

Karen L Maxwell

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/2495937/karen-l-maxwell-publications-by-year.pdf>

Version: 2024-04-26

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

68

papers

4,588

citations

36

h-index

67

g-index

74

ext. papers

5,682

ext. citations

14.4

avg, IF

5.92

L-index

#	Paper	IF	Citations
68	A Filamentous Bacteriophage Protein Inhibits Type IV Pili To Prevent Superinfection of <i>Pseudomonas aeruginosa</i> .. <i>MBio</i> , 2022 , e0244121	7.8	0
67	Cyclic pyrimidines jump on the anti-phage bandwagon. <i>Cell</i> , 2021 , 184, 5691-5693	56.2	0
66	Structural and mechanistic insight into CRISPR-Cas9 inhibition by anti-CRISPR protein AcrIIC4.. <i>Journal of Molecular Biology</i> , 2021 , 434, 167420	6.5	0
65	Anti-CRISPR AcrIE2 Binds the Type I-E CRISPR-Cas Complex But Does Not Block DNA Binding. <i>Journal of Molecular Biology</i> , 2021 , 433, 166759	6.5	4
64	A phage-encoded anti-activator inhibits quorum sensing in <i>Pseudomonas aeruginosa</i> . <i>Molecular Cell</i> , 2021 , 81, 571-583.e6	17.6	30
63	HK97 gp74 Possesses an α -Helical Insertion in the β -Fold That Affects Its Metal Binding, Site Digestion, and Activities. <i>Journal of Bacteriology</i> , 2020 , 202,	3.5	3
62	Retrons: Complementing CRISPR in Phage Defense. <i>CRISPR Journal</i> , 2020 , 3, 226-227	2.5	3
61	Inhibition of CRISPR-Cas9 ribonucleoprotein complex assembly by anti-CRISPR AcrIIC2. <i>Nature Communications</i> , 2019 , 10, 2806	17.4	30
60	Extrachromosomal circular elements targeted by CRISPR-Cas in <i>Dehalococcoides mccartyi</i> are linked to mobilization of reductive dehalogenase genes. <i>ISME Journal</i> , 2019 , 13, 24-38	11.9	10
59	Anti-CRISPR AcrIIA5 Potently Inhibits All Cas9 Homologs Used for Genome Editing. <i>Cell Reports</i> , 2019 , 29, 1739-1746.e5	10.6	20
58	Meet the Anti-CRISPRs: Widespread Protein Inhibitors of CRISPR-Cas Systems. <i>CRISPR Journal</i> , 2019 , 2, 23-30	2.5	53
57	Phages Tune in to Host Cell Quorum Sensing. <i>Cell</i> , 2019 , 176, 7-8	56.2	7
56	The Diverse Impacts of Phage Morons on Bacterial Fitness and Virulence. <i>Advances in Virus Research</i> , 2019 , 103, 1-31	10.7	55
55	Anti-CRISPR: discovery, mechanism and function. <i>Nature Reviews Microbiology</i> , 2018 , 16, 12-17	22.2	200
54	Type VI secretion system baseplate. <i>Nature Microbiology</i> , 2018 , 3, 1330-1331	26.6	1
53	Potent Cas9 Inhibition in Bacterial and Human Cells by AcrIIC4 and AcrIIC5 Anti-CRISPR Proteins. <i>MBio</i> , 2018 , 9,	7.8	51
52	A chemical defence against phage infection. <i>Nature</i> , 2018 , 564, 283-286	50.4	78

51	A Unified Resource for Tracking Anti-CRISPR Names. <i>CRISPR Journal</i> , 2018 , 1, 304-305	2.5	50
50	Phage-Encoded Anti-CRISPR Defenses. <i>Annual Review of Genetics</i> , 2018 , 52, 445-464	14.5	77
49	Phage Morons Play an Important Role in Pseudomonas aeruginosa Phenotypes. <i>Journal of Bacteriology</i> , 2018 , 200,	3.5	29
48	Structure Reveals Mechanisms of Viral Suppressors that Intercept a CRISPR RNA-Guided Surveillance Complex. <i>Cell</i> , 2017 , 169, 47-57.e11	56.2	131
47	The Anti-CRISPR Story: A Battle for Survival. <i>Molecular Cell</i> , 2017 , 68, 8-14	17.6	50
46	A Broad-Spectrum Inhibitor of CRISPR-Cas9. <i>Cell</i> , 2017 , 170, 1224-1233.e15	56.2	145
45	Disabling a Type I-E CRISPR-Cas Nuclease with a Bacteriophage-Encoded Anti-CRISPR Protein. <i>MBio</i> , 2017 , 8,	7.8	42
44	Baseplate assembly of phage Mu: Defining the conserved core components of contractile-tailed phages and related bacterial systems. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016 , 113, 10174-9	11.5	29
43	Inactivation of CRISPR-Cas systems by anti-CRISPR proteins in diverse bacterial species. <i>Nature Microbiology</i> , 2016 , 1, 16085	26.6	203
42	The solution structure of an anti-CRISPR protein. <i>Nature Communications</i> , 2016 , 7, 13134	17.4	41
41	Prophages mediate defense against phage infection through diverse mechanisms. <i>ISME Journal</i> , 2016 , 10, 2854-2866	11.9	176
40	Phages Fight Back: Inactivation of the CRISPR-Cas Bacterial Immune System by Anti-CRISPR Proteins. <i>PLoS Pathogens</i> , 2016 , 12, e1005282	7.6	39
39	Naturally Occurring Off-Switches for CRISPR-Cas9. <i>Cell</i> , 2016 , 167, 1829-1838.e9	56.2	260
38	Multiple mechanisms for CRISPR-Cas inhibition by anti-CRISPR proteins. <i>Nature</i> , 2015 , 526, 136-9	50.4	225
37	The phage tail tape measure protein, an inner membrane protein and a periplasmic chaperone play connected roles in the genome injection process of E. coli phage HK97. <i>Molecular Microbiology</i> , 2015 , 96, 437-47	4.1	58
36	A new group of phage anti-CRISPR genes inhibits the type I-E CRISPR-Cas system of Pseudomonas aeruginosa. <i>MBio</i> , 2014 , 5, e00896	7.8	180
35	HNH proteins are a widespread component of phage DNA packaging machines. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 6022-7	11.5	59
34	A shifty chaperone for phage tail assembly. <i>Journal of Molecular Biology</i> , 2014 , 426, 1001-3	6.5	4

33	Efficacy of bacteriophage treatment on <i>Pseudomonas aeruginosa</i> biofilms. <i>Journal of Endodontics</i> , 2013 , 39, 364-9	4.7	30
32	Rapid detection of <i>E. coli</i> bacteria using potassium-sensitive FETs in CMOS. <i>IEEE Transactions on Biomedical Circuits and Systems</i> , 2013 , 7, 621-30	5.1	27
31	Bacteriophage genes that inactivate the CRISPR/Cas bacterial immune system. <i>Nature</i> , 2013 , 493, 429-33	10.4	495
30	A conserved spiral structure for highly diverged phage tail assembly chaperones. <i>Journal of Molecular Biology</i> , 2013 , 425, 2436-49	6.5	12
29	Tail tip proteins related to bacteriophage λ gpL coordinate an iron-sulfur cluster. <i>Journal of Molecular Biology</i> , 2013 , 425, 2450-62	6.5	17
28	The solution structures of two prophage homologues of the bacteriophage λ Ea8.5 protein reveal a newly discovered hybrid homeodomain/zinc-finger fold. <i>Biochemistry</i> , 2013 , 52, 3612-4	3.2	8
27	Structural and functional studies of gpX of <i>Escherichia coli</i> phage P2 reveal a widespread role for LysM domains in the baseplates of contractile-tailed phages. <i>Journal of Bacteriology</i> , 2013 , 195, 5461-8	3.5	14
26	The bacteriophage HK97 gp15 moron element encodes a novel superinfection exclusion protein. <i>Journal of Bacteriology</i> , 2012 , 194, 5012-9	3.5	71
25	The protein gp74 from the bacteriophage HK97 functions as a HNH endonuclease. <i>Protein Science</i> , 2012 , 21, 809-18	6.3	23
24	Long noncontractile tail machines of bacteriophages. <i>Advances in Experimental Medicine and Biology</i> , 2012 , 726, 115-42	3.6	78
23	The moron comes of age. <i>Bacteriophage</i> , 2012 , 2, 225-228		39
22	Structural and biochemical characterization of phage λ FI protein (gpFI) reveals a novel mechanism of DNA packaging chaperone activity. <i>Journal of Biological Chemistry</i> , 2012 , 287, 32085-95	5.4	5
21	Assembly mechanism is the key determinant of the dosage sensitivity of a phage structural protein. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011 , 108, 10168-73	11.5	11
20	Phages have adapted the same protein fold to fulfill multiple functions in virion assembly. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010 , 107, 14384-9	11.5	29
19	The crystal structure of bacteriophage HK97 gp6: defining a large family of head-tail connector proteins. <i>Journal of Molecular Biology</i> , 2010 , 395, 754-68	6.5	49
18	The solution structure of the C-terminal Ig-like domain of the bacteriophage λ tail tube protein. <i>Journal of Molecular Biology</i> , 2010 , 403, 468-79	6.5	39
17	Viral proteomics. <i>Microbiology and Molecular Biology Reviews</i> , 2007 , 71, 398-411	13.2	103
16	Viral Proteomics. <i>Microbiology and Molecular Biology Reviews</i> , 2007 , 71, 549-549	13.2	78

15	The NMR structure of the gpU tail-terminator protein from bacteriophage lambda: identification of sites contributing to Mg(II)-mediated oligomerization and biological function. <i>Journal of Molecular Biology</i> , 2007 , 365, 175-86	6.5	27
14	Immunoglobulin-like domains on bacteriophage: weapons of modest damage?. <i>Current Opinion in Microbiology</i> , 2007 , 10, 382-7	7.9	69
13	Ig-like domains on bacteriophages: a tale of promiscuity and deceit. <i>Journal of Molecular Biology</i> , 2006 , 359, 496-507	6.5	136
12	Crystal structure of bacteriophage lambda cII and its DNA complex. <i>Molecular Cell</i> , 2005 , 19, 259-69	17.6	34
11	Protein folding: defining a "standard" set of experimental conditions and a preliminary kinetic data set of two-state proteins. <i>Protein Science</i> , 2005 , 14, 602-16	6.3	181
10	Refolding out of guanidine hydrochloride is an effective approach for high-throughput structural studies of small proteins. <i>Protein Science</i> , 2003 , 12, 2073-80	6.3	37
9	The solution structure of the bacteriophage lambda head-tail joining protein, gpFII. <i>Journal of Molecular Biology</i> , 2002 , 318, 1395-404	6.5	31
8	Protein folding kinetics beyond the phi value: using multiple amino acid substitutions to investigate the structure of the SH3 domain folding transition state. <i>Journal of Molecular Biology</i> , 2002 , 320, 389-402	6.5	71
7	The solution structure of bacteriophage lambda protein W, a small morphogenetic protein possessing a novel fold. <i>Journal of Molecular Biology</i> , 2001 , 308, 9-14	6.5	36
6	Structural proteomics of an archaeon. <i>Nature Structural Biology</i> , 2000 , 7, 903-9		247
5	Thermodynamic and functional characterization of protein W from bacteriophage lambda. The three C-terminal residues are critical for activity. <i>Journal of Biological Chemistry</i> , 2000 , 275, 18879-86	5.4	17
4	A simple in vivo assay for increased protein solubility. <i>Protein Science</i> , 1999 , 8, 1908-11	6.3	140
3	Mutagenesis of a buried polar interaction in an SH3 domain: sequence conservation provides the best prediction of stability effects. <i>Biochemistry</i> , 1998 , 37, 16172-82	3.2	85
2	One Anti-CRISPR to Rule Them All: Potent Inhibition of Cas9 Homologs Used for Genome Editing. <i>SSRN Electronic Journal</i> ,	1	1
1	Potent Cas9 inhibition in bacterial and human cells by new anti-CRISPR protein families		1