

Jeremy M Stark

List of Publications by Year in descending order

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63
papers

6,408
citations

101543

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docs citations

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times ranked

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citing authors

#	ARTICLE	IF	CITATIONS
1	Microarray screening reveals two non-conventional SUMO-binding modules linked to DNA repair by non-homologous end-joining. <i>Nucleic Acids Research</i> , 2022, 50, 4732-4754.	14.5	4
2	The importance of DNAPKcs for blunt DNA end joining is magnified when XLF is weakened. <i>Nature Communications</i> , 2022, 13, .	12.8	9
3	m ⁶ A deposition is regulated by PRMT1-mediated arginine methylation of METTL14 in its disordered C-terminal region. <i>EMBO Journal</i> , 2021, 40, e106309.	7.8	30
4	Pol η reverse transcribes RNA and promotes RNA-templated DNA repair. <i>Science Advances</i> , 2021, 7, .	10.3	51
5	R-Loop Associated Mitotic Stress Confers Vulnerabilities in Splicing Factor Mutant Leukemia. <i>Blood</i> , 2021, 138, 3713-3713.	1.4	0
6	Functional deficiency of DNA repair gene EXO5 results in androgen-induced genomic instability and prostate tumorigenesis. <i>Oncogene</i> , 2020, 39, 1246-1259.	5.9	8
7	53BP1 Enforces Distinct Pre- and Post-resection Blocks on Homologous Recombination. <i>Molecular Cell</i> , 2020, 77, 26-38.e7.	9.7	85
8	The canonical non-homologous end joining factor XLF promotes chromosomal deletion rearrangements in human cells. <i>Journal of Biological Chemistry</i> , 2020, 295, 125-137.	3.4	12
9	RNF8 has both KLU-dependent and independent roles in chromosomal break repair. <i>Nucleic Acids Research</i> , 2020, 48, 6032-6052.	14.5	15
10	The <i>RAD52</i> S346X variant reduces risk of developing breast cancer in carriers of pathogenic germline <i>BRCA2</i> mutations. <i>Molecular Oncology</i> , 2020, 14, 1124-1133.	4.6	13
11	USP22 Interacts with PALB2 and Promotes Chemotherapy Resistance via Homologous Recombination of DNA Double-Strand Breaks. <i>Molecular Cancer Research</i> , 2020, 18, 424-435.	3.4	12
12	BLM has Contrary Effects on Repeat-Mediated Deletions, based on the Distance of DNA DSBs to a Repeat and Repeat Divergence. <i>Cell Reports</i> , 2020, 30, 1342-1357.e4.	6.4	17
13	XLF acts as a flexible connector during non-homologous end joining. <i>ELife</i> , 2020, 9, .	6.0	22
14	Distinct roles of RAD52 and POLQ in chromosomal break repair and replication stress response. <i>PLoS Genetics</i> , 2019, 15, e1008319.	3.5	60
15	Electrophilic fatty acids impair RAD51 function and potentiate the effects of DNA-damaging agents on growth of triple-negative breast cells. <i>Journal of Biological Chemistry</i> , 2019, 294, 397-404.	3.4	16
16	Repeat-mediated deletions can be induced by a chromosomal break far from a repeat, but multiple pathways suppress such rearrangements. <i>Genes and Development</i> , 2018, 32, 524-536.	5.9	35
17	C-NHEJ without indels is robust and requires synergistic function of distinct XLF domains. <i>Nature Communications</i> , 2018, 9, 2484.	12.8	75
18	The Anticancer Activity of a First-in-class Small-molecule Targeting PCNA. <i>Clinical Cancer Research</i> , 2018, 24, 6053-6065.	7.0	27

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19	Contribution of canonical nonhomologous end joining to chromosomal rearrangements is enhanced by ATM kinase deficiency. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, 728-733.	7.1	28
20	PRPF8 is important for BRCA1-mediated homologous recombination. <i>Oncotarget</i> , 2017, 8, 93319-93337.	1.8	15
21	Myelodysplasia-associated mutations in serine/arginine-rich splicing factor SRSF2 lead to alternative splicing of CDC25C. <i>BMC Molecular Biology</i> , 2016, 17, 18.	3.0	7
22	Tetratricopeptide repeat factor XAB2 mediates the end resection step of homologous recombination. <i>Nucleic Acids Research</i> , 2016, 44, 5702-5716.	14.5	26
23	Regulation of Single-Strand Annealing and its Role in Genome Maintenance. <i>Trends in Genetics</i> , 2016, 32, 566-575.	6.7	344
24	RNF4 regulates DNA double-strand break repair in a cell cycle-dependent manner. <i>Cell Cycle</i> , 2016, 15, 787-798.	2.6	23
25	Heavy Metal Exposure Influences Double Strand Break DNA Repair Outcomes. <i>PLoS ONE</i> , 2016, 11, e0151367.	2.5	107
26	EXO1 is critical for embryogenesis and the DNA damage response in mice with a hypomorphic <i>Nbs1</i> allele. <i>Nucleic Acids Research</i> , 2015, 43, 7371-7387.	14.5	16
27	DNA Damage Response Factors from Diverse Pathways, Including DNA Crosslink Repair, Mediate Alternative End Joining. <i>PLoS Genetics</i> , 2015, 11, e1004943.	3.5	114
28	An RNF168 fragment defective for focal accumulation at DNA damage is proficient for inhibition of homologous recombination in BRCA1 deficient cells. <i>Nucleic Acids Research</i> , 2014, 42, 7720-7733.	14.5	28
29	TGF β ² Induces α -BRCAness and Sensitivity to PARP Inhibition in Breast Cancer by Regulating DNA-Repair Genes. <i>Molecular Cancer Research</i> , 2014, 12, 1597-1609.	3.4	56
30	A PCNA-Derived Cell Permeable Peptide Selectively Inhibits Neuroblastoma Cell Growth. <i>PLoS ONE</i> , 2014, 9, e94773.	2.5	22
31	Visualization of local DNA unwinding by Mre11/Rad50/Nbs1 using single-molecule FRET. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 18868-18873.	7.1	55
32	HP1 promotes tumor suppressor BRCA1 functions during the DNA damage response. <i>Nucleic Acids Research</i> , 2013, 41, 5784-5798.	14.5	83
33	The interaction between CtIP and BRCA1 is not essential for resection-mediated DNA repair or tumor suppression. <i>Journal of Cell Biology</i> , 2013, 201, 693-707.	5.2	71
34	The HSV-1 Exonuclease, UL12, Stimulates Recombination by a Single Strand Annealing Mechanism. <i>PLoS Pathogens</i> , 2012, 8, e1002862.	4.7	80
35	Suberoylanilide Hydroxamic Acid as a Radiosensitizer through Modulation of RAD51 Protein and Inhibition of Homology-Directed Repair in Multiple Myeloma. <i>Molecular Cancer Research</i> , 2012, 10, 1052-1064.	3.4	47
36	Examining the roles of DNA2 during mammalian end resection. <i>Cell Cycle</i> , 2012, 11, 4111-4111.	2.6	0

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37	Ring Finger Nuclear Factor RNF168 Is Important for Defects in Homologous Recombination Caused by Loss of the Breast Cancer Susceptibility Factor BRCA1. <i>Journal of Biological Chemistry</i> , 2012, 287, 40618-40628.	3.4	44
38	RI-1: a chemical inhibitor of RAD51 that disrupts homologous recombination in human cells. <i>Nucleic Acids Research</i> , 2012, 40, 7347-7357.	14.5	154
39	A Dual Role for UVRAG in Maintaining Chromosomal Stability Independent of Autophagy. <i>Developmental Cell</i> , 2012, 22, 1001-1016.	7.0	90
40	I-SceI-Based Assays to Examine Distinct Repair Outcomes of Mammalian Chromosomal Double Strand Breaks. <i>Methods in Molecular Biology</i> , 2012, 920, 379-391.	0.9	271
41	Chronic myelogenous leukemia stem and progenitor cells demonstrate chromosomal instability related to repeated breakage-fusion-bridge cycles mediated by increased nonhomologous end joining. <i>Blood</i> , 2012, 119, 6187-6197.	1.4	42
42	Subtelomeric regions in mammalian cells are deficient in DNA double-strand break repair. <i>DNA Repair</i> , 2011, 10, 536-544.	2.8	34
43	Trex2 Enables Spontaneous Sister Chromatid Exchanges Without Facilitating DNA Double-Strand Break Repair. <i>Genetics</i> , 2011, 188, 787-797.	2.9	15
44	The relative efficiency of homology-directed repair has distinct effects on proper anaphase chromosome separation. <i>Nucleic Acids Research</i> , 2011, 39, 5935-5944.	14.5	41
45	Interhomolog recombination and loss of heterozygosity in wild-type and Bloom syndrome helicase (BLM)-deficient mammalian cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 11971-11976.	7.1	72
46	Protein phosphatase PP6 is required for homology-directed repair of DNA double-strand breaks. <i>Cell Cycle</i> , 2011, 10, 1411-1419.	2.6	45
47	Correct End Use during End Joining of Multiple Chromosomal Double Strand Breaks Is Influenced by Repair Protein RAD50, DNA-dependent Protein Kinase DNA-PKcs, and Transcription Context. <i>Journal of Biological Chemistry</i> , 2011, 286, 42470-42482.	3.4	95
48	Mammalian Fbh1 is important to restore normal mitotic progression following decatenation stress. <i>DNA Repair</i> , 2010, 9, 708-717.	2.8	18
49	Physical Interaction of RECQ5 Helicase with RAD51 Facilitates Its Anti-recombinase Activity. <i>Journal of Biological Chemistry</i> , 2010, 285, 15739-15745.	3.4	78
50	ATM Limits Incorrect End Utilization during Non-Homologous End Joining of Multiple Chromosome Breaks. <i>PLoS Genetics</i> , 2010, 6, e1001194.	3.5	43
51	53BP1 Inhibits Homologous Recombination in Brca1-Deficient Cells by Blocking Resection of DNA Breaks. <i>Cell</i> , 2010, 141, 243-254.	28.9	1,406
52	Limiting the Persistence of a Chromosome Break Diminishes Its Mutagenic Potential. <i>PLoS Genetics</i> , 2009, 5, e1000683.	3.5	77
53	Alternative-NHEJ Is a Mechanistically Distinct Pathway of Mammalian Chromosome Break Repair. <i>PLoS Genetics</i> , 2008, 4, e1000110.	3.5	708
54	RECQL5/Recql5 helicase regulates homologous recombination and suppresses tumor formation via disruption of Rad51 presynaptic filaments. <i>Genes and Development</i> , 2007, 21, 3073-3084.	5.9	283

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55	Suppression of the DNA repair defects of BRCA2-deficient cells with heterologous protein fusions. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2006, 103, 8768-8773.	7.1	91
56	Genetic Steps of Mammalian Homologous Repair with Distinct Mutagenic Consequences. <i>Molecular and Cellular Biology</i> , 2004, 24, 9305-9316.	2.3	439
57	Rad51 overexpression promotes alternative double-strand break repair pathways and genome instability. <i>Oncogene</i> , 2004, 23, 546-553.	5.9	216
58	Extensive Loss of Heterozygosity Is Suppressed during Homologous Repair of Chromosomal Breaks. <i>Molecular and Cellular Biology</i> , 2003, 23, 733-743.	2.3	110
59	ATP Hydrolysis by Mammalian RAD51 Has a Key Role during Homology-directed DNA Repair. <i>Journal of Biological Chemistry</i> , 2002, 277, 20185-20194.	3.4	124
60	Variant XRCC3 implicated in cancer is functional in homology-directed repair of double-strand breaks. <i>Oncogene</i> , 2002, 21, 4176-4180.	5.9	107
61	Double-strand breaks and tumorigenesis. <i>Trends in Cell Biology</i> , 2001, 11, S52-S59.	7.9	135
62	The Relative Strengths of SR Protein-mediated Associations of Alternative and Constitutive Exons Can Influence Alternative Splicing. <i>Journal of Biological Chemistry</i> , 1999, 274, 29838-29842.	3.4	12
63	Multiple Regions of Yeast Ribosomal Protein L1 Are Important for Its Interaction with 5 S rRNA and Assembly into Ribosomes. <i>Journal of Biological Chemistry</i> , 1995, 270, 30148-30156.	3.4	43