid-Induced 2. <i>Cancer Discovery</i> , 2015, 5, 506-519.	7.7	45
14	Dual Targeting of Akt and mTORC1 Impairs Repair of DNA Double-Strand Breaks and Increases Radiation Sensitivity of Human Tumor Cells. <i>PLoS ONE</i> , 2016, 11, e0154745.	1.1	42
15	AKT3 regulates ErbB2, ErbB3 and estrogen receptor $\alpha$ expression and contributes to endocrine therapy resistance of ErbB2+ breast tumor cells from Balb-neuT mice. <i>Cellular Signalling</i> , 2014, 26, 1021-1029.	1.7	37
16	PTEN mediates the cross talk between breast and glial cells in brain metastases leading to rapid disease progression. <i>Oncotarget</i> , 2017, 8, 6155-6168.	0.8	35
17	Vertical Targeting of AKT and mTOR as Well as Dual Targeting of AKT and MEK Signaling Is Synergistic in Hepatocellular Carcinoma. <i>Journal of Cancer</i> , 2015, 6, 1195-1205.	1.2	34
18	Targeted PI3K/AKT-hyperactivation induces cell death in chronic lymphocytic leukemia. <i>Nature Communications</i> , 2021, 12, 3526.	5.8	34

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19	Akt isoform specific effects in ovarian cancer progression. <i>Oncotarget</i> , 2016, 7, 74820-74833.	0.8	33
20	Leukemia-associated mutations in SHIP1 inhibit its enzymatic activity, interaction with the GM-CSF receptor and Grb2, and its ability to inactivate PI3K/AKT signaling. <i>Cellular Signalling</i> , 2012, 24, 2095-2101.	1.7	31
21	Combined inhibition of GLI and FLT3 signaling leads to effective anti-leukemic effects in human acute myeloid leukemia. <i>Oncotarget</i> , 2017, 8, 29187-29201.	0.8	28
22	High Sensitivity of Circulating Tumor Cells Derived from a Colorectal Cancer Patient for Dual Inhibition with AKT and mTOR Inhibitors. <i>Cells</i> , 2020, 9, 2129.	1.8	26
23	The tumor suppressor SHIP1 colocalizes in nucleolar cavities with p53 and components of PML nuclear bodies. <i>Nucleus</i> , 2015, 6, 154-164.	0.6	24
24	The inositol 5-phosphatase SHIP1 is a nucleo-cytoplasmic shuttling protein and enzymatically active in cell nuclei. <i>Cellular Signalling</i> , 2012, 24, 621-628.	1.7	18
25	Circulating tumor cells as a promising target for individualized drug susceptibility tests in cancer therapy. <i>Biochemical Pharmacology</i> , 2021, 188, 114589.	2.0	18
26	Combined Targeting of AKT and mTOR Inhibits Proliferation of Human NF1-Associated Malignant Peripheral Nerve Sheath Tumour Cells In Vitro but not in a Xenograft Mouse Model In Vivo. <i>International Journal of Molecular Sciences</i> , 2020, 21, 1548.	1.8	15
27	Knockdown of AKT3 Activates HER2 and DDR Kinases in Bone-Seeking Breast Cancer Cells, Promotes Metastasis In Vivo and Attenuates the TGF $\beta$ <sup>2</sup> /CTGF Axis. <i>Cells</i> , 2021, 10, 430.	1.8	14
28	An increase in the expression and total activity of endogenous p60c-Src in several factor-independent mutants of a human GM-CSF-dependent leukemia cell line (TF-1). <i>Oncogene</i> , 2003, 22, 7170-7180.	2.6	12
29	AKT in Bone Metastasis of Solid Tumors: A Comprehensive Review. <i>Cancers</i> , 2021, 13, 2287.	1.7	10
30	Combined Targeting of AKT and mTOR Synergistically Inhibits Formation of Primary Colorectal Carcinoma Tumouroids <i>in Vitro</i> : A 3D Tumour Model for Pre-therapeutic Drug Screening. <i>Anticancer Research</i> , 2021, 41, 2257-2275.	0.5	8
31	Nuclear accumulation of SHIP1 mutants derived from AML patients leads to increased proliferation of leukemic cells. <i>Cellular Signalling</i> , 2018, 49, 87-94.	1.7	7
32	Truncated O-GalNAc glycans impact on fundamental signaling pathways in pancreatic cancer. <i>Glycobiology</i> , 2021, , .	1.3	6
33	Combined Targeting of AKT and mTOR Inhibits Tumor Formation of EpCAM+ and CD90+ Human Hepatocellular Carcinoma Cells in an Orthotopic Mouse Model. <i>Cancers</i> , 2022, 14, 1882.	1.7	6
34	AKT1 and PTEN show the highest affinities among phosphoinositide binding proteins for the second messengers PtdIns(3,4,5)P <sub>3</sub> and PtdIns(3,4)P <sub>2</sub> . <i>Biochemical and Biophysical Research Communications</i> , 2021, 568, 110-115.	1.0	5
35	Analysis of the FLVR motif of SHIP1 and its importance for the protein stability of SH2 containing signaling proteins. <i>Cellular Signalling</i> , 2019, 63, 109380.	1.7	4
36	Differential regulation of extracellular matrix proteins in three recurrent liver metastases of a single patient with colorectal cancer. <i>Clinical and Experimental Metastasis</i> , 2020, 37, 649-656.	1.7	4

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37	Investigation of the function of the PI3-Kinase / AKT signaling pathway for leukemogenesis and therapy of acute childhood lymphoblastic leukemia (ALL). Cellular Signalling, 2022, 93, 110301.	1.7	4
38	Discontinuing MEK inhibitors in tumor cells with an acquired resistance increases migration and invasion. Cellular Signalling, 2015, 27, 2191-2200.	1.7	3
39	Characterization of the substrate specificity of the inositol 5-phosphatase SHIP1. Biochemical and Biophysical Research Communications, 2020, 524, 366-370.	1.0	3
40	Ectopic Expression of Hematopoietic SHIP1 in Human Colorectal Cancer. Biomedicines, 2020, 8, 215.	1.4	2
41	JAK2-V617F is a negative regulation factor of SHIP1 protein and thus influences the AKT signaling pathway in patients with Myeloproliferative neoplasm (MPN). International Journal of Biochemistry and Cell Biology, 2022, 149, 106229.	1.2	2
42	"Alcohol and nicotine"--Concept and evaluation of an interdisciplinary elective course with OSPE in preclinical medical education. GMS Zeitschrift für Medizinische Ausbildung, 2014, 31, Doc9.	1.2	0