

# Yuxian He

## List of Publications by Year in descending order

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76  
papers

6,466  
citations

109264

35  
h-index

76872

74  
g-index

77  
all docs

77  
docs citations

77  
times ranked

8848  
citing authors

| #  | ARTICLE   | IF   | CITATIONS |
|----|---|------|-----------|
| 1  | The spike protein of SARS-CoV is a target for vaccine and therapeutic development. <i>Nature Reviews Microbiology</i> , 2009, 7, 226-236.   | 13.6 | 1,405     |
| 2  | Adaptation of SARS-CoV-2 in BALB/c mice for testing vaccine efficacy. <i>Science</i> , 2020, 369, 1603-1607.  | 6.0  | 678       |
| 3  | Interaction between heptad repeat 1 and 2 regions in spike protein of SARS-associated coronavirus: implications for virus fusogenic mechanism and identification of fusion inhibitors. <i>Lancet</i> , The, 2004, 363, 938-947.                     | 6.3  | 476       |
| 4  | Receptor-binding domain of SARS-CoV spike protein induces highly potent neutralizing antibodies: implication for developing subunit vaccine. <i>Biochemical and Biophysical Research Communications</i> , 2004, 324, 773-781.                       | 1.0  | 366       |
| 5  | N-Substituted Pyrrole Derivatives as Novel Human Immunodeficiency Virus Type 1 Entry Inhibitors That Interfere with the gp41 Six-Helix Bundle Formation and Block Virus Fusion. <i>Antimicrobial Agents and Chemotherapy</i> , 2004, 48, 4349-4359. | 1.4  | 253       |
| 6  | Receptor-Binding Domain of Severe Acute Respiratory Syndrome Coronavirus Spike Protein Contains Multiple Conformation-Dependent Epitopes that Induce Highly Potent Neutralizing Antibodies. <i>Journal of Immunology</i> , 2005, 174, 4908-4915.    | 0.4  | 230       |
| 7  | Design and Evaluation of Sifuvirtide, a Novel HIV-1 Fusion Inhibitor. <i>Journal of Biological Chemistry</i> , 2008, 283, 11126-11134.  | 1.6  | 200       |
| 8  | 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       |
| 9  | Identification of Immunodominant Sites on the Spike Protein of Severe Acute Respiratory Syndrome (SARS) Coronavirus: Implication for Developing SARS Diagnostics and Vaccines. <i>Journal of Immunology</i> , 2004, 173, 4050-4057.                 | 0.4  | 145       |
| 10 | Potent HIV fusion inhibitors against Enfuvirtide-resistant HIV-1 strains. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 16332-16337.  | 3.3  | 129       |
| 11 | Inactivated SARS-CoV vaccine elicits high titers of spike protein-specific antibodies that block receptor binding and virus entry. <i>Biochemical and Biophysical Research Communications</i> , 2004, 325, 445-452.                                 | 1.0  | 120       |
| 12 | Antigenic and Immunogenic Characterization of Recombinant Baculovirus-Expressed Severe Acute Respiratory Syndrome Coronavirus Spike Protein: Implication for Vaccine Design. <i>Journal of Virology</i> , 2006, 80, 5757-5767.                      | 1.5  | 113       |
| 13 | Cross-Neutralization of Human and Palm Civet Severe Acute Respiratory Syndrome Coronaviruses by Antibodies Targeting the Receptor-Binding Domain of Spike Protein. <i>Journal of Immunology</i> , 2006, 176, 6085-6092.                             | 0.4  | 108       |
| 14 | Identification of SARS-CoV RBD-targeting monoclonal antibodies with cross-reactive or neutralizing activity against SARS-CoV-2. <i>Antiviral Research</i> , 2020, 179, 104820.  | 1.9  | 106       |
| 15 | Identification of a critical neutralization determinant of severe acute respiratory syndrome (SARS)-associated coronavirus: importance for designing SARS vaccines. <i>Virology</i> , 2005, 334, 74-82.   | 1.1  | 103       |
| 16 | Identification of Immunodominant Epitopes on the Membrane Protein of the Severe Acute Respiratory Syndrome-Associated Coronavirus. <i>Journal of Clinical Microbiology</i> , 2005, 43, 3718-3726.   | 1.8  | 81        |
| 17 | Identification of a Critical Motif for the Human Immunodeficiency Virus Type 1 (HIV-1) gp41 Core Structure: Implications for Designing Novel Anti-HIV Fusion Inhibitors. <i>Journal of Virology</i> , 2008, 82, 6349-6358.                          | 1.5  | 81        |
| 18 | Conserved Residue Lys574 in the Cavity of HIV-1 Gp41 Coiled-coil Domain Is Critical for Six-helix Bundle Stability and Virus Entry. <i>Journal of Biological Chemistry</i> , 2007, 282, 25631-25639.  | 1.6  | 75        |

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|----|---|-----|-----------|
| 19 | Mapping of Antigenic Sites on the Nucleocapsid Protein of the Severe Acute Respiratory Syndrome Coronavirus. <i>Journal of Clinical Microbiology</i> , 2004, 42, 5309-5314.   | 1.8 | 70        |
| 20 | Long-Term Persistence of Robust Antibody and Cytotoxic T Cell Responses in Recovered Patients Infected with SARS Coronavirus. <i>PLoS ONE</i> , 2006, 1, e24.   | 1.1 | 69        |
| 21 | Potent and persistent antibody responses against the receptor-binding domain of SARS-CoV spike protein in recovered patients. <i>Virology Journal</i> , 2010, 7, 299.   | 1.4 | 69        |
| 22 | 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        |
| 23 | Conserved Salt Bridge between the N- and C-Terminal Heptad Repeat Regions of the Human Immunodeficiency Virus Type 1 gp41 Core Structure Is Critical for Virus Entry and Inhibition. <i>Journal of Virology</i> , 2008, 82, 11129-11139.          | 1.5 | 60        |
| 24 | Broad Antiviral Activity and Crystal Structure of HIV-1 Fusion Inhibitor Sifuvirtide. <i>Journal of Biological Chemistry</i> , 2012, 287, 6788-6796.  | 1.6 | 60        |
| 25 | Biophysical Property and Broad Anti-HIV Activity of Albuvirtide, a 3-Maleimimidopropionic Acid-Modified Peptide Fusion Inhibitor. <i>PLoS ONE</i> , 2012, 7, e32599.  | 1.1 | 57        |
| 26 | 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        |
| 27 | Identification and characterization of novel neutralizing epitopes in the receptor-binding domain of SARS-CoV spike protein: Revealing the critical antigenic determinants in inactivated SARS-CoV vaccine. <i>Vaccine</i> , 2006, 24, 5498-5508. | 1.7 | 55        |
| 28 | Short-peptide fusion inhibitors with high potency against wild-type and enfuvirtide-resistant HIV-1. <i>FASEB Journal</i> , 2013, 27, 1203-1213.  | 0.2 | 54        |
| 29 | Development of potent and long-acting HIV-1 fusion inhibitors. <i>Aids</i> , 2016, 30, 1187-1196.   | 1.0 | 53        |
| 30 | A Lipopeptide HIV-1/2 Fusion Inhibitor with Highly Potent In Vitro, Ex Vivo, and In Vivo Antiviral Activity. <i>Journal of Virology</i> , 2017, 91, .   | 1.5 | 53        |
| 31 | Synthesized Peptide Inhibitors of HIV-1 gp41-dependent Membrane Fusion. <i>Current Pharmaceutical Design</i> , 2013, 19, 1800-1809.   | 0.9 | 51        |
| 32 | In Vitro Selection and Characterization of HIV-1 Variants with Increased Resistance to Sifuvirtide, a Novel HIV-1 Fusion Inhibitor. <i>Journal of Biological Chemistry</i> , 2011, 286, 3277-3287.  | 1.6 | 47        |
| 33 | The M-T Hook Structure Is Critical for Design of HIV-1 Fusion Inhibitors. <i>Journal of Biological Chemistry</i> , 2012, 287, 34558-34568.  | 1.6 | 47        |
| 34 | Design of a highly potent HIV-1 fusion inhibitor targeting the gp41 pocket. <i>Aids</i> , 2015, 29, 13-21.  | 1.0 | 44        |
| 35 | Discovery of Critical Residues for Viral Entry and Inhibition through Structural Insight of HIV-1 Fusion Inhibitor CP621652. <i>Journal of Biological Chemistry</i> , 2012, 287, 20281-20289.   | 1.6 | 42        |
| 36 | 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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|----|--|------|-----------|
| 37 | Vaccine Design for Severe Acute Respiratory Syndrome Coronavirus. <i>Viral Immunology</i> , 2005, 18, 327-332.   | 0.6  | 36        |
| 38 | Longitudinal profiles of immunoglobulin G antibodies against severe acute respiratory syndrome coronavirus components and neutralizing activities in recovered patients. <i>Scandinavian Journal of Infectious Diseases</i> , 2011, 43, 515-521. | 1.5  | 36        |
| 39 | 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        |
| 40 | The M-T hook structure increases the potency of HIV-1 fusion inhibitor sifuvirtide and overcomes drug resistance. <i>Journal of Antimicrobial Chemotherapy</i> , 2014, 69, 2759-2769.  | 1.3  | 34        |
| 41 | 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        |
| 42 | Mechanism of HIV-1 Resistance to Short-Peptide Fusion Inhibitors Targeting the Gp41 Pocket. <i>Journal of Virology</i> , 2015, 89, 5801-5811.  | 1.5  | 30        |
| 43 | 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        |
| 44 | 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        |
| 45 | 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        |
| 46 | 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        |
| 47 | 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        |
| 48 | Two M-T hook residues greatly improve the antiviral activity and resistance profile of the HIV-1 fusion inhibitor SC29EK. <i>Retrovirology</i> , 2014, 11, 40.   | 0.9  | 21        |
| 49 | Genetic Pathway of HIV-1 Resistance to Novel Fusion Inhibitors Targeting the Gp41 Pocket. <i>Journal of Virology</i> , 2015, 89, 12467-12479.  | 1.5  | 21        |
| 50 | Identification and characterization of a subpocket on the N-trimer of HIV-1 Gp41. <i>Aids</i> , 2015, 29, 1015-1024.   | 1.0  | 20        |
| 51 | Molecular mechanism of HIV-1 resistance to sifuvirtide, a clinical trial-approved membrane fusion inhibitor. <i>Journal of Biological Chemistry</i> , 2018, 293, 12703-12718.  | 1.6  | 20        |
| 52 | 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        |
| 53 | Structural Basis of Potent and Broad HIV-1 Fusion Inhibitor CP32M. <i>Journal of Biological Chemistry</i> , 2012, 287, 26618-26629.  | 1.6  | 18        |
| 54 | 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        |

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|----|---|-----|-----------|
| 55 | 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        |
| 56 | 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        |
| 57 | 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        |
| 58 | Mechanism of HIV-1 Resistance to an Electronically Constrained $\alpha$ -Helical Peptide Membrane Fusion Inhibitor. <i>Journal of Virology</i> , 2018, 92, .  | 1.5 | 12        |
| 59 | Screening HLA-A-restricted T cell epitopes of SARS-CoV-2 and the induction of CD8+ T cell responses in HLA-A transgenic mice. <i>Cellular and Molecular Immunology</i> , 2021, 18, 2588-2608.                                     | 4.8 | 12        |
| 60 | The N-Terminal Tâ€“T Motif of a Third-Generation HIV-1 Fusion Inhibitor Is Not Required for Binding Affinity and Antiviral Activity. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 6378-6388.                                 | 2.9 | 11        |
| 61 | 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        |
| 62 | Preparation and evaluation of amphipathic lipopeptideâ€“loaded PLGA microspheres as sustainedâ€“release system for AIDS prevention. <i>Engineering in Life Sciences</i> , 2020, 20, 476-484.                                      | 2.0 | 9         |
| 63 | 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         |
| 64 | 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         |
| 65 | 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         |
| 66 | Isolation and characterization of a novel neutralizing antibody targeting the CD4-binding site of HIV-1 gp120. <i>Antiviral Research</i> , 2016, 132, 252-261.  | 1.9 | 5         |
| 67 | 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         |
| 68 | DNA Triplex-Based Complexes Display Anti-HIV-1-Cell Fusion Activity. <i>Nucleic Acid Therapeutics</i> , 2015, 25, 219-225.  | 2.0 | 4         |
| 69 | 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         |
| 70 | 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         |
| 71 | 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         |
| 72 | Defective HIV-1 envelope gene promotes the evolution of the infectious strain through recombination in vitro. <i>BMC Infectious Diseases</i> , 2020, 20, 569.   | 1.3 | 2         |

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|----|---|-----|-----------|
| 73 | 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         |
| 74 | Safety Assessment of Microbicide 2P23 on the Rectal and Vaginal Microbiota and Its Antiviral Activity on HIV Infection. <i>Frontiers in Immunology</i> , 2021, 12, 702172.                                  | 2.2 | 2         |
| 75 | In Vitro Selection and Characterization of HIV-1 Variants with Increased Resistance to LP-40, Enfuvirtide-Based Lipopeptide Inhibitor. <i>International Journal of Molecular Sciences</i> , 2022, 23, 6638. | 1.8 | 0         |
| 76 | Protocol for evaluating CD8+ T cell-mediated immunity in latently SHIV-infected rhesus macaques with HIV fusion-inhibitory lipopeptide monotherapy. <i>STAR Protocols</i> , 2022, 3, 101479.                | 0.5 | 0         |