

Anthony C Faber

List of Publications by Year in descending order

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Version: 2024-02-01

61
papers

5,860
citations

126907

33
h-index

149698

56
g-index

63
all docs

63
docs citations

63
times ranked

10898
citing authors

#	ARTICLE	IF	CITATIONS
1	MYCN upregulates the transsulfuration pathway to suppress the ferroptotic vulnerability in MYCN-amplified neuroblastoma. <i>Cell Stress</i> , 2022, 6, 21-29.	3.2	5
2	<i>MYCN</i> -Amplified Neuroblastoma Is Addicted to Iron and Vulnerable to Inhibition of the System Xc-/Glutathione Axis. <i>Cancer Research</i> , 2021, 81, 1896-1908.	0.9	73
3	Targeting transcription of MCL-1 sensitizes HER2-amplified breast cancers to HER2 inhibitors. <i>Cell Death and Disease</i> , 2021, 12, 179.	6.3	11
4	Catastrophic ATP loss underlies a metabolic combination therapy tailored for <i>MYCN</i> -amplified neuroblastoma. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	7.1	10
5	Unmasking BCL-2 Addiction in Synovial Sarcoma by Overcoming Low NOXA. <i>Cancers</i> , 2021, 13, 2310.	3.7	6
6	Venetoclax-based Rational Combinations are Effective in Models of <i>MYCN</i> -amplified Neuroblastoma. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 1400-1411.	4.1	10
7	Pharmaceutical Interference of the EWS-FLI1-driven Transcriptome By Cotargeting H3K27ac and RNA Polymerase Activity in Ewing Sarcoma. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 1868-1879.	4.1	8
8	Androgen-deprivation induced senescence in prostate cancer cells is permissive for the development of castration-resistance but susceptible to senolytic therapy. <i>Biochemical Pharmacology</i> , 2021, 193, 114765.	4.4	20
9	Investigating New Mechanisms of Acquired Resistance to Targeted Therapies: If You Hit Them Harder, Do They Get Up Differently?. <i>Cancer Research</i> , 2020, 80, 25-26.	0.9	4
10	Clearance of therapy-induced senescent tumor cells by the senolytic ABT-263 via interference with BCL-XL/BAX interaction. <i>Molecular Oncology</i> , 2020, 14, 2504-2519.	4.6	90
11	Targetable immune checkpoint molecules may be significantly differentially expressed in minority ethnicities.. <i>Journal of Clinical Oncology</i> , 2020, 38, 3576-3576.	1.6	4
12	Evaluation of combined BCL-2/MCL-1 inhibition as a therapeutic approach for synovial sarcoma.. <i>Journal of Clinical Oncology</i> , 2020, 38, e23561-e23561.	1.6	2
13	Epithelial-to-mesenchymal transition and drug resistance: transitioning away from death. <i>Journal of Thoracic Disease</i> , 2019, 11, E82-E85.	1.4	21
14	Paragangliomas in Carney-Stratakis Syndrome. <i>Hormone and Metabolic Research</i> , 2019, 51, 437-442.	1.5	5
15	Prognostic Factors of Malignant Pheochromocytoma and Paraganglioma: A Combined SEER and TCGA Databases Review. <i>Hormone and Metabolic Research</i> , 2019, 51, 451-457.	1.5	20
16	The Ewing Family of Tumors Relies on BCL-2 and BCL-XL to Escape PARP Inhibitor Toxicity. <i>Clinical Cancer Research</i> , 2019, 25, 1664-1675.	7.0	26
17	NOTCH1 Represses MCL-1 Levels in GSI-resistant T-ALL, Making them Susceptible to ABT-263. <i>Clinical Cancer Research</i> , 2019, 25, 312-324.	7.0	11
18	Tumor mutation burden and PD-L1 expression in SDH/FH mutated solid tumors.. <i>Journal of Clinical Oncology</i> , 2019, 37, 1524-1524.	1.6	1

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19	Coamplification of <i>miR-4728</i> protects <i>HER2</i> -amplified breast cancers from targeted therapy. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E2594-E2603.	7.1	23
20	Therapeutically targeting tumor microenvironment-mediated drug resistance in estrogen receptor-positive breast cancer. Journal of Experimental Medicine, 2018, 215, 895-910.	8.5	63
21	Distinct dependencies on receptor tyrosine kinases in the regulation of MAPK signaling between BRAF V600E and non-V600E mutant lung cancers. Oncogene, 2018, 37, 1775-1787.	5.9	28
22	Gastrointestinal Stromal Tumors: The GIST of Precision Medicine. Trends in Cancer, 2018, 4, 74-91.	7.4	71
23	Classification of gastrointestinal stromal tumor syndromes. Endocrine-Related Cancer, 2018, 25, R49-R58.	3.1	31
24	Epithelial-to-Mesenchymal Transition Antagonizes Response to Targeted Therapies in Lung Cancer by Suppressing BIM. Clinical Cancer Research, 2018, 24, 197-208.	7.0	74
25	Venetoclax Is Effective in Small-Cell Lung Cancers with High BCL-2 Expression. Clinical Cancer Research, 2018, 24, 360-369.	7.0	96
26	One gene to rule them all and in the darkness bind them. Molecular and Cellular Oncology, 2018, 5, e1465881.	0.7	0
27	Targeted inhibition of histone H3K27 demethylation is effective in high-risk neuroblastoma. Science Translational Medicine, 2018, 10, .	12.4	70
28	OVOL2 in metastasis prevention in NPC. Theranostics, 2018, 8, 2242-2244.	10.0	3
29	Increased Synthesis of MCL-1 Protein Underlies Initial Survival of <i>EGFR</i> -Mutant Lung Cancer to EGFR Inhibitors and Provides a Novel Drug Target. Clinical Cancer Research, 2018, 24, 5658-5672.	7.0	38
30	BCL-2 inhibition is a promising therapeutic strategy for small cell lung cancer. Oncoscience, 2018, 5, 218-219.	2.2	19
31	Sensitivity and Resistance to BH3 Mimetics in Cancer Therapy. Resistance To Targeted Anti-cancer Therapeutics, 2018, , 147-180.	0.1	0
32	An expanding role for osimertinib for the treatment of ErbB family driven NSCLC. Translational Cancer Research, 2018, 7, S787-S791.	1.0	1
33	Epithelial-to-Mesenchymal Transition Defines Feedback Activation of Receptor Tyrosine Kinase Signaling Induced by MEK Inhibition in <i>KRAS</i> -Mutant Lung Cancer. Cancer Discovery, 2016, 6, 754-769.	9.4	132
34	Tumor cells can follow distinct evolutionary paths to become resistant to epidermal growth factor receptor inhibition. Nature Medicine, 2016, 22, 262-269.	30.7	768
35	Exploitation of the Apoptosis-Primed State of MYCN-Amplified Neuroblastoma to Develop a Potent and Specific Targeted Therapy Combination. Cancer Cell, 2016, 29, 159-172.	16.8	104
36	Assessment of ABT-263 activity across a cancer cell line collection leads to a potent combination therapy for small-cell lung cancer. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E1288-96.	7.1	110

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37	Measurement of PIP3 Levels Reveals an Unexpected Role for p110 ^β in Early Adaptive Responses to p110 ^β -Specific Inhibitors in Luminal Breast Cancer. <i>Cancer Cell</i> , 2015, 27, 97-108.	16.8	165
38	The BCL2 Family: Key Mediators of the Apoptotic Response to Targeted Anticancer Therapeutics. <i>Cancer Discovery</i> , 2015, 5, 475-487.	9.4	501
39	Lack of Association between the BIM Deletion Polymorphism and the Risk of Lung Cancer with and without EGFR Mutations. <i>Journal of Thoracic Oncology</i> , 2015, 10, 59-66.	1.1	13
40	Inhibition of mutant EGFR in lung cancer cells triggers SOX2-FOXO6-dependent survival pathways. <i>ELife</i> , 2015, 4, .	6.0	37
41	Not just a RAS ping at flaws: Finding vulnerabilities to develop novel therapies for treating KRAS mutant cancers. <i>Cancer Science</i> , 2014, 105, 499-505.	3.9	19
42	mTOR Inhibition Specifically Sensitizes Colorectal Cancers with KRAS or BRAF Mutations to BCL-2/BCL-XL Inhibition by Suppressing MCL-1. <i>Cancer Discovery</i> , 2014, 4, 42-52.	9.4	116
43	Patient-derived models of acquired resistance can identify effective drug combinations for cancer. <i>Science</i> , 2014, 346, 1480-1486.	12.6	635
44	Combination PI3K/MEK inhibition promotes tumor apoptosis and regression in PIK3CA wild-type, KRAS mutant colorectal cancer. <i>Cancer Letters</i> , 2014, 347, 204-211.	7.2	36
45	Failure to Induce Apoptosis via BCL-2 Family Proteins Underlies Lack of Efficacy of Combined MEK and PI3K Inhibitors for KRAS-Mutant Lung Cancers. <i>Cancer Research</i> , 2014, 74, 3146-3156.	0.9	69
46	Synthetic Lethal Interaction of Combined BCL-XL and MEK Inhibition Promotes Tumor Regressions in KRAS Mutant Cancer Models. <i>Cancer Cell</i> , 2013, 23, 121-128.	16.8	343
47	Concomitant BRAF and PI3K/mTOR Blockade Is Required for Effective Treatment of BRAFV600E Colorectal Cancer. <i>Clinical Cancer Research</i> , 2013, 19, 2688-2698.	7.0	76
48	TORC1 Suppression Predicts Responsiveness to RAF and MEK Inhibition in BRAF-Mutant Melanoma. <i>Science Translational Medicine</i> , 2013, 5, 196ra98.	12.4	124
49	PI3K regulates MEK/ERK signaling in breast cancer via the Rac-GEF, P-Rex1. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 21124-21129.	7.1	175
50	Abstract A062: Sensitizing low BIM expressing breast cancers to targeted therapies. , 2013, , .		0
51	Activation of PI3K Signaling in Merkel Cell Carcinoma. <i>Clinical Cancer Research</i> , 2012, 18, 1227-1236.	7.0	97
52	Apoptosis In Targeted Therapy Responses. <i>Advances in Pharmacology</i> , 2012, 65, 519-542.	2.0	58
53	BIM Expression in Treatment-Naïve Cancers Predicts Responsiveness to Kinase Inhibitors. <i>Cancer Discovery</i> , 2011, 1, 352-365.	9.4	268
54	An ErbB3 Antibody, MM-121, Is Active in Cancers with Ligand-Dependent Activation. <i>Cancer Research</i> , 2010, 70, 2485-2494.	0.9	250

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55	Differences underlying EGFR and HER2 oncogene addiction. <i>Cell Cycle</i> , 2010, 9, 851-852.	2.6	39
56	Differential induction of apoptosis in HER2 and EGFR addicted cancers following PI3K inhibition. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 19503-19508.	7.1	286
57	Acquired resistance to EGFR tyrosine kinase inhibitors in cancer cells is mediated by loss of IGF-binding proteins. <i>Journal of Clinical Investigation</i> , 2008, 118, 2609-19.	8.2	443
58	Thematic Review Series: Sphingolipids. Ganglioside GM3 suppresses the proangiogenic effects of vascular endothelial growth factor and ganglioside GD1a. <i>Journal of Lipid Research</i> , 2008, 49, 929-938.	4.2	48
59	Inhibition of Cyclin-dependent Kinase-2 Induces Apoptosis in Human Diffuse Large B-cell Lymphomas. <i>Cell Cycle</i> , 2007, 6, 2982-2989.	2.6	37
60	Inhibition of phosphatidylinositol 3-kinase-mediated glucose metabolism coincides with resveratrol-induced cell cycle arrest in human diffuse large B-cell lymphomas. <i>Biochemical Pharmacology</i> , 2006, 72, 1246-1256.	4.4	67
61	Resveratrol Induces Cell Cycle Arrest in the OCI-LY18 B-Cell Lymphoma.. <i>Blood</i> , 2005, 106, 4680-4680.	1.4	0