

# Yuanmei Zhu

## List of Publications by Year in descending order

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28  
papers

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citations

643344

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591227

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times ranked

937  
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#	ARTICLE	IF	CITATIONS
1	Cell membrane-anchored anti-HIV single-chain antibodies and bifunctional inhibitors targeting the gp41 fusion protein: new strategies for HIV gene therapy. <i>Emerging Microbes and Infections</i> , 2022, 11, 30-49.	3.0	5
2	Efficient treatment and pre-exposure prophylaxis in rhesus macaques by an HIV fusion-inhibitory lipopeptide. <i>Cell</i> , 2022, 185, 131-144.e18.	13.5	24
3	Design of a Bispecific HIV Entry Inhibitor Targeting the Cell Receptor CD4 and Viral Fusion Protein Gp41. <i>Frontiers in Cellular and Infection Microbiology</i> , 2022, 12, .	1.8	4
4	SARS-CoV-2 fusion-inhibitory lipopeptides maintain high potency against divergent variants of concern including Omicron. <i>Emerging Microbes and Infections</i> , 2022, 11, 1819-1827.	3.0	10
5	Structure-based design and characterization of novel fusion-inhibitory lipopeptides against SARS-CoV-2 and emerging variants. <i>Emerging Microbes and Infections</i> , 2021, 10, 1227-1240.	3.0	17
6	Generation of HIV-resistant cells with a single-domain antibody: implications for HIV-1 gene therapy. <i>Cellular and Molecular Immunology</i> , 2021, 18, 660-674.	4.8	9
7	SARS-CoV-2-derived fusion inhibitor lipopeptides exhibit highly potent and broad-spectrum activity against divergent human coronaviruses. <i>Signal Transduction and Targeted Therapy</i> , 2021, 6, 294.	7.1	20
8	Pan-coronavirus fusion inhibitors possess potent inhibitory activity against HIV-1, HIV-2, and simian immunodeficiency virus. <i>Emerging Microbes and Infections</i> , 2021, 10, 810-821.	3.0	15
9	Cross-reactive neutralization of SARS-CoV-2 by serum antibodies from recovered SARS patients and immunized animals. <i>Science Advances</i> , 2020, 6, .	4.7	57
10	Design of Potent Membrane Fusion Inhibitors against SARS-CoV-2, an Emerging Coronavirus with High Fusogenic Activity. <i>Journal of Virology</i> , 2020, 94, .	1.5	164
11	Therapeutic Efficacy and Resistance Selection of a Lipopeptide Fusion Inhibitor in Simian Immunodeficiency Virus-Infected Rhesus Macaques. <i>Journal of Virology</i> , 2020, 94, .	1.5	3
12	Structural and Functional Characterization of the Secondary Mutation N126K Selected by Various HIV-1 Fusion Inhibitors. <i>Viruses</i> , 2020, 12, 326.	1.5	2
13	Conserved Residue Asn-145 in the C-Terminal Heptad Repeat Region of HIV-1 gp41 is Critical for Viral Fusion and Regulates the Antiviral Activity of Fusion Inhibitors. <i>Viruses</i> , 2019, 11, 609.	1.5	4
14	A Membrane-Anchored Short-Peptide Fusion Inhibitor Fully Protects Target Cells from Infections of Human Immunodeficiency Virus Type 1 (HIV-1), HIV-2, and Simian Immunodeficiency Virus. <i>Journal of Virology</i> , 2019, 93, .	1.5	15
15	Design and Characterization of Cholesterylated Peptide HIV-1/2 Fusion Inhibitors with Extremely Potent and Long-Lasting Antiviral Activity. <i>Journal of Virology</i> , 2019, 93, .	1.5	34
16	Monotherapy with a low-dose lipopeptide HIV fusion inhibitor maintains long-term viral suppression in rhesus macaques. <i>PLoS Pathogens</i> , 2019, 15, e1007552.	2.1	30
17	The Tryptophan-Rich Motif of HIV-1 gp41 Can Interact with the N-Terminal Deep Pocket Site: New Insights into the Structure and Function of gp41 and Its Inhibitors. <i>Journal of Virology</i> , 2019, 94, .	1.5	7
18	Structural and functional characterization of HIV-1 cell fusion inhibitor T20. <i>Aids</i> , 2019, 33, 1-11.	1.0	38

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19	Exceptional potency and structural basis of a T1249-derived lipopeptide fusion inhibitor against HIV-1, HIV-2, and simian immunodeficiency virus. <i>Journal of Biological Chemistry</i> , 2018, 293, 5323-5334.	1.6	27
20	Mechanism of HIV-1 Resistance to an Electronically Constrained Î±-Helical Peptide Membrane Fusion Inhibitor. <i>Journal of Virology</i> , 2018, 92, .	1.5	12
21	Molecular mechanism of HIV-1 resistance to sifuvirtide, a clinical trialâ€‘approved membrane fusion inhibitor. <i>Journal of Biological Chemistry</i> , 2018, 293, 12703-12718.	1.6	20
22	Structural Insights into the Mechanisms of Action of Short-Peptide HIV-1 Fusion Inhibitors Targeting the Gp41 Pocket. <i>Frontiers in Cellular and Infection Microbiology</i> , 2018, 8, 51.	1.8	14
23	Structural and Functional Characterization of Membrane Fusion Inhibitors with Extremely Potent Activity against Human Immunodeficiency Virus Type 1 (HIV-1), HIV-2, and Simian Immunodeficiency Virus. <i>Journal of Virology</i> , 2018, 92, .	1.5	30
24	Design of Novel HIV-1/2 Fusion Inhibitors with High Therapeutic Efficacy in Rhesus Monkey Models. <i>Journal of Virology</i> , 2018, 92, .	1.5	29
25	A Lipopeptide HIV-1/2 Fusion Inhibitor with Highly Potent <i>In Vitro</i> , <i>Ex Vivo</i> , and <i>In Vivo</i> Antiviral Activity. <i>Journal of Virology</i> , 2017, 91, .	1.5	53
26	Enfuvirtide (T20)-Based Lipopeptide Is a Potent HIV-1 Cell Fusion Inhibitor: Implications for Viral Entry and Inhibition. <i>Journal of Virology</i> , 2017, 91, .	1.5	65
27	A Helical Short-Peptide Fusion Inhibitor with Highly Potent Activity against Human Immunodeficiency Virus Type 1 (HIV-1), HIV-2, and Simian Immunodeficiency Virus. <i>Journal of Virology</i> , 2017, 91, .	1.5	35
28	Identification of a novel HIV-1-neutralizing antibody from a CRF07_BC-infected Chinese donor. <i>Oncotarget</i> , 2017, 8, 63047-63063.	0.8	6