Christiane Berger-Schaffitzel

List of Publications by Year in descending order

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CHRISTIANE

#	Article	IF	CITATIONS
1	The fatty acid site is coupled to functional motifs in the SARS-CoV-2 spike protein and modulates spike allosteric behaviour. Computational and Structural Biotechnology Journal, 2022, 20, 139-147.	4.1	19
2	Structural insights in cell-type specific evolution of intra-host diversity by SARS-CoV-2. Nature Communications, 2022, 13, 222.	12.8	23
3	Synthetic virions reveal fatty acid-coupled adaptive immunogenicity of SARS-CoV-2 spike glycoprotein. Nature Communications, 2022, 13, 868.	12.8	20
4	Structures of nonsense-mediated mRNA decay factors UPF3B and UPF3A in complex with UPF2 reveal molecular basis for competitive binding and for neurodevelopmental disorder-causing mutation. Nucleic Acids Research, 2022, 50, 5934-5947.	14.5	8
5	No-nonsense: insights into the functional interplay of nonsense-mediated mRNA decay factors. Biochemical Journal, 2022, 479, 973-993.	3.7	9
6	Pathogen-sugar interactions revealed by universal saturation transfer analysis. Science, 2022, 377, .	12.6	24
7	Highly efficient CRISPR-mediated large DNA docking and multiplexed prime editing using a single baculovirus. Nucleic Acids Research, 2022, 50, 7783-7799.	14.5	15
8	Molecular Simulations suggest Vitamins, Retinoids and Steroids as Ligands of the Free Fatty Acid Pocket of the SARSâ€CoVâ€2 Spike Protein**. Angewandte Chemie - International Edition, 2021, 60, 7098-7110.	13.8	77
9	Molecular Simulations suggest Vitamins, Retinoids and Steroids as Ligands of the Free Fatty Acid Pocket of the SARSâ€CoVâ€2 Spike Protein**. Angewandte Chemie, 2021, 133, 7174-7186.	2.0	6
10	Frontispiz: Molecular Simulations suggest Vitamins, Retinoids and Steroids as Ligands of the Free Fatty Acid Pocket of the SARS oVâ€2 Spike Protein. Angewandte Chemie, 2021, 133, .	2.0	7
11	Frontispiece: Molecular Simulations suggest Vitamins, Retinoids and Steroids as Ligands of the Free Fatty Acid Pocket of the SARSâ€CoVâ€2 Spike Protein. Angewandte Chemie - International Edition, 2021, 60, .	13.8	0
12	VLPâ€factoryâ,,¢ and ADDomer [©] : Selfâ€assembling Virusâ€Like Particle (VLP) Technologies for Multiple Protein and Peptide Epitope Display. Current Protocols, 2021, 1, e55.	2.9	9
13	Blasticidin S inhibits mammalian translation and enhances production of protein encoded by nonsense mRNA. Nucleic Acids Research, 2021, 49, 7665-7679.	14.5	13
14	Identification and Phenotypic Characterization of Hsp90 Phosphorylation Sites That Modulate Virulence Traits in the Major Human Fungal Pathogen Candida albicans. Frontiers in Cellular and Infection Microbiology, 2021, 11, 637836.	3.9	9
15	Structural biology in the fight against COVID-19. Nature Structural and Molecular Biology, 2021, 28, 2-7.	8.2	20
16	Production of Multi-subunit Membrane Protein Complexes. Methods in Molecular Biology, 2021, 2247, 3-16.	0.9	1
17	New insights into no-go, non-stop and nonsense-mediated mRNA decay complexes. Current Opinion in Structural Biology, 2020, 65, 110-118.	5.7	40
18	Free fatty acid binding pocket in the locked structure of SARS-CoV-2 spike protein. Science, 2020, 370, 725-730.	12.6	348

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19	The SARS-CoV-2 spike protein: balancing stability and infectivity. Cell Research, 2020, 30, 1059-1060.	12.0	82
20	High-Throughput Production of Influenza Virus-Like Particle (VLP) Array by Using VLP-factoryâ"¢, a MultiBac Baculoviral Genome Customized for Enveloped VLP Expression. Methods in Molecular Biology, 2019, 2025, 213-226.	0.9	7
21	Synthetic self-assembling ADDomer platform for highly efficient vaccination by genetically encoded multiepitope display. Science Advances, 2019, 5, eaaw2853.	10.3	29
22	Structure and Dynamics of the Central Lipid Pool and Proteins of the Bacterial Holo-Translocon. Biophysical Journal, 2019, 116, 1931-1940.	0.5	22
23	MultiBac: Baculovirus-Mediated Multigene DNA Cargo Delivery in Insect and Mammalian Cells. Viruses, 2019, 11, 198.	3.3	25
24	Cloning, expression, and purification of intact polyketide synthase modules. Methods in Enzymology, 2019, 617, 63-82.	1.0	3
25	New insights into the interplay between the translation machinery and nonsense-mediated mRNA decay factors. Biochemical Society Transactions, 2018, 46, 503-512.	3.4	38
26	Structure of a human cap-dependent 48S translation pre-initiation complex. Nucleic Acids Research, 2018, 46, 2678-2689.	14.5	76
27	Efficient production of a mature and functional gamma secretase protease. Scientific Reports, 2018, 8, 12834.	3.3	5
28	Multiprotein Complex Production in E. coli: The SecYEG-SecDFYajC-YidC Holotranslocon. Methods in Molecular Biology, 2017, 1586, 279-290.	0.9	2
29	Dual function of UPF3B in early and late translation termination. EMBO Journal, 2017, 36, 2968-2986.	7.8	89
30	Cryo-EM structure of Saccharomyces cerevisiae target of rapamycin complex 2. Nature Communications, 2017, 8, 1729.	12.8	46
31	A central cavity within the holo-translocon suggests a mechanism for membrane protein insertion. Scientific Reports, 2016, 6, 38399.	3.3	54
32	Membrane protein insertion and assembly by the bacterial holo-translocon SecYEG–SecDF–YajC–YidC. Biochemical Journal, 2016, 473, 3341-3354.	3.7	61
33	Cell-Free Synthesis of Macromolecular Complexes. Advances in Experimental Medicine and Biology, 2016, 896, 79-95.	1.6	0
34	ACEMBL Tool-Kits for High-Throughput Multigene Delivery and Expression in Prokaryotic and Eukaryotic Hosts. Advances in Experimental Medicine and Biology, 2016, 896, 27-42.	1.6	17
35	PABP enhances release factor recruitment and stop codon recognition during translation termination. Nucleic Acids Research, 2016, 44, 7766-7776.	14.5	99
36	MCM2-7 conformational changes in the presence of DNA. Cell Cycle, 2016, 15, 2391-2392.	2.6	0

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37	Molecular Basis of the Rapamycin Insensitivity of Target Of Rapamycin Complex 2. Molecular Cell, 2015, 58, 977-988.	9.7	120
38	Cytoplasmic TAF2–TAF8–TAF10 complex provides evidence for nuclear holo–TFIID assembly from preformed submodules. Nature Communications, 2015, 6, 6011.	12.8	77
39	ACEMBLing a Multiprotein Transmembrane Complex. Methods in Enzymology, 2015, 556, 23-49.	1.0	9
40	Advances and challenges of membrane–protein complex production. Current Opinion in Structural Biology, 2015, 32, 123-130.	5.7	32
41	Ribosome–SRP–FtsY cotranslational targeting complex in the closed state. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 3943-3948.	7.1	26
42	A network of SMG-8, SMG-9 and SMG-1 C-terminal insertion domain regulates UPF1 substrate recruitment and phosphorylation. Nucleic Acids Research, 2015, 43, 7600-7611.	14.5	51
43	Structural and functional analysis of the three MIF4G domains of nonsense-mediated decay factor UPF2. Nucleic Acids Research, 2014, 42, 2673-2686.	14.5	43
44	Membrane protein insertion and proton-motive-force-dependent secretion through the bacterial holo-translocon SecYEG–SecDF–YajC–YidC. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 4844-4849.	7.1	124
45	Continuous fluorescence-based measurement of redox-driven sodium ion translocation. Analytical Biochemistry, 2014, 459, 53-55.	2.4	2
46	The architecture of human general transcription factor TFIID core complex. Nature, 2013, 493, 699-702.	27.8	142
47	Robots, pipelines, polyproteins: Enabling multiprotein expression in prokaryotic and eukaryotic cells. Journal of Structural Biology, 2011, 175, 198-208.	2.8	92
48	Automated unrestricted multigene recombineering for multiprotein complex production. Nature Methods, 2009, 6, 447-450.	19.0	98
49	Multiprotein Expression Strategy for Structural Biology of Eukaryotic Complexes. Structure, 2007, 15, 275-279.	3.3	50
50	Protein complex expression by using multigene baculoviral vectors. Nature Methods, 2006, 3, 1021-1032.	19.0	330
51	Ribosome display: an in vitro method for selection and evolution of antibodies from libraries. Journal of Immunological Methods, 1999, 231, 119-135.	1.4	202