

Hongkai Zhang

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/136027/hongkai-zhang-publications-by-year.pdf>

Version: 2024-04-25

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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

29
papers

466
citations

11
h-index

21
g-index

36
ext. papers

610
ext. citations

7.9
avg, IF

3.2
L-index

#	Paper	IF	Citations
29	Structures of Omicron spike complexes and implications for neutralizing antibody development.. <i>Cell Reports</i> , 2022 , 110770	10.6	4
28	Antigen-Specific Stimulation and Expansion of CAR-T Cells Using Membrane Vesicles as Target Cell Surrogates. <i>Small</i> , 2021 , 17, e2102643	11	1
27	Molecular deconvolution of the neutralizing antibodies induced by an inactivated SARS-CoV-2 virus vaccine. <i>Protein and Cell</i> , 2021 , 12, 818-823	7.2	5
26	High-throughput functional screening for next-generation cancer immunotherapy using droplet-based microfluidics. <i>Science Advances</i> , 2021 , 7,	14.3	12
25	Reshaping the Immune Microenvironment by Oncolytic Herpes Simplex Virus in Murine Pancreatic Ductal Adenocarcinoma. <i>Molecular Therapy</i> , 2021 , 29, 744-761	11.7	11
24	A general Fc engineering platform for the next generation of antibody therapeutics. <i>Theranostics</i> , 2021 , 11, 1901-1917	12.1	5
23	Targeting FSTL1 for Multiple Fibrotic and Systemic Autoimmune Diseases. <i>Molecular Therapy</i> , 2021 , 29, 347-364	11.7	5
22	Phenotypic selection with an intrabody library reveals an anti-apoptotic function of PKM2 requiring Mitofusin-1. <i>PLoS Biology</i> , 2019 , 17, e2004413	9.7	6
21	The interaction between Vav1 and EBNA1 promotes survival of Burkitt's lymphoma cells by down-regulating the expression of Bim. <i>Biochemical and Biophysical Research Communications</i> , 2019 , 511, 787-793	3.4	3
20	An agonist antibody prefers relapsed AML for induction of cells that kill each other. <i>Scientific Reports</i> , 2019 , 9, 3494	4.9	
19	Identification of novel Kv1.3 targeting venom peptides by a single round of autocrine-based selection. <i>Biochemical and Biophysical Research Communications</i> , 2019 , 509, 954-959	3.4	
18	Affinity maturation of an TpoR targeting antibody in full-length IgG form for enhanced agonist activity. <i>Protein Engineering, Design and Selection</i> , 2018 , 31, 233-241	1.9	6
17	Interleukin-5 suppresses Vascular Endothelial Growth Factor-induced angiogenesis through STAT5 signaling. <i>Cytokine</i> , 2018 , 110, 397-403	4	7
16	High-throughput reformatting of phage-displayed antibody fragments to IgGs by one-step emulsion PCR. <i>Protein Engineering, Design and Selection</i> , 2018 , 31, 427-436	1.9	2
15	A Proximity-Based Assay for Identification of Ligand and Membrane Protein Interaction in Living Cells. <i>Methods in Molecular Biology</i> , 2017 , 1575, 215-222	1.4	
14	Antibody-Mediated Inhibition of Tspan12 Ameliorates Vasoproliferative Retinopathy Through Suppression of E-Catenin Signaling. <i>Circulation</i> , 2017 , 136, 180-195	16.7	12
13	Autocrine-Based Selection of Drugs That Target Ion Channels from Combinatorial Venom Peptide Libraries. <i>Angewandte Chemie</i> , 2016 , 128, 9452-9456	3.6	

12	Titelbild: Autocrine-Based Selection of Drugs That Target Ion Channels from Combinatorial Venom Peptide Libraries (Angew. Chem. 32/2016). <i>Angewandte Chemie</i> , 2016 , 128, 9245-9245	3.6	
11	Autocrine-Based Selection of Drugs That Target Ion Channels from Combinatorial Venom Peptide Libraries. <i>Angewandte Chemie - International Edition</i> , 2016 , 55, 9306-10	16.4	12
10	Agonist antibody that induces human malignant cells to kill one another. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015 , 112, E6158-65	11.5	11
9	Antibodies from combinatorial libraries use functional receptor pleiotropism to regulate cell fates. <i>Quarterly Reviews of Biophysics</i> , 2015 , 48, 389-94	7	12
8	Autocrine selection of a GLP-1R G-protein biased agonist with potent antidiabetic effects. <i>Nature Communications</i> , 2015 , 6, 8918	17.4	90
7	Selection of multiple agonist antibodies from intracellular combinatorial libraries reveals that cellular receptors are functionally pleiotropic. <i>Current Opinion in Chemical Biology</i> , 2015 , 26, 1-7	9.7	16
6	A proximity based general method for identification of ligand and receptor interactions in living cells. <i>Biochemical and Biophysical Research Communications</i> , 2014 , 454, 251-5	3.4	13
5	Prevention of cell death by antibodies selected from intracellular combinatorial libraries. <i>Chemistry and Biology</i> , 2014 , 21, 274-83		32
4	Selecting agonists from single cells infected with combinatorial antibody libraries. <i>Chemistry and Biology</i> , 2013 , 20, 734-41		39
3	Selection of antibodies that regulate phenotype from intracellular combinatorial antibody libraries. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012 , 109, 15728-33	11.5	53
2	Magnetic and pH-responsive nanocarriers with multilayer core-shell architecture for anticancer drug delivery. <i>Journal of Materials Chemistry</i> , 2008 , 18, 5104		105
1	In vivo selection of phage sequences and characterization of peptide-specific binding to breast cancer cells. <i>Chinese Journal of Clinical Oncology</i> , 2008 , 5, 128-131		1