

Peter Friedhoff

List of Publications by Year in descending order

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59
papers

2,256
citations

201385

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docs citations

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times ranked

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citing authors

#	ARTICLE	IF	CITATIONS
1	Cryogenic electron microscopy structures reveal how ATP and DNA binding in MutS coordinates sequential steps of DNA mismatch repair. <i>Nature Structural and Molecular Biology</i> , 2022, 29, 59-66.	3.6	12
2	Reactive Acrylamide-Modified DNA Traps for Accurate Cross-Linking with Cysteine Residues in DNA-Protein Complexes Using Mismatch Repair Protein MutS as a Model. <i>Molecules</i> , 2022, 27, 2438.	1.7	1
3	The selection process of licensing a DNA mismatch for repair. <i>Nature Structural and Molecular Biology</i> , 2021, 28, 373-381.	3.6	22
4	Probing the DNA-binding center of the MutL protein from the Escherichia coli mismatch repair system via crosslinking and Förster resonance energy transfer. <i>Biochimie</i> , 2020, 171-172, 43-54.	1.3	9
5	The unstructured linker arms of MutL enable GATC site incision beyond roadblocks during initiation of DNA mismatch repair. <i>Nucleic Acids Research</i> , 2019, 47, 11667-11680.	6.5	26
6	Combinatorial recognition of clustered RNA elements by the multidomain RNA-binding protein IMP3. <i>Nature Communications</i> , 2019, 10, 2266.	5.8	53
7	Mechanism and Regulation of Co-transcriptional mRNP Assembly and Nuclear mRNA Export. <i>Advances in Experimental Medicine and Biology</i> , 2019, 1203, 1-31.	0.8	27
8	Use of Single-Cysteine Variants for Trapping Transient States in DNA Mismatch Repair. <i>Methods in Enzymology</i> , 2017, 592, 77-101.	0.4	4
9	Dual daughter strand incision is processive and increases the efficiency of DNA mismatch repair. <i>Nucleic Acids Research</i> , 2016, 44, 6770-6786.	6.5	18
10	Protein-protein interactions in DNA mismatch repair. <i>DNA Repair</i> , 2016, 38, 50-57.	1.3	38
11	Editorial: Alfred Pingoud (1945-2015). <i>Nucleic Acids Research</i> , 2015, 43, 7661-7663.	6.5	0
12	Chromatographic isolation of the functionally active MutS protein covalently linked to deoxyribonucleic acid. <i>Journal of Chromatography A</i> , 2015, 1389, 19-27.	1.8	12
13	MutS/MutL crystal structure reveals that the MutS sliding clamp loads MutL onto DNA. <i>ELife</i> , 2015, 4, e06744.	2.8	91
14	Is Thymidine Glycol Containing DNA a Substrate of E. coli DNA Mismatch Repair System?. <i>PLoS ONE</i> , 2014, 9, e104963.	1.1	4
15	Using stable MutS dimers and tetramers to quantitatively analyze DNA mismatch recognition and sliding clamp formation. <i>Nucleic Acids Research</i> , 2013, 41, 8166-8181.	6.5	36
16	Site- and strand-specific nicking of DNA by fusion proteins derived from MutH and I-SceI or TALE repeats. <i>Nucleic Acids Research</i> , 2013, 41, e83-e83.	6.5	31
17	Single-molecule multiparameter fluorescence spectroscopy reveals directional MutS binding to mismatched bases in DNA. <i>Nucleic Acids Research</i> , 2012, 40, 5448-5464.	6.5	44
18	Chemical Rescue of Active Site Mutants of <i>S. pneumoniae</i> Surface Endonuclease EndA and Other Nucleases of the HNH Family by Imidazole. <i>ChemBioChem</i> , 2012, 13, 713-721.	1.3	15

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19	Covalently trapping MutS on DNA to study DNA mismatch recognition and signaling. <i>Molecular BioSystems</i> , 2012, 8, 1861.	2.9	10
20	Generation of DNA nanocircles containing mismatched bases. <i>BioTechniques</i> , 2011, 51, 259-265.	0.8	5
21	Changes in cytosolic Mg ²⁺ levels can regulate the activity of the plasma membrane H ⁺ -ATPase in maize. <i>Biochemical Journal</i> , 2011, 435, 93-101.	1.7	26
22	Chemical Trapping of the Dynamic MutS-MutL Complex Formed in DNA Mismatch Repair in <i>Escherichia coli</i> . <i>Journal of Biological Chemistry</i> , 2011, 286, 17326-17337.	1.6	45
23	Native mass spectrometry provides direct evidence for DNA mismatch-induced regulation of asymmetric nucleotide binding in mismatch repair protein MutS. <i>Nucleic Acids Research</i> , 2011, 39, 8052-8064.	6.5	30
24	Chemical trapping of the dynamic MutS-MutL complex formed in DNA mismatch repair in <i>Escherichia coli</i> . <i>Journal of Biological Chemistry</i> , 2011, 286, 22706.	1.6	0
25	Crosslinking of (Cytosine-5)-DNA Methyltransferase SsoII and its Complexes with Specific DNA Duplexes Provides an Insight into Their Structures. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 2011, 30, 632-650.	0.4	8
26	Identification of Lynch syndrome mutations in the MLH1-PMS2 interface that disturb dimerization and mismatch repair. <i>Human Mutation</i> , 2010, 31, 975-982.	1.1	52
27	The C-Terminal Domain of the MutL Homolog from <i>Neisseria gonorrhoeae</i> Forms an Inverted Homodimer. <i>PLoS ONE</i> , 2010, 5, e13726.	1.1	29
28	Maintaining a sense of direction during long-range communication on DNA. <i>Biochemical Society Transactions</i> , 2010, 38, 404-409.	1.6	24
29	Structure of the Endonuclease Domain of MutL: Unlicensed to Cut. <i>Molecular Cell</i> , 2010, 39, 145-151.	4.5	122
30	Physical and functional interactions between <i>Escherichia coli</i> MutL and the Vsr repair endonuclease. <i>Nucleic Acids Research</i> , 2009, 37, 4453-4463.	6.5	24
31	On the Divalent Metal Ion Dependence of DNA Cleavage by Restriction Endonucleases of the EcoRI Family. <i>Journal of Molecular Biology</i> , 2009, 393, 140-160.	2.0	59
32	The C-terminal domain is sufficient for endonuclease activity of <i>Neisseria gonorrhoeae</i> MutL. <i>Biochemical Journal</i> , 2009, 423, 265-277.	1.7	46
33	The PMS2 Subunit of Human MutL ⁺ Contains a Metal Ion Binding Domain of the Iron-Dependent Repressor Protein Family. <i>Journal of Molecular Biology</i> , 2008, 382, 610-627.	2.0	55
34	Bioinformatics-Guided Experimental Characterization of Mismatch-Repair Enzymes and Their Relatives. <i>Nucleic Acids and Molecular Biology</i> , 2008, , 221-241.	0.2	0
35	Engineering Site-Specific Endonucleases. , 2007, 352, 111-124.		3
36	Mutations in the MutS ⁺ interaction interface of MLH1 can abolish DNA mismatch repair. <i>Nucleic Acids Research</i> , 2006, 34, 6574-6586.	6.5	61

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37	Structural and functional analysis of the MutS C-terminal tetramerization domain. <i>Nucleic Acids Research</i> , 2006, 34, 5270-5279.	6.5	32
38	Identifying an interaction site between MutH and the C-terminal domain of MutL by crosslinking, affinity purification, chemical coding and mass spectrometry. <i>Nucleic Acids Research</i> , 2006, 34, 3169-3180.	6.5	41
39	Mapping protein-protein interactions by bioinformatics and cross-linking. <i>Analytical and Bioanalytical Chemistry</i> , 2005, 381, 78-80.	1.9	21
40	Analysis of the Quaternary Structure of the MutL C-terminal Domain. <i>Journal of Molecular Biology</i> , 2005, 351, 895-909.	2.0	53
41	Mapping Protein-Protein Interactions between MutL and MutH by Cross-linking. <i>Journal of Biological Chemistry</i> , 2004, 279, 49338-49345.	1.6	49
42	Tyr212: A Key Residue Involved in Strand Discrimination by the DNA Mismatch Repair Endonuclease MutH. <i>Journal of Molecular Biology</i> , 2003, 325, 285-297.	2.0	27
43	Application of the C4'-Alkylated Deoxyribose Primer System (CAPS) in Allele-Specific Real-Time PCR for Increased Selectivity in Discrimination of Single Nucleotide Sequence Variants. <i>Biological Chemistry</i> , 2003, 384, 1533-41.	1.2	14
44	Site-specific protein modification to identify the MutL interface of MutH. <i>Nucleic Acids Research</i> , 2003, 31, 819-825.	6.5	18
45	Evolutionary Relationship between Different Subgroups of Restriction Endonucleases. <i>Journal of Biological Chemistry</i> , 2002, 277, 14306-14314.	1.6	53
46	An Efficient Method for the Preparation of Long Heteroduplex DNA as Substrate for Mismatch Repair by the Escherichia coli MutHLS System. <i>Biological Chemistry</i> , 2002, 383, 1459-1462.	1.2	17
47	Haemophilus influenzae and Vibrio cholerae genes for mutH are able to fully complement a mutH defect in Escherichia coli. <i>FEMS Microbiology Letters</i> , 2002, 208, 123-128.	0.7	15
48	Sau3AI, a Monomeric Type II Restriction Endonuclease That Dimerizes on the DNA and Thereby Induces DNA Loops. <i>Journal of Biological Chemistry</i> , 2001, 276, 23581-23588.	1.6	55
49	Microtiter-Plate Assay and Related Assays for Nonspecific Endonucleases. , 2001, 160, 037-048.		2
50	Structure of tau protein and assembly into paired helical filaments. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2000, 1502, 122-132.	1.8	130
51	A similar active site for non-specific and specific endonucleases. <i>Nature Structural Biology</i> , 1999, 6, 112-113.	9.7	59
52	Cleavage experiments with deoxythymidine 3'-5'-bis-(p-nitrophenyl phosphate) suggest that the homing endonuclease I-PpoI follows the same mechanism of phosphodiester bond hydrolysis as the non-specific Serratia nuclease 1. <i>FEBS Letters</i> , 1999, 443, 209-214.	1.3	36
53	Application of Oligonucleoside Methylphosphonates in the Studies on Phosphodiester Hydrolysis by Serratia Endonuclease 1. <i>Nucleosides & Nucleotides</i> , 1999, 18, 1945-1960.	0.5	4
54	Rapid Assembly of Alzheimer-like Paired Helical Filaments from Microtubule-Associated Protein Tau Monitored by Fluorescence in Solution. <i>Biochemistry</i> , 1998, 37, 10223-10230.	1.2	378

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55	Analysis of the reaction mechanism of the non-specific endonuclease of <i>Serratia marcescens</i> using an artificial minimal substrate. <i>FEBS Letters</i> , 1996, 397, 343-346.	1.3	27
56	Kinetic Analysis of the Cleavage of Natural and Synthetic Substrates by the <i>Serratia</i> Nuclease. <i>FEBS Journal</i> , 1996, 241, 572-580.	0.2	47
57	A Quantitative Microtiter Plate Nuclease Assay Based on Ethidium/DNA Fluorescence. <i>Analytical Biochemistry</i> , 1996, 240, 283-288.	1.1	11
58	Sequence preferences in cleavage of dsDNA and ssDNA by the extracellular <i>Serratia marcescens</i> endonuclease. <i>Biochemistry</i> , 1995, 34, 11979-11988.	1.2	48
59	Identification of catalytically relevant amino acids of the extracellular <i>Serratia marcescens</i> endonuclease by alignment-guided mutagenesis. <i>Nucleic Acids Research</i> , 1994, 22, 3280-3287.	6.5	72