

Anang A Shelat

List of Publications by Year in descending order

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Version: 2024-02-01

101
papers

5,249
citations

94433

37
h-index

98798

67
g-index

110
all docs

110
docs citations

110
times ranked

9707
citing authors

#	ARTICLE	IF	CITATIONS
1	An Evolutionarily Conserved Residue in ABCC2 Regulates the Transport Cycle and Inhibitor Activity. <i>FASEB Journal</i> , 2022, 36, .	0.5	0
2	INSP-09. Using genetically engineered mouse models and patient-derived orthotopic xenografts to develop new therapies for pediatric brain tumors.. <i>Neuro-Oncology</i> , 2022, 24, i188-i188.	1.2	0
3	MAIP: a web service for predicting blood-stage malaria inhibitors. <i>Journal of Cheminformatics</i> , 2021, 13, 13.	6.1	20
4	Modeling and targeting of erythroleukemia by hematopoietic genome editing. <i>Blood</i> , 2021, 137, 1628-1640.	1.4	25
5	Evaluating and evolving a screening library in academia: the St Jude approach. <i>Drug Discovery Today</i> , 2021, 26, 1060-1069.	6.4	6
6	Model-based evaluation of image-guided fractionated whole-brain radiation therapy in pediatric diffuse intrinsic pontine glioma xenografts. <i>CPT: Pharmacometrics and Systems Pharmacology</i> , 2021, 10, 599-610.	2.5	3
7	Patient-derived models recapitulate heterogeneity of molecular signatures and drug response in pediatric high-grade glioma. <i>Nature Communications</i> , 2021, 12, 4089.	12.8	27
8	SLFN11 is Widely Expressed in Pediatric Sarcoma and Induces Variable Sensitization to Replicative Stress Caused By DNA-Damaging Agents. <i>Molecular Cancer Therapeutics</i> , 2021, 20, 2151-2165.	4.1	6
9	Using response surface models to analyze drug combinations. <i>Drug Discovery Today</i> , 2021, 26, 2014-2024.	6.4	4
10	Small Molecule Sequestration of the Intrinsically Disordered Protein, p27Kip1, Within Soluble Oligomers. <i>Journal of Molecular Biology</i> , 2021, 433, 167120.	4.2	16
11	Data vignettes for the application of response surface models in drug combination analysis. <i>Data in Brief</i> , 2021, 38, 107400.	1.0	0
12	17-DMAG dually inhibits Hsp90 and histone lysine demethylases in alveolar rhabdomyosarcoma. <i>IScience</i> , 2021, 24, 101996.	4.1	7
13	Phenyl-Glutarimides: Alternative Cereblon Binders for the Design of PROTACs. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 26663-26670.	13.8	45
14	Loss of PTEN in Pediatric AML Confers Sensitivity to PARP Inhibition. <i>Blood</i> , 2021, 138, 3446-3446.	1.4	0
15	A phase I trial of talazoparib and irinotecan with and without temozolomide in children and young adults with recurrent or refractory solid malignancies. <i>European Journal of Cancer</i> , 2020, 137, 204-213.	2.8	39
16	Bromodomain-Selective BET Inhibitors Are Potent Antitumor Agents against MYC-Driven Pediatric Cancer. <i>Cancer Research</i> , 2020, 80, 3507-3518.	0.9	28
17	MYCN amplification and ATRX mutations are incompatible in neuroblastoma. <i>Nature Communications</i> , 2020, 11, 913.	12.8	66
18	A critical evaluation of methods to interpret drug combinations. <i>Scientific Reports</i> , 2020, 10, 5144.	3.3	10

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19	ABCG2 requires a single aromatic amino acid to clamp substrates and inhibitors into the binding pocket. <i>FASEB Journal</i> , 2020, 34, 4890-4903.	0.5	30
20	A high-throughput screen indicates gemcitabine and JAK inhibitors may be useful for treating pediatric AML. <i>Nature Communications</i> , 2019, 10, 2189.	12.8	26
21	Identification of substituted 5-membered heterocyclic compounds as potential anti-leukemic agents. <i>European Journal of Medicinal Chemistry</i> , 2019, 164, 391-398.	5.5	7
22	Combinatorial screening using orthotopic patient derived xenograft-expanded early phase cultures of osteosarcoma identify novel therapeutic drug combinations. <i>Cancer Letters</i> , 2019, 442, 262-270.	7.2	23
23	Determining the Molecular Characteristics of How Ligands Interact with an ABC Transporter. <i>FASEB Journal</i> , 2019, 33, 507.4.	0.5	0
24	Novel vitexin-inspired scaffold against leukemia. <i>European Journal of Medicinal Chemistry</i> , 2018, 146, 501-510.	5.5	17
25	Establishing a Preclinical Multidisciplinary Board for Brain Tumors. <i>Clinical Cancer Research</i> , 2018, 24, 1654-1666.	7.0	12
26	The role of ZA channel water-mediated interactions in the design of bromodomain-selective BET inhibitors. <i>Journal of Molecular Graphics and Modelling</i> , 2018, 81, 197-210.	2.4	18
27	Exploiting a water network to achieve enthalpy-driven, bromodomain-selective BET inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2018, 26, 25-36.	3.0	23
28	Virtual-screening workflow tutorials and prospective results from the Teach-Discover-Treat competition 2014 against malaria. <i>F1000Research</i> , 2018, 6, 1136.	1.6	6
29	Identification of Therapeutic Targets in Rhabdomyosarcoma through Integrated Genomic, Epigenomic, and Proteomic Analyses. <i>Cancer Cell</i> , 2018, 34, 411-426.e19.	16.8	106
30	Alteration of RNA Splicing by Small-Molecule Inhibitors of the Interaction between NHP2L1 and U4. <i>SLAS Discovery</i> , 2018, 23, 164-173.	2.7	14
31	Is Inhibitor Binding the Sole Requirement in Determining Inhibition of ABCG2 Mediated Transport?. <i>FASEB Journal</i> , 2018, 32, 693.9.	0.5	0
32	Multiplex CRISPR/Cas9-Based Genome Editing of Mouse Hematopoietic Stem Cells Recapitulates Acute Erythroid Leukemia and Identifies Therapeutic Targets. <i>Blood</i> , 2018, 132, 5-5.	1.4	0
33	Shared Consensus Machine Learning Models for Predicting Blood Stage Malaria Inhibition. <i>Journal of Chemical Information and Modeling</i> , 2017, 57, 445-453.	5.4	15
34	Targeting Metabolic Reprogramming by Influenza Infection for Therapeutic Intervention. <i>Cell Reports</i> , 2017, 19, 1640-1653.	6.4	127
35	Orthotopic patient-derived xenografts of paediatric solid tumours. <i>Nature</i> , 2017, 549, 96-100.	27.8	223
36	Ventromorphins: A New Class of Small Molecule Activators of the Canonical BMP Signaling Pathway. <i>ACS Chemical Biology</i> , 2017, 12, 2436-2447.	3.4	20

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37	Targeting Histone Demethylases in MYC-Driven Neuroblastomas with Ciclopirox. <i>Cancer Research</i> , 2017, 77, 4626-4638.	0.9	42
38	Discovery of novel, orally bioavailable, antileishmanial compounds using phenotypic screening. <i>PLoS Neglected Tropical Diseases</i> , 2017, 11, e0006157.	3.0	23
39	Blocking an N-terminal acetylation-dependent protein interaction inhibits an E3 ligase. <i>Nature Chemical Biology</i> , 2017, 13, 850-857.	8.0	80
40	Targeting the cell cycle for cancer therapy in rhabdomyosarcoma.. <i>Journal of Clinical Oncology</i> , 2017, 35, 10535-10535.	1.6	1
41	PARP inhibitor combination therapy in desmoplastic small round cell tumors.. <i>Journal of Clinical Oncology</i> , 2017, 35, e23212-e23212.	1.6	4
42	Virtual-screening workflow tutorials and prospective results from the Teach-Discover-Treat competition 2014 against malaria. <i>F1000Research</i> , 2017, 6, 1136.	1.6	7
43	A biomarker-guided approach to combining PARP inhibitors with radiotherapy in pediatric solid tumors.. <i>Journal of Clinical Oncology</i> , 2017, 35, 10556-10556.	1.6	1
44	The second European interdisciplinary Ewing sarcoma research summit - A joint effort to deconstructing the multiple layers of a complex disease. <i>Oncotarget</i> , 2016, 7, 8613-8624.	1.8	55
45	BRAID: A Unifying Paradigm for the Analysis of Combined Drug Action. <i>Scientific Reports</i> , 2016, 6, 25523.	3.3	63
46	Evaluation of artemisinins for the treatment of acute myeloid leukemia. <i>Cancer Chemotherapy and Pharmacology</i> , 2016, 77, 1231-1243.	2.3	41
47	Diversity-oriented natural product platform identifies plant constituents targeting <i>Plasmodium falciparum</i> . <i>Malaria Journal</i> , 2016, 15, 270.	2.3	4
48	Synthesis and evaluation of colletoic acid core derivatives. <i>European Journal of Medicinal Chemistry</i> , 2016, 110, 126-132.	5.5	8
49	Preclinical studies of 5-fluoro-2-deoxycytidine and tetrahydrouridine in pediatric brain tumors. <i>Journal of Neuro-Oncology</i> , 2016, 126, 225-234.	2.9	11
50	The Childhood Solid Tumor Network: A new resource for the developmental biology and oncology research communities. <i>Developmental Biology</i> , 2016, 411, 287-293.	2.0	63
51	Nuclear ULK1 promotes cell death in response to oxidative stress through PARP1. <i>Cell Death and Differentiation</i> , 2016, 23, 216-230.	11.2	61
52	Robust Classification of Small-Molecule Mechanism of Action Using a Minimalist High-Content Microscopy Screen and Multidimensional Phenotypic Trajectory Analysis. <i>PLoS ONE</i> , 2016, 11, e0149439.	2.5	17
53	Discovery of Small Molecules that Inhibit the Disordered Protein, p27Kip1. <i>Scientific Reports</i> , 2015, 5, 15686.	3.3	70
54	A High-Throughput Screen Reveals New Small-Molecule Activators and Inhibitors of Pantothenate Kinases. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 1563-1568.	6.4	28

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55	Development and characterization of a human orthotopic neuroblastoma xenograft. <i>Developmental Biology</i> , 2015, 407, 344-355.	2.0	30
56	Systematic Screening Identifies Dual PI3K and mTOR Inhibition as a Conserved Therapeutic Vulnerability in Osteosarcoma. <i>Clinical Cancer Research</i> , 2015, 21, 3216-3229.	7.0	58
57	ABCG2 Transporter Expression Impacts Group 3 Medulloblastoma Response to Chemotherapy. <i>Cancer Research</i> , 2015, 75, 3879-3889.	0.9	30
58	Targeting the DNA Repair Pathway in Ewing Sarcoma. <i>Cell Reports</i> , 2014, 9, 829-840.	6.4	141
59	(+)-SJ733, a clinical candidate for malaria that acts through ATP4 to induce rapid host-mediated clearance of <i>Plasmodium</i> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, E5455-62.	7.1	199
60	A Screening-Based Approach to Circumvent Tumor Microenvironment-Driven Intrinsic Resistance to BCR-ABL+ Inhibitors in Ph+ Acute Lymphoblastic Leukemia. <i>Journal of Biomolecular Screening</i> , 2014, 19, 158-167.	2.6	10
61	Pemetrexed and Gemcitabine as Combination Therapy for the Treatment of Group3 Medulloblastoma. <i>Cancer Cell</i> , 2014, 25, 516-529.	16.8	128
62	Development of a Cell-Based, High-Throughput Screening Assay for ATM Kinase Inhibitors. <i>Journal of Biomolecular Screening</i> , 2014, 19, 538-546.	2.6	32
63	UPLC-MS-ELSD-PDA as a Powerful Dereplication Tool to Facilitate Compound Identification from Small-Molecule Natural Product Libraries. <i>Journal of Natural Products</i> , 2014, 77, 902-909.	3.0	41
64	Ligand Binding Mode Prediction by Docking: Mdm2/Mdmx Inhibitors as a Case Study. <i>Journal of Chemical Information and Modeling</i> , 2014, 54, 648-659.	5.4	35
65	Targeted Inhibition of the MLL Transcriptional Complex By Proteasome Inhibitors Elicits a High Response Rate in Relapsed/Refractory MLL Rearranged Leukemia. <i>Blood</i> , 2014, 124, 972-972.	1.4	8
66	Incorporating PARP inhibitors in translational studies for Ewing's sarcoma. <i>Journal of Clinical Oncology</i> , 2014, 32, e21025-e21025.	1.6	0
67	Preclinical evaluation of PARP inhibitors in combination with DNA-damaging agents in a Ewing sarcoma orthotopic xenograft model. <i>Journal of Clinical Oncology</i> , 2014, 32, 10073-10073.	1.6	0
68	Molecular analysis of solid tumors (MAST): A protocol for comprehensive preclinical evaluation of pediatric solid tumors. <i>Journal of Clinical Oncology</i> , 2014, 32, 10036-10036.	1.6	0
69	Metabolic Activation of CaMKII by Coenzyme A. <i>Molecular Cell</i> , 2013, 52, 325-339.	9.7	35
70	Metabolic Activation of CaMKII by Coenzyme A. <i>Molecular Cell</i> , 2013, 52, 468.	9.7	1
71	Targeting Oxidative Stress in Embryonal Rhabdomyosarcoma. <i>Cancer Cell</i> , 2013, 24, 710-724.	16.8	252
72	Identification of Small Molecule Activators of BMP Signaling. <i>PLoS ONE</i> , 2013, 8, e59045.	2.5	61

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73	Global Phenotypic Screening for Antimalarials. <i>Chemistry and Biology</i> , 2012, 19, 116-129.	6.0	120
74	An Integrated InÂVitro and InÂVivo High-Throughput Screen Identifies Treatment Leads for Ependymoma. <i>Cancer Cell</i> , 2011, 20, 384-399.	16.8	105
75	A High-Content Screen Identifies Inhibitors of Nuclear Export of Forkhead Transcription Factors. <i>Journal of Biomolecular Screening</i> , 2011, 16, 394-404.	2.6	12
76	An Automated Approach to Efficiently Reformat a Large Collection of Compounds. <i>Current Chemical Genomics</i> , 2011, 5, 42-47.	2.0	5
77	Chemical genetics of <i>Plasmodium falciparum</i> . <i>Nature</i> , 2010, 465, 311-315.	27.8	515
78	Discovery of Potent and Selective Inhibitors of <i>Trypanosoma brucei</i> Ornithine Decarboxylase. <i>Journal of Biological Chemistry</i> , 2010, 285, 16771-16781.	3.4	33
79	Optimization of a Non-Radioactive High-Throughput Assay for Decarboxylase Enzymes. <i>Assay and Drug Development Technologies</i> , 2010, 8, 175-185.	1.2	15
80	Identification and Characterization of the First Small Molecule Inhibitor of MDMX. <i>Journal of Biological Chemistry</i> , 2010, 285, 10786-10796.	3.4	171
81	Automated High-Throughput System to Fractionate Plant Natural Products for Drug Discovery. <i>Journal of Natural Products</i> , 2010, 73, 751-754.	3.0	79
82	A road less traveled by: Exploring a decade of Ellman chemistry. <i>Bioorganic and Medicinal Chemistry</i> , 2009, 17, 1088-1093.	3.0	4
83	Structure-Guided Development of Selective TbcA Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 6489-6493.	6.4	25
84	Discovery of trypanocidal thiosemicarbazone inhibitors of rhodesain and TbcA. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 2883-2885.	2.2	52
85	Development of Potent Purine-Derived Nitrile Inhibitors of the Trypanosomal Protease TbcA. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 545-552.	6.4	58
86	Assay Optimization and Screening of RNA-Protein Interactions by AlphaScreen. <i>Journal of Biomolecular Screening</i> , 2007, 12, 946-955.	2.6	38
87	Design of a selective chemical probe for class I PDZ domains. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2007, 17, 546-548.	2.2	13
88	The interdependence between screening methods and screening libraries. <i>Current Opinion in Chemical Biology</i> , 2007, 11, 244-251.	6.1	54
89	Scaffold composition and biological relevance of screening libraries. <i>Nature Chemical Biology</i> , 2007, 3, 442-446.	8.0	164
90	Discovery of Trypanocidal Compounds by Whole Cell HTS of <i>Trypanosoma brucei</i> . <i>Chemical Biology and Drug Design</i> , 2006, 67, 355-363.	3.2	97

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91	Searching for New Antimalarial Therapeutics amongst Known Drugs. <i>Chemical Biology and Drug Design</i> , 2006, 67, 409-416.	3.2	110
92	A High-Throughput Screening Method to Identify Small Molecule Inhibitors of Thyroid Hormone Receptor Coactivator Binding. <i>Science Signaling</i> , 2006, 2006, p13-p13.	3.6	27
93	High-throughput assays for promiscuous inhibitors. , 2005, 1, 146-148.		300
94	Discovery of Small Molecule Inhibitors of the Interaction of the Thyroid Hormone Receptor with Transcriptional Coregulators. <i>Journal of Biological Chemistry</i> , 2005, 280, 43048-43055.	3.4	96
95	A Solid-Phase Approach to the Phallotoxins: Total Synthesis of [Ala7]-Phalloidin. <i>Journal of Organic Chemistry</i> , 2005, 70, 4578-4584.	3.2	38
96	Catalysis, Specificity, and ACP Docking Site of <i>Streptomyces coelicolor</i> Malonyl-CoA:ACP Transacylase. <i>Structure</i> , 2003, 11, 147-154.	3.3	125
97	Benzoflavone activators of the cystic fibrosis transmembrane conductance regulator: towards a pharmacophore model for the nucleotide-binding domain. <i>Bioorganic and Medicinal Chemistry</i> , 2003, 11, 4113-4120.	3.0	62
98	Structurally Conserved Amino Acid W501 Is Required for RNA Helicase Activity but Is Not Essential for DNA Helicase Activity of Hepatitis C Virus NS3 Protein. <i>Journal of Virology</i> , 2003, 77, 571-582.	3.4	23
99	Nanomolar Affinity Small Molecule Correctors of Defective $\Delta F508$ -CFTR Chloride Channel Gating. <i>Journal of Biological Chemistry</i> , 2003, 278, 35079-35085.	3.4	192
100	Phenyl-Glutarimides: Alternative Cereblon Binders for the Design of PROTACs. <i>Angewandte Chemie</i> , 0, , .	2.0	1
101	Integrated High-Throughput Screen to Identify Novel Treatment Leads for Pediatric Acute Myeloid Leukemia. <i>SSRN Electronic Journal</i> , 0, , .	0.4	0