

Paula E Cohen

List of Publications by Year in descending order

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57
papers

5,915
citations

101543

36
h-index

155660

55
g-index

64
all docs

64
docs citations

64
times ranked

5563
citing authors

#	ARTICLE	IF	CITATIONS
1	Meiotic Pachytene Arrest in MLH1-Deficient Mice. <i>Cell</i> , 1996, 85, 1125-1134.	28.9	528
2	Mammalian MutS homologue 5 is required for chromosome pairing in meiosis. <i>Nature Genetics</i> , 1999, 21, 123-127.	21.4	367
3	Meiotic arrest and aneuploidy in MLH3-deficient mice. <i>Nature Genetics</i> , 2002, 31, 385-390.	21.4	332
4	Mutation in the Mismatch Repair Gene Msh6 Causes Cancer Susceptibility. <i>Cell</i> , 1997, 91, 467-477.	28.9	326
5	Control of Meiotic Crossovers: From Double-Strand Break Formation to Designation. <i>Annual Review of Genetics</i> , 2016, 50, 175-210.	7.6	311
6	Inactivation of Exonuclease 1 in mice results in DNA mismatch repair defects, increased cancer susceptibility, and male and female sterility. <i>Genes and Development</i> , 2003, 17, 603-614.	5.9	282
7	The time course and chromosomal localization of recombination-related proteins at meiosis in the mouse are compatible with models that can resolve the early DNA-DNA interactions without reciprocal recombination. <i>Journal of Cell Science</i> , 2002, 115, 1611-22.	2.0	282
8	RNF212 is a dosage-sensitive regulator of crossing-over during mammalian meiosis. <i>Nature Genetics</i> , 2013, 45, 269-278.	21.4	231
9	Essential Role of Fkbp6 in Male Fertility and Homologous Chromosome Pairing in Meiosis. <i>Science</i> , 2003, 300, 1291-1295.	12.6	200
10	Genetic Recombination Is Targeted towards Gene Promoter Regions in Dogs. <i>PLoS Genetics</i> , 2013, 9, e1003984.	3.5	198
11	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
12	Antagonistic roles of ubiquitin ligase HEI10 and SUMO ligase RNF212 regulate meiotic recombination. <i>Nature Genetics</i> , 2014, 46, 194-199.	21.4	172
13	MUS81 Generates a Subset of MLH1-MLH3-Independent Crossovers in Mammalian Meiosis. <i>PLoS Genetics</i> , 2008, 4, e1000186.	3.5	164
14	Extreme Heterogeneity in the Molecular Events Leading to the Establishment of Chiasmata during Meiosis I in Human Oocytes. <i>American Journal of Human Genetics</i> , 2005, 76, 112-127.	6.2	151
15	Absence of Colony-Stimulating Factor-1 in Osteopetrotic (csfmoP/csfmOP) Mice Results in Male Fertility Defects1. <i>Biology of Reproduction</i> , 1996, 55, 310-317.	2.7	132
16	Macrophages: important accessory cells for reproductive function. <i>Journal of Leukocyte Biology</i> , 1999, 66, 765-772.	3.3	128
17	Absence of Colony Stimulating Factor-1 in Osteopetrotic (csfmop/csfmop) Mice Disrupts Estrous Cycles and Ovulation1. <i>Biology of Reproduction</i> , 1997, 56, 110-118.	2.7	127
18	Localization of MMR proteins on meiotic chromosomes in mice indicates distinct functions during prophase I. <i>Journal of Cell Biology</i> , 2005, 171, 447-458.	5.2	123

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19	Regulation of meiotic recombination and prophase I progression in mammals. <i>BioEssays</i> , 2001, 23, 996-1009.	2.5	105
20	Control of Spermatogenesis in Mice by the Cyclin D-Dependent Kinase Inhibitors p18 Ink4c and p19 Ink4d. <i>Molecular and Cellular Biology</i> , 2001, 21, 3244-3255.	2.3	103
21	Ubiquitylated PCNA plays a role in somatic hypermutation and class-switch recombination and is required for meiotic progression. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 16248-16253.	7.1	99
22	Chromosome Cohesion Established by Rec8-Cohesin in Fetal Oocytes Is Maintained without Detectable Turnover in Oocytes Arrested for Months in Mice. <i>Current Biology</i> , 2016, 26, 678-685.	3.9	92
23	AGO4 Regulates Entry into Meiosis and Influences Silencing of Sex Chromosomes in the Male Mouse Germline. <i>Developmental Cell</i> , 2012, 23, 251-264.	7.0	88
24	Colony-Stimulating Factor-1 Plays a Major Role in the Development of Reproductive Function in Male Mice. <i>Molecular Endocrinology</i> , 1997, 11, 1636-1650.	3.7	81
25	Mammalian CNTD1 is critical for meiotic crossover maturation and deselection of excess precrossover sites. <i>Journal of Cell Biology</i> , 2014, 205, 633-641.	5.2	80
26	Dynamic transcriptome profiles within spermatogonial and spermatocyte populations during postnatal testis maturation revealed by single-cell sequencing. <i>PLoS Genetics</i> , 2019, 15, e1007810.	3.5	80
27	Mammalian BLM helicase is critical for integrating multiple pathways of meiotic recombination. <i>Journal of Cell Biology</i> , 2010, 188, 779-789.	5.2	75
28	SFMBT1 functions with LSD1 to regulate expression of canonical histone genes and chromatin-related factors. <i>Genes and Development</i> , 2013, 27, 749-766.	5.9	73
29	Reproduction in Men with Klinefelter Syndrome: The Past, the Present, and the Future. <i>Seminars in Reproductive Medicine</i> , 2009, 27, 137-148.	1.1	71
30	Mismatch repair proteins, meiosis, and mice: understanding the complexities of mammalian meiosis. <i>Experimental Cell Research</i> , 2004, 296, 71-79.	2.6	66
31	Mammalian BTBD12 (SLX4) Protects against Genomic Instability during Mammalian Spermatogenesis. <i>PLoS Genetics</i> , 2011, 7, e1002094.	3.5	65
32	Distinct Functions of MLH3 at Recombination Hot Spots in the Mouse. <i>Genetics</i> , 2008, 178, 1937-1945.	2.9	56
33	Effect of the Colony-Stimulating Factor-1 Null Mutation, Osteopetrotic (csfmoP), on the Distribution of Macrophages in the Male Mouse Reproductive Tract ¹ . <i>Biology of Reproduction</i> , 1997, 56, 1290-1300.	2.7	55
34	Conditional Inactivation of the DNA Damage Response Gene Hus1 in Mouse Testis Reveals Separable Roles for Components of the RAD9-RAD1-HUS1 Complex in Meiotic Chromosome Maintenance. <i>PLoS Genetics</i> , 2013, 9, e1003320.	3.5	48
35	<i>Dgcr8</i> and <i>Dicer</i> are essential for sex chromosome integrity during meiosis in males. <i>Journal of Cell Science</i> , 2015, 128, 2314-2327.	2.0	47
36	Comparative Analysis of Meiotic Progression in Female Mice Bearing Mutations in Genes of the DNA Mismatch Repair Pathway ¹ . <i>Biology of Reproduction</i> , 2008, 78, 462-471.	2.7	40

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37	Distinct effects of the recurrent Mlh1 ^{G67R} mutation on MMR functions, cancer, and meiosis. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 4247-4252.	7.1	39
38	Fancj (Brip1) loss-of-function allele results in spermatogonial cell depletion during embryogenesis and altered processing of crossover sites during meiotic prophase I in mice. Chromosoma, 2016, 125, 237-252.	2.2	33
39	Cohesin Removal along the Chromosome Arms during the First Meiotic Division Depends on a NEK1-PP1 ³ -WAPL Axis in the Mouse. Cell Reports, 2016, 17, 977-986.	6.4	31
40	A mutation in the endonuclease domain of mouse MLH3 reveals novel roles for MutL ³ during crossover formation in meiotic prophase I. PLoS Genetics, 2019, 15, e1008177.	3.5	25
41	Analysis of meiotic prophase I in live mouse spermatocytes. Chromosome Research, 2008, 16, 743-760.	2.2	24
42	A role for Mlh3 in somatic hypermutation. DNA Repair, 2006, 5, 675-682.	2.8	22
43	Transcriptome profiling of the developing male germ line identifies the miR-29 family as a global regulator during meiosis. RNA Biology, 2017, 14, 219-235.	3.1	21
44	Predicting Gene Networks in Human Oocyte Meiosis. Biology of Reproduction, 2010, 82, 469-472.	2.7	20
45	Cyclin N-Terminal Domain-Containing-1 Coordinates Meiotic Crossover Formation with Cell-Cycle Progression in a Cyclin-Independent Manner. Cell Reports, 2020, 32, 107858.	6.4	20
46	Somatic CAG expansion in Huntington's disease is dependent on the MLH3 endonuclease domain, which can be excluded via splice redirection. Nucleic Acids Research, 2021, 49, 3907-3918.	14.5	20
47	Mutation of the ATPase Domain of MutS Homolog-5 (MSH5) Reveals a Requirement for a Functional MutS ³ Complex for All Crossovers in Mammalian Meiosis. G3: Genes, Genomes, Genetics, 2019, 9, 1839-1850.	1.8	19
48	NIMA-related kinase 1 (NEK1) regulates meiosis I spindle assembly by altering the balance between β -Adducin and Myosin X. PLoS ONE, 2017, 12, e0185780.	2.5	16
49	NEK1 Facilitates Cohesin Removal during Mammalian Spermatogenesis. Genes, 2011, 2, 260-279.	2.4	15
50	Phosphoproteomics of ATR signaling in mouse testes. ELife, 2022, 11, .	6.0	12
51	Small RNAs and RNAi pathways in meiotic prophase I. Chromosome Research, 2007, 15, 653-665.	2.2	11
52	Double trouble in human aneuploidy. Nature Genetics, 2015, 47, 696-698.	21.4	10
53	Characterization of telomeric repeat-containing RNA (TERRA) localization and protein interactions in primordial germ cells of the mouse. Biology of Reproduction, 2019, 100, 950-962.	2.7	9
54	Multiple 9-1-1 complexes promote homolog synapsis, DSB repair, and ATR signaling during mammalian meiosis. ELife, 2022, 11, .	6.0	7

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55	Special issue on "Recent advances in meiotic chromosome structure, recombination and segregation". Chromosoma, 2016, 125, 173-175.	2.2	3
56	Non-human primates exhibit errors in meiosis I. Molecular Reproduction and Development, 2012, 79, 665-665.	2.0	0
57	Gene networks in oocyte meiosis. , 0, , 24-37.		0