## **Richard Fishel**

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	The human mutator gene homolog MSH2 and its association with hereditary nonpolyposis colon cancer. Cell, 1993, 75, 1027-1038.	28.9	2,706
2	Mutation in the DNA mismatch repair gene homologue hMLH 1 is associated with hereditary non-polyposis colon cancer. Nature, 1994, 368, 258-261.	27.8	2,001
3	Cytochrome c and dATP-mediated Oligomerization of Apaf-1 Is a Prerequisite for Procaspase-9 Activation. Journal of Biological Chemistry, 1999, 274, 17941-17945.	3.4	432
4	hMSH4-hMSH5 Recognizes Holliday Junctions and Forms a Meiosis-Specific Sliding Clamp that Embraces Homologous Chromosomes. Molecular Cell, 2004, 15, 437-451.	9.7	361
5	hMSH2–hMSH6 Forms a Hydrolysis-Independent Sliding Clamp on Mismatched DNA. Molecular Cell, 1999, 3, 255-261.	9.7	338
6	The Human Mismatch Recognition Complex hMSH2-hMSH6 Functions as a Novel Molecular Switch. Cell, 1997, 91, 995-1005.	28.9	336
7	MicroRNA-21 induces resistance to 5-fluorouracil by down-regulating human DNA MutS homolog 2 (hMSH2). Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 21098-21103.	7.1	333
8	DNA Damage-Dependent Acetylation and Ubiquitination of H2AX Enhances Chromatin Dynamics. Molecular and Cellular Biology, 2007, 27, 7028-7040.	2.3	327
9	Modulation of mismatch repair and genomic stability by miR-155. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 6982-6987.	7.1	306
10	Identification of mismatch repair genes and their role in the development of cancer. Current Opinion in Genetics and Development, 1995, 5, 382-395.	3.3	293
11	BCR/ABL Regulates Mammalian RecA Homologs, Resulting in Drug Resistance. Molecular Cell, 2001, 8, 795-806.	9.7	290
12	Deficiencies in Mouse <b> <i>Myh</i> </b> and <b> <i>Ogg1</i> </b> Result in Tumor Predisposition and G to T Mutations in Codon 12 of the <b> <i>K-Ras</i> </b> Oncogene in Lung Tumors. Cancer Research, 2004, 64, 3096-3102.	0.9	271
13	The Coordinated Functions of the E. coli MutS and MutL Proteins in Mismatch Repair. Molecular Cell, 2003, 12, 233-246.	9.7	259
14	Fusion Tyrosine Kinases Induce Drug Resistance by Stimulation of Homology-Dependent Recombination Repair, Prolongation of G 2 /M Phase, and Protection from Apoptosis. Molecular and Cellular Biology, 2002, 22, 4189-4201.	2.3	188
15	Mismatch Repair. Journal of Biological Chemistry, 2015, 290, 26395-26403.	3.4	181
16	Increased Hypermutation at G and C Nucleotides in Immunoglobulin Variable Genes from Mice Deficient in the MSH2 Mismatch Repair Protein. Journal of Experimental Medicine, 1998, 187, 1745-1751.	8.5	170
17	A Human REV7 Homolog That Interacts with the Polymerase ζ Catalytic Subunit hREV3 and the Spindle Assembly Checkpoint Protein hMAD2. Journal of Biological Chemistry, 2000, 275, 4391-4397.	3.4	170
18	The mismatch-repair protein hMSH2 binds selectively to DNA adducts of the anticancer drug cisplatin. Chemistry and Biology, 1996, 3, 579-589.	6.0	167

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19	The Interaction of the Human MutL Homologues in Hereditary Nonpolyposis Colon Cancer. Journal of Biological Chemistry, 1999, 274, 6336-6341.	3.4	153
20	Extreme Heterogeneity in the Molecular Events Leading to the Establishment of Chiasmata during Meiosis I in Human Oocytes. American Journal of Human Genetics, 2005, 76, 112-127.	6.2	151
21	MutS homologs in mammalian cells. Current Opinion in Genetics and Development, 1997, 7, 105-113.	3.3	150
22	Activation of Human MutS Homologs by 8-Oxo-guanine DNA Damage. Journal of Biological Chemistry, 2002, 277, 8260-8266.	3.4	149
23	Mismatch Repair during Homologous and Homeologous Recombination. Cold Spring Harbor Perspectives in Biology, 2015, 7, a022657.	5.5	146
24	The Interaction of DNA Mismatch Repair Proteins with Human Exonuclease I. Journal of Biological Chemistry, 2001, 276, 33011-33018.	3.4	133
25	Signaling mismatch repair in cancer. Nature Medicine, 1999, 5, 1239-1241.	30.7	126
26	Interactions of Human hMSH2 with hMSH3 and hMSH2 with hMSH6: Examination of Mutations Found in Hereditary Nonpolyposis Colorectal Cancer. Molecular and Cellular Biology, 1998, 18, 6616-6623.	2.3	123
27	MutS switches between two fundamentally distinct clamps during mismatch repair. Nature Structural and Molecular Biology, 2011, 18, 379-385.	8.2	120
28	Mismatch Repair and the Hereditary Non-polyposis Colorectal Cancer Syndrome (HNPCC). Cancer Investigation, 2002, 20, 102-109.	1.3	117
29	DNA Mismatch Repair-dependent Response to Fluoropyrimidine-generated Damage. Journal of Biological Chemistry, 2005, 280, 5516-5526.	3.4	108
30	The Role of Mismatched Nucleotides in Activating the hMSH2-hMSH6 Molecular Switch. Journal of Biological Chemistry, 2000, 275, 3922-3930.	3.4	103
31	DNA Repair and Tumorigenesis: Lessons from Hereditary Cancer Syndromes. Cancer Biology and Therapy, 2002, 1, 477-485.	3.4	101
32	Current and emerging trends in Lynch syndrome identification in women with endometrial cancer. Gynecologic Oncology, 2009, 114, 128-134.	1.4	97
33	Cascading MutS and MutL sliding clamps control DNA diffusion to activate mismatch repair. Nature, 2016, 539, 583-587.	27.8	91
34	Dissociation of Mismatch Recognition and ATPase Activity by hMSH2-hMSH3. Journal of Biological Chemistry, 1999, 274, 21659-21664.	3.4	90
35	ATP Alters the Diffusion Mechanics of MutS on Mismatched DNA. Structure, 2012, 20, 1264-1274.	3.3	87
36	Severe Attenuation of the B Cell Immune Response in Msh2-deficient Mice. Journal of Experimental Medicine, 1999, 189, 471-482.	8.5	80

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37	BRCA1 and cell signaling. Oncogene, 2000, 19, 6152-6158.	5.9	80
38	Colorectal carcinomas with high microsatellite instability: Defining a distinct immunologic and molecular entity with respect to prognostic markers. Human Pathology, 2000, 31, 1506-1514.	2.0	76
39	Biochemical Characterization of the Human RAD51 Protein. Journal of Biological Chemistry, 2002, 277, 14417-14425.	3.4	71
40	Female embryonic lethality in mice nullizygous for both Msh2 and p53. Nature Genetics, 1997, 17, 114-118.	21.4	67
41	DNA mismatch repair (MMR)â€dependent 5â€fluorouracil cytotoxicity and the potential for new therapeutic targets. British Journal of Pharmacology, 2009, 158, 679-692.	5.4	66
42	The mechanism of mismatch repair and the functional analysis of mismatch repair defects in Lynch syndrome. Familial Cancer, 2013, 12, 159-168.	1.9	65
43	Nucleosome Remodeling by hMSH2-hMSH6. Molecular Cell, 2009, 36, 1086-1094.	9.7	62
44	HNPCC mutations in hMSH2 result in reduced hMSH2-hMSH6 molecular switch functions. Cancer Cell, 2002, 1, 469-478.	16.8	59
45	Sequence context effect for hMSH2-hMSH6 mismatch-dependent activation. Proceedings of the National Academy of Sciences of the United States of America, 2009, 106, 4177-4182.	7.1	59
46	Adenosine nucleotide modulates the physical interaction between hMSH2 and BRCA1. Oncogene, 2001, 20, 4640-4649.	5.9	57
47	Sequence analysis of the mismatch repair gene hMSH6 in the germline of patients with familial and sporadic colorectal cancer. , 2000, 85, 606-613.		53
48	A quantitative model of nucleosome dynamics. Nucleic Acids Research, 2011, 39, 8306-8313.	14.5	49
49	Nitric Oxide–Donating Aspirin Derivatives Suppress Microsatellite Instability in Mismatch Repair–Deficient and Hereditary Nonpolyposis Colorectal Cancer Cells. Cancer Research, 2007, 67, 10966-10975.	0.9	47
50	hXRCC2 Enhances ADP/ATP Processing and Strand Exchange by hRAD51. Journal of Biological Chemistry, 2004, 279, 30385-30394.	3.4	44
51	hMSH4-hMSH5 Adenosine Nucleotide Processing and Interactions with Homologous Recombination Machinery. Journal of Biological Chemistry, 2008, 283, 145-154.	3.4	44
52	The Effect of O6-Methylguanine DNA Adducts on the Adenosine Nucleotide Switch Functions of hMSH2-hMSH6 and hMSH2-hMSH3. Journal of Biological Chemistry, 2000, 275, 27851-27857.	3.4	38
53	The Base Excision Repair Pathway Is Required for Efficient Lentivirus Integration. PLoS ONE, 2011, 6, e17862.	2.5	38
54	Mismatch repair protein hMSH2–hMSH6 recognizes mismatches and forms sliding clamps within a D-loop recombination intermediate. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, E316-25.	7.1	37

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55	Single-molecule views of MutS on mismatched DNA. DNA Repair, 2014, 20, 82-93.	2.8	37
56	Dynamic control of strand excision during human DNA mismatch repair. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 3281-3286.	7.1	35
57	Human MSH2 (hMSH2) Protein Controls ATP Processing by hMSH2-hMSH6. Journal of Biological Chemistry, 2011, 286, 40287-40295.	3.4	33
58	Single-Molecule Analysis Reveals the Kinetics and Physiological Relevance of MutL-ssDNA Binding. PLoS ONE, 2010, 5, e15496.	2.5	32
59	Molecular diagnostics of cancer predisposition: hereditary non-polyposis colorectal carcinoma and mismatch repair defects. Biochimica Et Biophysica Acta: Reviews on Cancer, 1999, 1423, O1-O10.	7.4	30
60	Retroviral intasomes search for a target DNA by 1D diffusion which rarely results in integration. Nature Communications, 2016, 7, 11409.	12.8	29
61	Homologous Recombination in Eukaryotes. Progress in Molecular Biology and Translational Science, 2012, 110, 155-206.	1.7	28
62	Biochemical Characterization of the Human RAD51 Protein. Journal of Biological Chemistry, 2002, 277, 14426-14433.	3.4	25
63	Lynch Syndrome: Form, Function, Proteins, and Basketball. Gastroenterology, 2005, 129, 751-755.	1.3	25
64	Lynch Syndrome: Form, Function, Proteins, and Basketball. Gastroenterology, 2005, 129, 751-755.	1.3	25
65	Evidence that hMLH3 functions primarily in meiosis and in hMSH2-hMSH3 mismatch repair. Cancer Biology and Therapy, 2009, 8, 1411-1420.	3.4	24
66	Widespread nuclease contamination in commonly used oxygen-scavenging systems. Nature Methods, 2015, 12, 901-902.	19.0	24
67	Pseudomonas aeruginosa AmrZ Binds to Four Sites in the <i>algD</i> Promoter, Inducing DNA-AmrZ Complex Formation and Transcriptional Activation. Journal of Bacteriology, 2016, 198, 2673-2681.	2.2	22
68	Long repeating (TTAGGG) single-stranded DNA self-condenses into compact beaded filaments stabilized by G-quadruplex formation. Journal of Biological Chemistry, 2018, 293, 9473-9485.	3.4	22
69	Multiple Pathways Leading to Genomic Instabiligy and Tumorigenesis. Annals of the New York Academy of Sciences, 1994, 726, 165-177.	3.8	20
70	An Efficient Site-Specific Method for Irreversible Covalent Labeling of Proteins with a Fluorophore. Scientific Reports, 2015, 5, 16883.	3.3	20
71	MutL sliding clamps coordinate exonuclease-independent Escherichia coli mismatch repair. Nature Communications, 2019, 10, 5294.	12.8	20
72	Female embryonic lethality in Msh2–Trp53 nullizygous mice is strain dependent. Mammalian Genome, 1999, 10, 1020-1022.	2.2	16

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73	Strategies for Targeting Retroviral Integration for Safer Gene Therapy: Advances and Challenges. Frontiers in Molecular Biosciences, 2021, 8, 662331.	3.5	16
74	The P seudomonas aeruginosa †AmrZ C â€ŧerminal domain mediates tetramerization and is required for its activator and repressor functions. Environmental Microbiology Reports, 2016, 8, 85-90.	2.4	15
75	The GTP hydrolysis defect of theSaccharomyces cerevisiae mutant G-protein Gpa1G50V. Yeast, 2000, 16, 387-400.	1.7	13
76	Stochastic Processes and Component Plasticity Governing DNA Mismatch Repair. Journal of Molecular Biology, 2018, 430, 4456-4468.	4.2	13
77	Linker domain function predicts pathogenic MLH1 missense variants. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	13
78	Repair of Oxidative DNA Base Damage in the Host Genome Influences the HIV Integration Site Sequence Preference. PLoS ONE, 2014, 9, e103164.	2.5	12
79	Magnesium influences the discrimination and release of ADP by human RAD51. DNA Repair, 2006, 5, 704-717.	2.8	11
80	Coordinating Multi-Protein Mismatch Repair by Managing Diffusion Mechanics on the DNA. Journal of Molecular Biology, 2018, 430, 4469-4480.	4.2	11
81	Expression and purification of nuclease-free protocatechuate 3,4-dioxygenase for prolonged single-molecule fluorescence imaging. Analytical Biochemistry, 2018, 556, 78-84.	2.4	11
82	MSI-Testing in Hereditary Non-Polyposis Colorectal Carcinoma (HNPCC). Disease Markers, 2004, 20, 225-236.	1.3	10
83	Mutation of TGFÎ <sup>2</sup> -RII eliminates NSAID cancer chemoprevention. Oncotarget, 2018, 9, 12554-12561.	1.8	10
84	Enrichment and characterization of histones by two-dimensional hydroxyapatite/reversed-phase liquid chromatography–mass spectrometry. Analytical Biochemistry, 2009, 388, 47-55.	2.4	9
85	The hMSH2(M688R) Lynch syndrome mutation may function as a dominant negative. Carcinogenesis, 2012, 33, 1647-1654.	2.8	8
86	Dynamic unwrapping of nucleosomes by HsRAD51 that includes sliding and rotational motion of histone octamers. Nucleic Acids Research, 2017, 45, 685-698.	14.5	8
87	Nucleosome DNA unwrapping does not affect prototype foamy virus integration efficiency or site selection. PLoS ONE, 2019, 14, e0212764.	2.5	8
88	MutS homolog sliding clamps shield the DNA from binding proteins. Journal of Biological Chemistry, 2018, 293, 14285-14294.	3.4	5
89	Retroviral prototype foamy virus intasome binding to a nucleosome target does not determine integration efficiency. Journal of Biological Chemistry, 2021, 296, 100550.	3.4	5
90	Plasticity of Multi-Protein Complexes. Journal of Molecular Biology, 2018, 430, 4441-4442.	4.2	3

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91	Evolutionary advantage of a dissociative search mechanism in DNA mismatch repair. Physical Review E, 2021, 103, 052404.	2.1	2
92	Mass COVID-19 patient screening using UvsX and UvsY mediated DNA recombination and high throughput parallel sequencing. Scientific Reports, 2022, 12, 4082.	3.3	2
93	Harnessing mismatch repair to model sporadic cancers. Nature Methods, 2008, 5, 225-226.	19.0	1
94	Enhanced gene targeting to evaluate Lynch syndrome alterations. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 3918-3920.	7.1	1
95	Mismatch Repair. , 2016, , 305-339.		1
96	The mechanics of DNA mismatch repair sliding clamp progression in the prediction of clinically relevant HsMLH1 missense mutations. FASEB Journal, 2021, 35, .	0.5	0
97	Mechanism of DNA Mismatch Repair from Bacteria to Human. , 2005, , .		0