

Guo-Min Li

List of Publications by Year in descending order

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86
papers

6,470
citations

94433

37
h-index

66911

78
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88
all docs

88
docs citations

88
times ranked

7969
citing authors

#	ARTICLE	IF	CITATIONS
1	Mechanisms and functions of DNA mismatch repair. <i>Cell Research</i> , 2008, 18, 85-98.	12.0	1,081
2	Hypermutability and mismatch repair deficiency in RER+ tumor cells. <i>Cell</i> , 1993, 75, 1227-1236.	28.9	1,081
3	The Histone Mark H3K36me3 Regulates Human DNA Mismatch Repair through Its Interaction with MutS \pm . <i>Cell</i> , 2013, 153, 590-600.	28.9	504
4	ARID1A deficiency promotes mutability and potentiates therapeutic antitumor immunity unleashed by immune checkpoint blockade. <i>Nature Medicine</i> , 2018, 24, 556-562.	30.7	372
5	Reconstitution of 5 $\hat{\text{A}}^{\text{€}}$ -Directed Human Mismatch Repair in a Purified System. <i>Cell</i> , 2005, 122, 693-705.	28.9	316
6	Mismatch Repair Genes Mlh1 and Mlh3 Modify CAG Instability in Huntington's Disease Mice: Genome-Wide and Candidate Approaches. <i>PLoS Genetics</i> , 2013, 9, e1003930.	3.5	175
7	DNA Sensing in Mismatch Repair-Deficient Tumor Cells Is Essential for Anti-tumor Immunity. <i>Cancer Cell</i> , 2021, 39, 96-108.e6.	16.8	153
8	Mismatch Repair Processing of Carcinogen-DNA Adducts Triggers Apoptosis. <i>Molecular and Cellular Biology</i> , 1999, 19, 8292-8301.	2.3	115
9	Evidence for Involvement of HMGB1 Protein in Human DNA Mismatch Repair. <i>Journal of Biological Chemistry</i> , 2004, 279, 20935-20940.	3.4	112
10	HDAC6 Deacetylates and Ubiquitinates MSH2 to Maintain Proper Levels of MutS \pm . <i>Molecular Cell</i> , 2014, 55, 31-46.	9.7	112
11	MLH1 Deficiency-Triggered DNA Hyperexcision by Exonuclease 1 Activates the cGAS-STING Pathway. <i>Cancer Cell</i> , 2021, 39, 109-121.e5.	16.8	108
12	Identification and characterization of OGG1 mutations in patients with Alzheimer's disease. <i>Nucleic Acids Research</i> , 2007, 35, 2759-2766.	14.5	105
13	Altered 8-oxoguanine glycosylase in mild cognitive impairment and late-stage Alzheimer's disease brain. <i>Free Radical Biology and Medicine</i> , 2008, 45, 813-819.	2.9	99
14	Specific Binding of Human MSH2 $\hat{\text{A}}$ -MSH6 Mismatch-Repair Protein Heterodimers to DNA Incorporating Thymine- or Uracil-containing UV Light Photoproducts Opposite Mismatched Bases. <i>Journal of Biological Chemistry</i> , 1999, 274, 16894-16900.	3.4	93
15	Prereplicative repair of oxidized bases in the human genome is mediated by NEIL1 DNA glycosylase together with replication proteins. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, E3090-9.	7.1	90
16	Cancer-driving H3G34V/R/D mutations block H3K36 methylation and H3K36me3 $\hat{\text{A}}$ MutS \pm interaction. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, 9598-9603.	7.1	87
17	Genetic and epigenetic modification of mismatch repair genes hMSH2 and hMLH1 in sporadic breast cancer with microsatellite instability. <i>Oncogene</i> , 2002, 21, 5696-5703.	5.9	80
18	Partial Reconstitution of Human DNA Mismatch Repair In Vitro: Characterization of the Role of Human Replication Protein A. <i>Molecular and Cellular Biology</i> , 2002, 22, 2037-2046.	2.3	75

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19	Effect of Carcinogenic Acrolein on DNA Repair and Mutagenic Susceptibility. <i>Journal of Biological Chemistry</i> , 2012, 287, 12379-12386.	3.4	75
20	Targeted DNA damage at individual telomeres disrupts their integrity and triggers cell death. <i>Nucleic Acids Research</i> , 2015, 43, 6334-6347.	14.5	68
21	Truncating mutation in the autophagy gene UVRAG confers oncogenic properties and chemosensitivity in colorectal cancers. <i>Nature Communications</i> , 2015, 6, 7839.	12.8	67
22	DNA mismatch repair and cancer. <i>Frontiers in Bioscience - Landmark</i> , 2003, 8, d997-1017.	3.0	66
23	Phosphorylation of PCNA by EGFR inhibits mismatch repair and promotes misincorporation during DNA synthesis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 5667-5672.	7.1	60
24	Evidence That Nucleosomes Inhibit Mismatch Repair in Eukaryotic Cells. <i>Journal of Biological Chemistry</i> , 2009, 284, 33056-33061.	3.4	59
25	Human MutS α Specifically Binds to DNA Containing Aminofluorene and Acetylaminofluorene Adducts. <i>Journal of Biological Chemistry</i> , 1996, 271, 24084-24088.	3.4	54
26	hMRE11 deficiency leads to microsatellite instability and defective DNA mismatch repair. <i>EMBO Reports</i> , 2005, 6, 438-444.	4.5	49
27	Folate deficiency, mismatch repair-dependent apoptosis, and human disease. <i>Journal of Nutritional Biochemistry</i> , 2003, 14, 568-575.	4.2	47
28	Differential Requirement for Proliferating Cell Nuclear Antigen in 5' and 3' Nick-directed Excision in Human Mismatch Repair. <i>Journal of Biological Chemistry</i> , 2004, 279, 16912-16917.	3.4	47
29	Roles of mismatch repair proteins hMSH2 and hMLH1 in the development of sporadic breast cancer. <i>Cancer Letters</i> , 2005, 223, 143-150.	7.2	47
30	Distinct Nucleotide Binding/Hydrolysis Properties and Molar Ratio of MutS α and MutS β Determine Their Differential Mismatch Binding Activities. <i>Journal of Biological Chemistry</i> , 2009, 284, 11557-11562.	3.4	47
31	Incision-dependent and error-free repair of (CAG) $_n$ /(CTG) $_n$ hairpins in human cell extracts. <i>Nature Structural and Molecular Biology</i> , 2009, 16, 869-875.	8.2	47
32	Mismatch Recognition Protein MutS β Does Not Hijack (CAG) Hairpin Repair in Vitro. <i>Journal of Biological Chemistry</i> , 2009, 284, 20452-20456.	3.4	45
33	MutS β promotes trinucleotide repeat expansion by recruiting DNA polymerase β to nascent (CAG) $_n$ or (CTG) $_n$ hairpins for error-prone DNA synthesis. <i>Cell Research</i> , 2016, 26, 775-786.	12.0	43
34	Mismatch repair deficiency in hematological malignancies with microsatellite instability. <i>Oncogene</i> , 2002, 21, 5758-5764.	5.9	41
35	Bi-directional Processing of DNA Loops by Mismatch Repair-dependent and -independent Pathways in Human Cells. <i>Journal of Biological Chemistry</i> , 2003, 278, 3891-3896.	3.4	41
36	The C-terminal Domain (CTD) of Human DNA Glycosylase NEIL1 Is Required for Forming BERosome Repair Complex with DNA Replication Proteins at the Replicating Genome. <i>Journal of Biological Chemistry</i> , 2015, 290, 20919-20933.	3.4	41

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37	Ubiquitin-specific Peptidase 10 (USP10) Deubiquitinates and Stabilizes MutS Homolog 2 (MSH2) to Regulate Cellular Sensitivity to DNA Damage. <i>Journal of Biological Chemistry</i> , 2016, 291, 10783-10791.	3.4	41
38	Molecular Cooperation between the Werner Syndrome Protein and Replication Protein A in Relation to Replication Fork Blockage. <i>Journal of Biological Chemistry</i> , 2011, 286, 3497-3508.	3.4	39
39	Modulation of microRNA processing by mismatch repair protein MutL β . <i>Cell Research</i> , 2012, 22, 973-985.	12.0	38
40	Involvement of DNA mismatch repair in folate deficiency-induced apoptosis β . <i>Journal of Nutritional Biochemistry</i> , 2002, 13, 355-363.	4.2	36
41	Decoding the Histone Code: Role of H3K36me3 in Mismatch Repair and Implications for Cancer Susceptibility and Therapy. <i>Cancer Research</i> , 2013, 73, 6379-6383.	0.9	36
42	Regulation of Replication Protein A Functions in DNA Mismatch Repair by Phosphorylation. <i>Journal of Biological Chemistry</i> , 2006, 281, 21607-21616.	3.4	34
43	New insights and challenges in mismatch repair: Getting over the chromatin hurdle. <i>DNA Repair</i> , 2014, 19, 48-54.	2.8	33
44	Arsenic Inhibits DNA Mismatch Repair by Promoting EGFR Expression and PCNA Phosphorylation. <i>Journal of Biological Chemistry</i> , 2015, 290, 14536-14541.	3.4	33
45	Okazaki fragment maturation involves β -segment error editing by the mammalian β FEN1/MutS β functional complex. <i>EMBO Journal</i> , 2015, 34, 1829-1843.	7.8	28
46	HDAC6 regulates DNA damage response via deacetylating MLH1. <i>Journal of Biological Chemistry</i> , 2019, 294, 5813-5826.	3.4	28
47	MutS β abundance and Msh3 ATP hydrolysis activity are important drivers of CTG β CAG repeat expansions. <i>Nucleic Acids Research</i> , 2017, 45, 10068-10078.	14.5	27
48	Mismatch-bound human MutS β -MutL complex triggers DNA incisions and activates mismatch repair. <i>Cell Research</i> , 2021, 31, 542-553.	12.0	26
49	DNA instability in replicating Huntington's disease lymphoblasts. <i>BMC Medical Genetics</i> , 2009, 10, 11.	2.1	24
50	DNA mismatch repair preferentially safeguards actively transcribed genes. <i>DNA Repair</i> , 2018, 71, 82-86.	2.8	24
51	Regulation of mismatch repair by histone code and posttranslational modifications in eukaryotic cells. <i>DNA Repair</i> , 2016, 38, 68-74.	2.8	22
52	In vitro repair of DNA hairpins containing various numbers of CAG/CTG trinucleotide repeats. <i>DNA Repair</i> , 2012, 11, 201-209.	2.8	20
53	A human MUTYH variant linking colonic polyposis to redox degradation of the [4Fe4S] β cluster. <i>Nature Chemistry</i> , 2018, 10, 873-880.	13.6	20
54	Phosphorylation of proliferating cell nuclear antigen promotes cancer progression by activating the ATM/Akt/GSK3 β /Snail signaling pathway. <i>Journal of Biological Chemistry</i> , 2019, 294, 7037-7045.	3.4	20

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55	MIF is a 3' flap nuclease that facilitates DNA replication and promotes tumor growth. <i>Nature Communications</i> , 2021, 12, 2954.	12.8	20
56	Nick-dependent and -independent Processing of Large DNA Loops in Human Cells. <i>Journal of Biological Chemistry</i> , 2003, 278, 50803-50809.	3.4	19
57	HDAC3 deacetylates the DNA mismatch repair factor MutS β to stimulate triplet repeat expansions. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 23597-23605.	7.1	19
58	Coordinated Processing of 3' Slipped (CAG) $_n$ /(CTG) $_n$ Hairpins by DNA Polymerases β and γ Preferentially Induces Repeat Expansions. <i>Journal of Biological Chemistry</i> , 2013, 288, 15015-15022.	3.4	18
59	The Werner Syndrome Protein Promotes CAG/CTG Repeat Stability by Resolving Large (CAG) / (CTG) Hairpins. <i>Journal of Biological Chemistry</i> , 2012, 287, 30151-30156.	3.4	17
60	Trinucleotide repeat expansions catalyzed by human cell-free extracts. <i>Cell Research</i> , 2013, 23, 565-572.	12.0	16
61	Identification of novel genetic variants predisposing to familial oral squamous cell carcinomas. <i>Cell Discovery</i> , 2019, 5, 57.	6.7	16
62	In vitro and in vivo modulations of benzo[c]phenanthrene-DNA adducts by DNA mismatch repair system. <i>Nucleic Acids Research</i> , 2003, 31, 6428-6434.	14.5	15
63	DNA mismatch repair in the context of chromatin. <i>Cell and Bioscience</i> , 2020, 10, 10.	4.8	15
64	OTUB1 stabilizes mismatch repair protein MSH2 by blocking ubiquitination. <i>Journal of Biological Chemistry</i> , 2021, 296, 100466.	3.4	15
65	The Role of XPG in Processing (CAG) $_n$ /(CTG) $_n$ DNA Hairpins. <i>Cell and Bioscience</i> , 2011, 1, 11.	4.8	14
66	Increased transversions in a novel mutator colon cancer cell line. <i>Oncogene</i> , 1998, 16, 1125-1130.	5.9	13
67	DNA mismatch repair in the chromatin context: Mechanisms and therapeutic potential. <i>DNA Repair</i> , 2020, 93, 102918.	2.8	10
68	Identification of Regulatory Factor X as a Novel Mismatch Repair Stimulatory Factor. <i>Journal of Biological Chemistry</i> , 2008, 283, 12730-12735.	3.4	9
69	The hMSH2(M688R) Lynch syndrome mutation may function as a dominant negative. <i>Carcinogenesis</i> , 2012, 33, 1647-1654.	2.8	8
70	A special issue on DNA damage response and genome stability. <i>Cell and Bioscience</i> , 2012, 2, 4.	4.8	8
71	NBS1-CtIP-mediated DNA end resection suppresses cGAS binding to micronuclei. <i>Nucleic Acids Research</i> , 2022, 50, 2681-2699.	14.5	8
72	DNA-PKcs-dependent phosphorylation of RECQL4 promotes NHEJ by stabilizing the NHEJ machinery at DNA double-strand breaks. <i>Nucleic Acids Research</i> , 2022, 50, 5635-5651.	14.5	8

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73	Novel molecular insights into the mechanism of GO removal by MutM. Cell Research, 2010, 20, 116-118.	12.0	7
74	DNA polymerase δ promotes CAG \leftrightarrow CTG repeat expansions in Huntington's disease via insertion sequences of its catalytic domain. Journal of Biological Chemistry, 2021, 297, 101144.	3.4	7
75	A special issue on DNA damage responses and genome maintenance. Cell Research, 2008, 18, 1-2.	12.0	6
76	DNA mismatch repair in trinucleotide repeat instability. Science China Life Sciences, 2017, 60, 1087-1092.	4.9	6
77	Analysis of DNA Mismatch Repair in Cellular Response to DNA Damage. Methods in Enzymology, 2006, 408, 303-317.	1.0	4
78	Proteomic analysis of mismatch repair-mediated alkylating agent-induced DNA damage response. Cell and Bioscience, 2013, 3, 37.	4.8	4
79	Interplay between H3K36me3, methyltransferase SETD2, and mismatch recognition protein MutS \pm facilitates processing of oxidative DNA damage in human cells. Journal of Biological Chemistry, 2022, 298, 102102.	3.4	4
80	Mismatch Repair. , 2014, , 1-14.		0
81	DNA repair DNA Mismatch Repair and the DNA Damage Response. , 2021, , 232-235.		0
82	DNA Mismatch Repair: Biological Functions and Molecular Mechanisms. , 2007, , 87-117.		0
83	Regulation of mismatch repair protein MutS \pm functions by its Walker A and Walker B motifs. FASEB Journal, 2013, 27, 758.9.	0.5	0
84	Coordinated processing of 3 \times slipped (CAG) n /(CTG) n hairpins by DNA polymerases preferentially induces repeat expansions. FASEB Journal, 2013, 27, 758.4.	0.5	0
85	Clamping down on mismatches. ELife, 2016, 5, .	6.0	0
86	Mismatch Repair. , 2018, , 683-695.		0