

# Christopher A Alabi

## List of Publications by Year in descending order

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34  
papers

964  
citations

566801

15  
h-index

454577

30  
g-index

38  
all docs

38  
docs citations

38  
times ranked

1240  
citing authors

#	ARTICLE	IF	CITATIONS
1	Sequence-Defined Polymers via Orthogonal Allyl Acrylamide Building Blocks. <i>Journal of the American Chemical Society</i> , 2014, 136, 13162-13165.	6.6	183
2	Intranasal fusion inhibitory lipopeptide prevents direct-contact SARS-CoV-2 transmission in ferrets. <i>Science</i> , 2021, 371, 1379-1382.	6.0	158
3	Sequence-defined bioactive macrocycles via an acid-catalysed cascade reaction. <i>Nature Chemistry</i> , 2016, 8, 590-596.	6.6	126
4	Sequence-Defined Backbone Modifications Regulate Antibacterial Activity of OligoTEAs. <i>ACS Chemical Biology</i> , 2017, 12, 715-723.	1.6	44
5	Biomimetic Electronic Devices for Measuring Bacterial Membrane Disruption. <i>Advanced Materials</i> , 2018, 30, e1803130.	11.1	43
6	<i>In Vivo</i> Efficacy of Measles Virus Fusion Protein-Derived Peptides Is Modulated by the Properties of Self-Assembly and Membrane Residence. <i>Journal of Virology</i> , 2017, 91, .	1.5	40
7	Thiol-ene Networks from Sequence-Defined Polyurethane Macromers. <i>Journal of the American Chemical Society</i> , 2020, 142, 6729-6736.	6.6	35
8	Dual Site-Specific Antibody Conjugates for Sequential and Orthogonal Cargo Release. <i>Bioconjugate Chemistry</i> , 2019, 30, 1702-1710.	1.8	29
9	Sequence-Defined Oligothioetheramides. <i>Synlett</i> , 2015, 26, 565-571.	1.0	26
10	Effective <i>In Vivo</i> Targeting of Influenza Virus through a Cell-Penetrating/Fusion Inhibitor Tandem Peptide Anchored to the Plasma Membrane. <i>Bioconjugate Chemistry</i> , 2018, 29, 3362-3376.	1.8	26
11	Substrate Design Enables Heterobifunctional, Dual <i>Click</i> -Antibody Modification via Microbial Transglutaminase. <i>Bioconjugate Chemistry</i> , 2019, 30, 2452-2457.	1.8	23
12	Versatile Platform for the Synthesis of Orthogonally Cleavable Heteromultifunctional Cross-Linkers. <i>Bioconjugate Chemistry</i> , 2017, 28, 907-912.	1.8	20
13	Antibacterial isoamphiphathic oligomers highlight the importance of multimeric lipid aggregation for antibacterial potency. <i>Communications Biology</i> , 2018, 1, 220.	2.0	19
14	Effect of Composition on Antibacterial Activity of Sequence-Defined Cationic Oligothioetheramides. <i>ACS Infectious Diseases</i> , 2018, 4, 1257-1263.	1.8	19
15	Hydrophilic Sequence-Defined Cross-Linkers for Antibody-Drug Conjugates. <i>Bioconjugate Chemistry</i> , 2019, 30, 2982-2988.	1.8	15
16	Hijacking the Fusion Complex of Human Parainfluenza Virus as an Antiviral Strategy. <i>MBio</i> , 2020, 11, .	1.8	15
17	Responsive Antibody Conjugates Enable Quantitative Determination of Intracellular Bond Degradation Rate. <i>Cell Chemical Biology</i> , 2019, 26, 1643-1651.e4.	2.5	14
18	Antibody-Mediated Endocytosis of Polysialic Acid Enables Intracellular Delivery and Cytotoxicity of a Glycan-Directed Antibody-Drug Conjugate. <i>Cancer Research</i> , 2019, 79, 1810-1821.	0.4	14

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19	Synthesis and Solution-Phase Characterization of Sulfonated Oligothioetheramides. <i>Macromolecules</i> , 2017, 50, 8731-8738.	2.2	12
20	Intracellular Delivery via Noncharged Sequence-Defined Cell-Penetrating Oligomers. <i>Bioconjugate Chemistry</i> , 2018, 29, 2628-2635.	1.8	11
21	Sensitivity of Antibacterial Activity to Backbone Sequence in Constitutionally Isomeric OligoTEAs. <i>Macromolecular Bioscience</i> , 2018, 18, 1800241.	2.1	9
22	Biophysical Characterization of Cationic Antibacterial Oligothioetheramides. <i>Analytical Chemistry</i> , 2019, 91, 3118-3124.	3.2	9
23	Inhibiting Human Parainfluenza Virus Infection by Preactivating the Cell Entry Mechanism. <i>MBio</i> , 2019, 10, .	1.8	9
24	Inhibition of Measles Viral Fusion Is Enhanced by Targeting Multiple Domains of the Fusion Protein. <i>ACS Nano</i> , 2021, 15, 12794-12803.	7.3	9
25	Effect of backbone and end-group regioisomerism on thermomechanical properties of vanillin-based polyurethane networks. <i>Polymer Chemistry</i> , 2021, 12, 1526-1532.	1.9	8
26	Characterization of 14-Crown-4 Ethers for the Extraction of Lithium from Natural Brines: Synthesis, Solubility Measurements in Supercritical Carbon Dioxide, and Thermodynamic Modeling. <i>Industrial &amp; Engineering Chemistry Research</i> , 2021, 60, 7926-7934.	1.8	7
27	Predictive Platforms of Bond Cleavage and Drug Release Kinetics for Macromolecule-Drug Conjugates. <i>Annual Review of Chemical and Biomolecular Engineering</i> , 2021, 12, 241-261.	3.3	7
28	Decomplexation as a rate limitation in the thiol-Michael addition of <i>N</i> -acrylamides. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 6364-6377.	1.5	6
29	Design of a PEGylated Antimicrobial Prodrug with Species-Specific Activation. <i>Biomacromolecules</i> , 2021, 22, 984-992.	2.6	5
30	Mechanism of Action and Resistance Evasion of an Antimicrobial Oligomer against Multidrug-Resistant Gram-Negative Bacteria. <i>ACS Applied Bio Materials</i> , 2022, 5, 1159-1168.	2.3	5
31	Repurposing an In Vitro Measles Virus Dissemination Assay for Screening of Antiviral Compounds. <i>Viruses</i> , 2022, 14, 1186.	1.5	4
32	Design of protein-based molecular probes for intracellular bond cleavage. <i>Molecular Systems Design and Engineering</i> , 2020, 5, 385-391.	1.7	2
33	PEGylated Oligothioetheramide Prodrugs Activated by Host Serum Proteases. <i>ChemBioChem</i> , 2021, 22, 2697-2702.	1.3	2
34	Quantitative Determination of Intracellular Bond Cleavage. <i>Methods in Pharmacology and Toxicology</i> , 2021, , 305-330.	0.1	1