

Guo-Min Li

List of Publications by Year in descending order

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6,470
citations

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docs citations

88
times ranked

7969
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#	ARTICLE	IF	CITATIONS
1	NBS1-CtIP-mediated DNA end resection suppresses cGAS binding to micronuclei. <i>Nucleic Acids Research</i> , 2022, 50, 2681-2699.	14.5	8
2	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
3	Interplay between H3K36me3, methyltransferase SETD2, and mismatch recognition protein MutS± facilitates processing of oxidative DNA damage in human cells. <i>Journal of Biological Chemistry</i> , 2022, 298, 102102.	3.4	4
4	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
5	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
6	OTUB1 stabilizes mismatch repair protein MSH2 by blocking ubiquitination. <i>Journal of Biological Chemistry</i> , 2021, 296, 100466.	3.4	15
7	DNA repair DNA Mismatch Repair and the DNA Damage Response. , 2021, , 232-235.		0
8	MIF is a 3â€™ flap nuclease that facilitates DNA replication and promotes tumor growth. <i>Nature Communications</i> , 2021, 12, 2954.	12.8	20
9	DNA polymerase Î± promotes CAGâ€™CTG repeat expansions in Huntingtonâ€™s disease via insertion sequences of its catalytic domain. <i>Journal of Biological Chemistry</i> , 2021, 297, 101144.	3.4	7
10	Mismatch-bound human MutSâ€™MutL complex triggers DNA incisions and activates mismatch repair. <i>Cell Research</i> , 2021, 31, 542-553.	12.0	26
11	DNA mismatch repair in the chromatin context: Mechanisms and therapeutic potential. <i>DNA Repair</i> , 2020, 93, 102918.	2.8	10
12	HDAC3 deacetylates the DNA mismatch repair factor MutS2 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
13	DNA mismatch repair in the context of chromatin. <i>Cell and Bioscience</i> , 2020, 10, 10.	4.8	15
14	Phosphorylation of proliferating cell nuclear antigen promotes cancer progression by activating the ATM/Akt/GSK3Î²/Snail signaling pathway. <i>Journal of Biological Chemistry</i> , 2019, 294, 7037-7045.	3.4	20
15	HDAC6 regulates DNA damage response via deacetylating MLH1. <i>Journal of Biological Chemistry</i> , 2019, 294, 5813-5826.	3.4	28
16	Identification of novel genetic variants predisposing to familial oral squamous cell carcinomas. <i>Cell Discovery</i> , 2019, 5, 57.	6.7	16
17	Cancer-driving H3G34V/R/D mutations block H3K36 methylation and H3K36me3â€™MutS± interaction. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, 9598-9603.	7.1	87
18	DNA mismatch repair preferentially safeguards actively transcribed genes. <i>DNA Repair</i> , 2018, 71, 82-86.	2.8	24

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19	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
20	A human MUTYH variant linking colonic polyposis to redox degradation of the [4Fe4S] ₂₊ cluster. <i>Nature Chemistry</i> , 2018, 10, 873-880.	13.6	20
21	Mismatch Repair. , 2018, , 683-695.		0
22	DNA mismatch repair in trinucleotide repeat instability. <i>Science China Life Sciences</i> , 2017, 60, 1087-1092.	4.9	6
23	MutS ^{Δ2} abundance and Msh3 ATP hydrolysis activity are important drivers of CTG/CAG repeat expansions. <i>Nucleic Acids Research</i> , 2017, 45, 10068-10078.	14.5	27
24	MutS ^{Δ2} 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
25	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
26	Regulation of mismatch repair by histone code and posttranslational modifications in eukaryotic cells. <i>DNA Repair</i> , 2016, 38, 68-74.	2.8	22
27	Clamping down on mismatches. <i>ELife</i> , 2016, 5, .	6.0	0
28	Okazaki fragment maturation involves β -segment error editing by the mammalian FEN1/MutS ^{Δ2} functional complex. <i>EMBO Journal</i> , 2015, 34, 1829-1843.	7.8	28
29	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
30	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
31	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
32	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
33	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
34	Mismatch Repair. , 2014, , 1-14.		0
35	New insights and challenges in mismatch repair: Getting over the chromatin hurdle. <i>DNA Repair</i> , 2014, 19, 48-54.	2.8	33
36	HDAC6 Deacetylates and Ubiquitinates MSH2 to Maintain Proper Levels of MutS ^{Δ2} . <i>Molecular Cell</i> , 2014, 55, 31-46.	9.7	112

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37	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
38	Proteomic analysis of mismatch repair-mediated alkylating agent-induced DNA damage response. <i>Cell and Bioscience</i> , 2013, 3, 37.	4.8	4
39	The Histone Mark H3K36me3 Regulates Human DNA Mismatch Repair through Its Interaction with MutS β . <i>Cell</i> , 2013, 153, 590-600.	28.9	504
40	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
41	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
42	Coordinated Processing of 3 α -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
43	Trinucleotide repeat expansions catalyzed by human cell-free extracts. <i>Cell Research</i> , 2013, 23, 565-572.	12.0	16
44	Regulation of mismatch repair protein MutS β functions by its Walker A and Walker B motifs. <i>FASEB Journal</i> , 2013, 27, 758.9.	0.5	0
45	Coordinated processing of 3 α -slipped (CAG) $_n$ /(CTG) $_n$ hairpins by DNA polymerases preferentially induces repeat expansions. <i>FASEB Journal</i> , 2013, 27, 758.4.	0.5	0
46	The hMSH2(M688R) Lynch syndrome mutation may function as a dominant negative. <i>Carcinogenesis</i> , 2012, 33, 1647-1654.	2.8	8
47	The Werner Syndrome Protein Promotes CAG/CTG Repeat Stability by Resolving Large (CAG) $_n$ /(CTG) $_n$ Hairpins. <i>Journal of Biological Chemistry</i> , 2012, 287, 30151-30156.	3.4	17
48	Modulation of microRNA processing by mismatch repair protein MutL β . <i>Cell Research</i> , 2012, 22, 973-985.	12.0	38
49	Effect of Carcinogenic Acrolein on DNA Repair and Mutagenic Susceptibility. <i>Journal of Biological Chemistry</i> , 2012, 287, 12379-12386.	3.4	75
50	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
51	A special issue on DNA damage response and genome stability. <i>Cell and Bioscience</i> , 2012, 2, 4.	4.8	8
52	The Role of XPG in Processing (CAG) $_n$ /(CTG) $_n$ DNA Hairpins. <i>Cell and Bioscience</i> , 2011, 1, 11.	4.8	14
53	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
54	Novel molecular insights into the mechanism of GO removal by MutM. <i>Cell Research</i> , 2010, 20, 116-118.	12.0	7

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55	Evidence That Nucleosomes Inhibit Mismatch Repair in Eukaryotic Cells. <i>Journal of Biological Chemistry</i> , 2009, 284, 33056-33061.	3.4	59
56	Distinct Nucleotide Binding/Hydrolysis Properties and Molar Ratio of MutS [±] and MutS ² Determine Their Differential Mismatch Binding Activities. <i>Journal of Biological Chemistry</i> , 2009, 284, 11557-11562.	3.4	47
57	Mismatch Recognition Protein MutS ² Does Not Hijack (CAG) Hairpin Repair in Vitro. <i>Journal of Biological Chemistry</i> , 2009, 284, 20452-20456.	3.4	45
58	DNA instability in replicating Huntington's disease lymphoblasts. <i>BMC Medical Genetics</i> , 2009, 10, 11.	2.1	24
59	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
60	Mechanisms and functions of DNA mismatch repair. <i>Cell Research</i> , 2008, 18, 85-98.	12.0	1,081
61	A special issue on DNA damage responses and genome maintenance. <i>Cell Research</i> , 2008, 18, 1-2.	12.0	6
62	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
63	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
64	Identification and characterization of OGG1 mutations in patients with Alzheimer's disease. <i>Nucleic Acids Research</i> , 2007, 35, 2759-2766.	14.5	105
65	DNA Mismatch Repair: Biological Functions and Molecular Mechanisms. , 2007, , 87-117.		0
66	Analysis of DNA Mismatch Repair in Cellular Response to DNA Damage. <i>Methods in Enzymology</i> , 2006, 408, 303-317.	1.0	4
67	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
68	hMRE11 deficiency leads to microsatellite instability and defective DNA mismatch repair. <i>EMBO Reports</i> , 2005, 6, 438-444.	4.5	49
69	Reconstitution of 5'→3'-Directed Human Mismatch Repair in a Purified System. <i>Cell</i> , 2005, 122, 693-705.	28.9	316
70	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
71	Evidence for Involvement of HMGB1 Protein in Human DNA Mismatch Repair. <i>Journal of Biological Chemistry</i> , 2004, 279, 20935-20940.	3.4	112
72	Differential Requirement for Proliferating Cell Nuclear Antigen in 5'→3' and 3'→5' Nick-directed Excision in Human Mismatch Repair. <i>Journal of Biological Chemistry</i> , 2004, 279, 16912-16917.	3.4	47

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73	Folate deficiency, mismatch repair-dependent apoptosis, and human disease. <i>Journal of Nutritional Biochemistry</i> , 2003, 14, 568-575.	4.2	47
74	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
75	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
76	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
77	DNA mismatch repair and cancer. <i>Frontiers in Bioscience - Landmark</i> , 2003, 8, d997-1017.	3.0	66
78	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
79	Involvement of DNA mismatch repair in folate deficiency-induced apoptosis. <i>Journal of Nutritional Biochemistry</i> , 2002, 13, 355-363.	4.2	36
80	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
81	Mismatch repair deficiency in hematological malignancies with microsatellite instability. <i>Oncogene</i> , 2002, 21, 5758-5764.	5.9	41
82	Specific Binding of Human MSH2-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
83	Mismatch Repair Processing of Carcinogen-DNA Adducts Triggers Apoptosis. <i>Molecular and Cellular Biology</i> , 1999, 19, 8292-8301.	2.3	115
84	Increased transversions in a novel mutator colon cancer cell line. <i>Oncogene</i> , 1998, 16, 1125-1130.	5.9	13
85	Human MutS Specifically Binds to DNA Containing Aminofluorene and Acetylaminofluorene Adducts. <i>Journal of Biological Chemistry</i> , 1996, 271, 24084-24088.	3.4	54
86	Hypermutability and mismatch repair deficiency in RER+ tumor cells. <i>Cell</i> , 1993, 75, 1227-1236.	28.9	1,031