

David Cortez

List of Publications by Year in descending order

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Version: 2024-02-01

64
papers

10,219
citations

81743

39
h-index

114278

63
g-index

75
all docs

75
docs citations

75
times ranked

9316
citing authors

#	ARTICLE	IF	CITATIONS
1	Oligomerization of DNA replication regulatory protein RADX is essential to maintain replication fork stability. <i>Journal of Biological Chemistry</i> , 2022, 298, 101672.	1.6	5
2	CHK1 phosphorylates PRIMPOL to promote replication stress tolerance. <i>Science Advances</i> , 2022, 8, eabm0314.	4.7	22
3	Topoisomerase II poisons inhibit vertebrate DNA replication through distinct mechanisms. <i>EMBO Journal</i> , 2022, 41, e110632.	3.5	7
4	ATR activation is regulated by dimerization of ATR activating proteins. <i>Journal of Biological Chemistry</i> , 2021, 296, 100455.	1.6	12
5	RADX controls RAD51 filament dynamics to regulate replication fork stability. <i>Molecular Cell</i> , 2021, 81, 1074-1083.e5.	4.5	26
6	RADX prevents genome instability by confining replication fork reversal to stalled forks. <i>Molecular Cell</i> , 2021, 81, 3007-3017.e5.	4.5	19
7	Two replication fork remodeling pathways generate nuclease substrates for distinct fork protection factors. <i>Science Advances</i> , 2020, 6, .	4.7	53
8	HMCES Maintains Replication Fork Progression and Prevents Double-Strand Breaks in Response to APOBEC Deamination and Abasic Site Formation. <i>Cell Reports</i> , 2020, 31, 107705.	2.9	51
9	The plasticity of DNA replication forks in response to clinically relevant genotoxic stress. <i>Nature Reviews Molecular Cell Biology</i> , 2020, 21, 633-651.	16.1	198
10	New insights into abasic site repair and tolerance. <i>DNA Repair</i> , 2020, 90, 102866.	1.3	86
11	Perturbing cohesin dynamics drives MRE11 nuclease-dependent replication fork slowing. <i>Nucleic Acids Research</i> , 2019, 47, 1294-1310.	6.5	38
12	Functional Analysis of the Replication Fork Proteome Identifies BET Proteins as PCNA Regulators. <i>Cell Reports</i> , 2019, 28, 3497-3509.e4.	2.9	75
13	Protection of abasic sites during DNA replication by a stable thiazolidine protein-DNA cross-link. <i>Nature Structural and Molecular Biology</i> , 2019, 26, 613-618.	3.6	69
14	Replication-Coupled DNA Repair. <i>Molecular Cell</i> , 2019, 74, 866-876.	4.5	178
15	Common motifs in ETAA1 and TOPBP1 required for ATR kinase activation. <i>Journal of Biological Chemistry</i> , 2019, 294, 8395-8402.	1.6	29
16	Quantitative phosphoproteomics reveals mitotic function of the ATR activator ETAA1. <i>Journal of Cell Biology</i> , 2019, 218, 1235-1249.	2.3	45
17	HMCES Maintains Genome Integrity by Shielding Abasic Sites in Single-Strand DNA. <i>Cell</i> , 2019, 176, 144-153.e13.	13.5	127
18	RPA and RAD51: fork reversal, fork protection, and genome stability. <i>Nature Structural and Molecular Biology</i> , 2018, 25, 446-453.	3.6	264

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19	RADX Modulates RAD51 Activity to Control Replication Fork Protection. <i>Cell Reports</i> , 2018, 24, 538-545.	2.9	88
20	An intrinsic S/G ₂ checkpoint enforced by ATR. <i>Science</i> , 2018, 361, 806-810.	6.0	215
21	Mutations in DONSON disrupt replication fork stability and cause microcephalic dwarfism. <i>Nature Genetics</i> , 2017, 49, 537-549.	9.4	81
22	Proteomic Analyses of the Eukaryotic Replication Machinery. <i>Methods in Enzymology</i> , 2017, 591, 33-53.	0.4	25
23	Functions of SMARCAL1, ZRANB3, and HLTf in maintaining genome stability. <i>Critical Reviews in Biochemistry and Molecular Biology</i> , 2017, 52, 696-714.	2.3	105
24	Replication Fork Slowing and Reversal upon DNA Damage Require PCNA Polyubiquitination and ZRANB3 DNA Translocase Activity. <i>Molecular Cell</i> , 2017, 67, 882-890.e5.	4.5	190
25	RADX Promotes Genome Stability and Modulates Chemosensitivity by Regulating RAD51 at Replication Forks. <i>Molecular Cell</i> , 2017, 67, 374-386.e5.	4.5	153
26	The essential kinase ATR: ensuring faithful duplication of a challenging genome. <i>Nature Reviews Molecular Cell Biology</i> , 2017, 18, 622-636.	16.1	589
27	ETAA1 acts at stalled replication forks to maintain genome integrity. <i>Nature Cell Biology</i> , 2016, 18, 1185-1195.	4.6	204
28	Replication fork stability confers chemoresistance in BRCA-deficient cells. <i>Nature</i> , 2016, 535, 382-387.	13.7	685
29	Identification of a Substrate Recognition Domain in the Replication Stress Response Protein Zinc Finger Ran-binding Domain-containing Protein 3 (ZRANB3). <i>Journal of Biological Chemistry</i> , 2016, 291, 8251-8257.	1.6	18
30	SMARCAL1 and telomeres: Replicating the troublesome ends. <i>Nucleus</i> , 2016, 7, 270-274.	0.6	11
31	A novel splice site mutation in <i>SMARCAL1</i> results in aberrant exon definition in a child with schimke immunosseous dysplasia. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 2260-2264.	0.7	15
32	Simian Virus Large T Antigen Interacts with the N-Terminal Domain of the 70 kD Subunit of Replication Protein A in the Same Mode as Multiple DNA Damage Response Factors. <i>PLoS ONE</i> , 2015, 10, e0116093.	1.1	7
33	High-affinity DNA-binding Domains of Replication Protein A (RPA) Direct SMARCAL1-dependent Replication Fork Remodeling. <i>Journal of Biological Chemistry</i> , 2015, 290, 4110-4117.	1.6	55
34	SMARCAL1 maintains telomere integrity during DNA replication. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 14864-14869.	3.3	67
35	Stephen Elledge and the DNA damage response. <i>DNA Repair</i> , 2015, 35, 156-157.	1.3	0
36	Enhancer of Rudimentary Homolog Affects the Replication Stress Response through Regulation of RNA Processing. <i>Molecular and Cellular Biology</i> , 2015, 35, 2979-2990.	1.1	26

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37	Preventing replication fork collapse to maintain genome integrity. <i>DNA Repair</i> , 2015, 32, 149-157.	1.3	169
38	A whole genome RNAi screen identifies replication stress response genes. <i>DNA Repair</i> , 2015, 35, 55-62.	1.3	15
39	The Replication Checkpoint Prevents Two Types of Fork Collapse without Regulating Replisome Stability. <i>Molecular Cell</i> , 2015, 59, 998-1010.	4.5	301
40	A Synthetic Lethal Screen Identifies DNA Repair Pathways that Sensitize Cancer Cells to Combined ATR Inhibition and Cisplatin Treatments. <i>PLoS ONE</i> , 2015, 10, e0125482.	1.1	99
41	Fork reversal, too much of a good thing. <i>Cell Cycle</i> , 2014, 13, 1049-1050.	1.3	17
42	SV40 Utilizes ATM Kinase Activity to Prevent Non-homologous End Joining of Broken Viral DNA Replication Products. <i>PLoS Pathogens</i> , 2014, 10, e1004536.	2.1	21
43	ATR Pathway Inhibition Is Synthetically Lethal in Cancer Cells with ERCC1 Deficiency. <i>Cancer Research</i> , 2014, 74, 2835-2845.	0.4	112
44	Mutation of Serine 1333 in the ATR HEAT Repeats Creates a Hyperactive Kinase. <i>PLoS ONE</i> , 2014, 9, e99397.	1.1	14
45	Substrate-Selective Repair and Restart of Replication Forks by DNA Translocases. <i>Cell Reports</i> , 2013, 3, 1958-1969.	2.9	136
46	ATR phosphorylates SMARCAL1 to prevent replication fork collapse. <i>Genes and Development</i> , 2013, 27, 1610-1623.	2.7	343
47	Identification and Characterization of SMARCAL1 Protein Complexes. <i>PLoS ONE</i> , 2013, 8, e63149.	1.1	26
48	SMARCAL1 catalyzes fork regression and Holliday junction migration to maintain genome stability during DNA replication. <i>Genes and Development</i> , 2012, 26, 151-162.	2.7	235
49	Defining genome maintenance pathways using functional genomic approaches. <i>Critical Reviews in Biochemistry and Molecular Biology</i> , 2011, 46, 327-341.	2.3	5
50	Thr-1989 Phosphorylation Is a Marker of Active Ataxia Telangiectasia-mutated and Rad3-related (ATR) Kinase. <i>Journal of Biological Chemistry</i> , 2011, 286, 28707-28714.	1.6	101
51	Identification of Genome Maintenance Proteins with Functional Genomic Screens. <i>FASEB Journal</i> , 2010, 24, 304.1.	0.2	0
52	The annealing helicase SMARCAL1 maintains genome integrity at stalled replication forks. <i>Genes and Development</i> , 2009, 23, 2405-2414.	2.7	216
53	Common mechanisms of PIKK regulation. <i>DNA Repair</i> , 2009, 8, 1004-1008.	1.3	233
54	ATR: an essential regulator of genome integrity. <i>Nature Reviews Molecular Cell Biology</i> , 2008, 9, 616-627.	16.1	1,497

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55	Dpb11 activates the Mec1â€“Ddc2 complex. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 18730-18734.	3.3	110
56	TopBP1 activates ATR through ATRIP and a PIKK regulatory domain. Genes and Development, 2008, 22, 1478-1489.	2.7	293
57	Function of the ATR N-terminal domain revealed by an ATM/ATR chimera. Experimental Cell Research, 2007, 313, 1667-1674.	1.2	15
58	ATRIP Oligomerization Is Required for ATR-dependent Checkpoint Signaling. Journal of Biological Chemistry, 2005, 280, 31390-31396.	1.6	69
59	Unwind and slow down: checkpoint activation by helicase and polymerase uncoupling. Genes and Development, 2005, 19, 1007-1012.	2.7	76
60	From The Cover: Minichromosome maintenance proteins are direct targets of the ATM and ATR checkpoint kinases. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 10078-10083.	3.3	295
61	Caffeine Inhibits Checkpoint Responses without Inhibiting the Ataxia-Telangiectasia-mutated (ATM) and ATM- and Rad3-related (ATR) Protein Kinases. Journal of Biological Chemistry, 2003, 278, 37139-37145.	1.6	132
62	ATR and ATRIP: Partners in Checkpoint Signaling. Science, 2001, 294, 1713-1716.	6.0	867
63	Conducting the mitotic symphony. Nature, 2000, 406, 354-356.	13.7	28
64	Requirement of ATM-Dependent Phosphorylation of Brca1 in the DNA Damage Response to Double-Strand Breaks. Science, 1999, 286, 1162-1166.	6.0	941