

# Daniel Lauster

## List of Publications by Year in descending order

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28  
papers

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citations

567281  
15  
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docs citations

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times ranked

1418  
citing authors

#	ARTICLE	IF	CITATIONS
1	Atomistic insight into the essential binding event of ACE2-derived peptides to the SARS-CoV-2 spike protein. <i>Biological Chemistry</i> , 2022, 403, 615-624.	2.5	2
2	Design and Functional Analysis of Heterobifunctional Multivalent Phage Capsid Inhibitors Blocking the Entry of Influenza Virus. <i>Bioconjugate Chemistry</i> , 2022, 33, 1269-1278.	3.6	1
3	Synthetic $\beta$ -Helical Peptides as Potential Inhibitors of the ACE2 SARS-CoV-2 Interaction. <i>ChemBioChem</i> , 2022, 23, .	2.6	6
4	Wrapping and Blocking of Influenza A Viruses by Sialylated 2D Nanoplatforms. <i>Advanced Materials Interfaces</i> , 2021, 8, 2100285.	3.7	17
5	Polysulfate hemmen durch elektrostatische Wechselwirkungen die SARS-CoV-2-Infektion**. <i>Angewandte Chemie</i> , 2021, 133, 16005-16014.	2.0	0
6	Polysulfates Block SARS-CoV-2 Uptake through Electrostatic Interactions**. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 15870-15878.	13.8	49
7	Evaluation of Multivalent Sialylated Polyglycerols for Resistance Induction in and Broad Antiviral Activity against Influenza A Viruses. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 12774-12789.	6.4	11
8	Inhibition of SARS-CoV-2 Replication by a Small Interfering RNA Targeting the Leader Sequence. <i>Viruses</i> , 2021, 13, 2030.	3.3	23
9	Quantification of Multivalent Interactions between Sialic Acid and Influenza A Virus Spike Proteins by Single-Molecule Force Spectroscopy. <i>Journal of the American Chemical Society</i> , 2020, 142, 12181-12192.	13.7	43
10	Adaptive Flexible Sialylated Nanogels as Highly Potent Influenza A Virus Inhibitors. <i>Angewandte Chemie</i> , 2020, 132, 12517-12522.	2.0	5
11	Phage capsid nanoparticles with defined ligand arrangement block influenza virus entry. <i>Nature Nanotechnology</i> , 2020, 15, 373-379.	31.5	96
12	Adaptive Flexible Sialylated Nanogels as Highly Potent Influenza A Virus Inhibitors. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 12417-12422.	13.8	36
13	Mobility-Based Quantification of Multivalent Virus-Receptor Interactions: New Insights Into Influenza A Virus Binding Mode. <i>Nano Letters</i> , 2019, 19, 1875-1882.	9.1	60
14	Force Spectroscopy Shows Dynamic Binding of Influenza Hemagglutinin and Neuraminidase to Sialic Acid. <i>Biophysical Journal</i> , 2019, 116, 1037-1048.	0.5	33
15	Hooking on Viral Glycoproteins with Single Molecule Force Spectroscopy to Study Single and Multiple Bond Formations. <i>Biophysical Journal</i> , 2019, 116, 428a.	0.5	0
16	The kinetochore module Okp1 $\text{CENP}^{\text{Q}}$ /Ame1 $\text{CENP}^{\text{U}}$ is a reader for N-terminal modifications on the centromeric histone Cse4 $\text{CENP}^{\text{A}}$ . <i>EMBO Journal</i> , 2019, 38, .	7.8	34
17	Interactions of Fullerene-Polyglycerol Sulfates at Viral and Cellular Interfaces. <i>Small</i> , 2018, 14, e1800189.	10.0	30
18	Sialyl-LacNAc-PNA <sup>TM</sup> DNA concatamers by rolling circle amplification as multivalent inhibitors for Influenza A virus particles. <i>ChemBioChem</i> , 2018, 20, 159-165.	2.6	15

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19	Exploring Rigid and Flexible Core Trivalent Sialosides for Influenza Virus Inhibition. <i>Chemistry - A European Journal</i> , 2018, 24, 19373-19385.	3.3	14
20	Functionalized Graphene as Extracellular Matrix Mimics: Toward Well-defined 2D Nanomaterials for Multivalent Virus Interactions. <i>Advanced Functional Materials</i> , 2017, 27, 1606477.	14.9	65
21	Multivalent Peptide-“Nanoparticle Conjugates for Influenza Virus Inhibition. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 5931-5936.	13.8	86
22	Linear polysialoside outperforms dendritic analogs for inhibition of influenza virus infection in vitro and in vivo. <i>Biomaterials</i> , 2017, 138, 22-34.	11.4	83
23	Multivalente Peptid-“Nanopartikel-Konjugate zur Hemmung des Influenzavirus. <i>Angewandte Chemie</i> , 2017, 129, 6025-6030.	2.0	8
24	Spatial Screening of Hemagglutinin on Influenza A Virus Particles: Sialyl-LacNAc Displays on DNA and PEG Scaffolds Reveal the Requirements for Bivalence Enhanced Interactions with Weak Monovalent Binders. <i>Journal of the American Chemical Society</i> , 2017, 139, 16389-16397.	13.7	70
25	Anti-Hemagglutinin Antibody Derived Lead Peptides for Inhibitors of Influenza Virus Binding. <i>PLoS ONE</i> , 2016, 11, e0159074.	2.5	25
26	Potential of acylated peptides to target the influenza A virus. <i>Beilstein Journal of Organic Chemistry</i> , 2015, 11, 589-595.	2.2	6
27	Potential of Proapoptotic Peptides to Induce the Formation of Giant Plasma Membrane Vesicles with Lipid Domains. <i>ChemBioChem</i> , 2015, 16, 1288-1292.	2.6	2
28	Autophagy restricts <i>Chlamydia trachomatis</i> growth in human macrophages via IFNG-inducible guanylate binding proteins. <i>Autophagy</i> , 2013, 9, 50-62.	9.1	108