Penelope A Jeggo

List of Publications by Year
in descending order

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145 papers	19,867 citations	11608 70 h-index	10708 138 g-index
147 all docs	147 docs citations	147 times ranked	17097 citing authors

#	Article	IF	Citations
1	Editorial for special issue â€˜women in radiobiologyâ€™. International Journal of Radiation Biology, 2022, , 1-1.	1.0	0
2	Haematopoietic Stem Cell Transplantation for DNA Ligase 1 Deficiency. Journal of Clinical Immunology, 2021, 41, 238-242.	2.0	2
3	Cutting-edge perspectives in genomic maintenance V11. DNA Repair, 2021, 97, 103004.	1.3	0
4	TP53 modulates radiotherapy fraction size sensitivity in normal and malignant cells. Scientific Reports, 2021, 11, 7119.	1.6	11
5	ATMâ€™s Role in the Repair of DNA Double-Strand Breaks. Genes, 2021, 12, 1370.	1.0	38
6	Establishing mechanisms affecting the individual response to ionizing radiation. International Journal of Radiation Biology, 2020, 96, 297-323.	1.0	34
7	Roles for 53BP1 in the repair of radiation-induced DNA double strand breaks. DNA Repair, 2020, 93, 102915.	1.3	61
8	Roles for the DNA-PK complex and 53BP1 in protecting ends from resection during DNA double-strand break repair. Journal of Radiation Research, 2020, 61, 718-726.	0.8	17
9	DNA double-strand break end resection: a critical relay point for determining the pathway of repair and signaling. Genome Instability & Disease, 2020, 1, 155-171.	0.5	18
10	Advances in Radiation Biology â€“ Highlights from the 16th ICRR special feature: introductory editorial. British Journal of Radiology, 2020, 93, 20209006.	1.0	1
11	Canonical DNA non-homologous end-joining; capacity versus fidelity. British Journal of Radiology, 2020, 93, 20190966.	1.0	24
12	Regulation of programmed deathâ€ligand 1 expression in response to <scp>DNA</scp> damage in cancer cells: Implications for precision medicine. Cancer Science, 2019, 110, 3415-3423.	1.7	42
13	Distinct response of adult neural stem cells to low versus high dose ionising radiation. DNA Repair, 2019, 76, 70-75.	1.3	15
14	Analysis of cilia dysfunction phenotypes in zebrafish embryos depleted of Origin recognition complex factors. European Journal of Human Genetics, 2019, 27, 772-782.	1.4	12
15	The Antiresection Activity of the X Protein Encoded by Hepatitis Virus B. Hepatology, 2019, 69, 2546-2561.	3.6	20
16	Repression of Transcription at DNA Breaks Requires Cohesin throughout Interphase and Prevents Genome Instability. Molecular Cell, 2019, 73, 212-223.e7.	4.5	83
17	A historical reflection on our understanding of radiation-induced DNA double strand break repair in somatic mammalian cells; interfacing the past with the present. International Journal of Radiation Biology, 2019, 95, 945-956.	1.0	31
18	Resting cells rely on the DNA helicase component MCM2 to build cilia. Nucleic Acids Research, 2019, 47, 134-151.	6.5	29

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19	Hazards of human spaceflight. Science, 2019, 364, 127-128.	6.0	46
20	ATM: Its Recruitment, Activation, Signalling and Contribution to Tumour Suppression. Cancer Drug Discovery and Development, 2018, , 129-154.	0.2	0
21	The pendulum of the Ku-Ku clock. DNA Repair, 2018, 71, 164-171.	1.3	52
22	Novel function of HATs and HDACs in homologous recombination through acetylation of human RAD52 at double-strand break sites. PLoS Genetics, 2018, 14, e1007277.	1.5	25
23	Ionizing radiation biomarkers in epidemiological studies â€“ An update. Mutation Research - Reviews in Mutation Research, 2017, 771, 59-84.	2.4	118
24	DNA Double-Strand Break Resection Occurs during Non-homologous End Joining in G1 but Is Distinct from Resection during Homologous Recombination. Molecular Cell, 2017, 65, 671-684.e5.	4.5	184
25	Chromatin modifiers and remodellers in DNA repair and signalling. Philosophical Transactions of the Royal Society B: Biological Sciences, 2017, 372, 20160279.	1.8	18
26	A restatement of the natural science evidence base concerning the health effects of low-level ionizing radiation. Proceedings of the Royal Society B: Biological Sciences, 2017, 284, 20171070.	1.2	68
27	A Process of Resection-Dependent Nonhomologous End Joining Involving the Goddess Artemis. Trends in Biochemical Sciences, 2017, 42, 690-701.	3.7	86
28	DNA non-homologous end-joining enters the resection arena. Oncotarget, 2017, 8, 93317-93318.	0.8	5
29	A coordinated DNA damage response promotes adult quiescent neural stem cell activation. PLoS Biology, 2017, 15, e2001264.	2.6	36
30	In vivosensitivity of the embryonic and adult neural stem cell compartments to low-dose radiation. Journal of Radiation Research, 2016, 57, i2-i10.	0.8	18
31	DNA repair, genome stability and cancer: a historical perspective. Nature Reviews Cancer, 2016, 16, 35-42.	12.8	575
32	ATR promotes cilia signalling: links to developmental impacts. Human Molecular Genetics, 2016, 25, 1574-1587.	1.4	24
33	ATM Localization and Heterochromatin Repair Depend on Direct Interaction of the 53BP1-BRCT 2 Domain with Î³H2AX. Cell Reports, 2015, 13, 2081-2089.	2.9	61
34	Endogenous and X-ray-induced DNA double strand breaks sensitively activate apoptosis in adult neural stem cells. Journal of Cell Science, 2015, 128, 3597-606.	1.2	24
35	SETDB1, HP1 and SUV39 promote repositioning of 53BP1 to extend resection during homologous recombination in G2 cells. Nucleic Acids Research, 2015, 43, 7931-7944.	6.5	69
36	How cancer cells hijack DNA double-strand break repair pathways to gain genomic instability. Biochemical Journal, 2015, 471, 1-11.	1.7	81

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37	XRCC4 deficiency in human subjects causes a marked neurological phenotype but no overt immunodeficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 1007-1017.	1.5	44
38	The rate of X-ray-induced DNA double-strand break repair in the embryonic mouse brain is unaffected by exposure to 50 Hz magnetic fields. International Journal of Radiation Biology, 2015, 91, 495-499.	1.0	9
39	Evaluation of Severe Combined Immunodeficiency and Combined Immunodeficiency Pediatric Patients on the Basis of Cellular Radiosensitivity. Journal of Molecular Diagnostics, 2015, 17, 560-575.	1.2	16
40	The PBAF chromatin remodeling complex represses transcription and promotes rapid repair at DNA double-strand breaks. Molecular and Cellular Oncology, 2015, 2, e970072.	0.3	13
41	Low levels of endogenous or X-ray-induced DNA double-strand breaks activate apoptosis in adult neural stem cells. Development (Cambridge), 2015, 142, e1.2-e1.2.	1.2	0
42	Requirement for PBAF in Transcriptional Repression and Repair at DNA Breaks in Actively Transcribed Regions of Chromatin. Molecular Cell, 2014, 55, 723-732.	4.5	230
43	Increased apoptosis and DNA double-strand breaks in the embryonic mouse brain in response to very low-dose X-rays but not 50 Hz magnetic fields. Journal of the Royal Society Interface, 2014, 11, 20140783.	1.5	35
44	DNA Double-strand Break Repair in a Cellular Context. Clinical Oncology, 2014, 26, 243-249.	0.6	126
45	DNA Double-Strand Break Repair Pathway Choice Is Directed by Distinct MRE11 Nuclease Activities. Molecular Cell, 2014, 53, 7-18.	4.5	466
46	Roles of chromatin remodellers in DNA double strand break repair. Experimental Cell Research, 2014, 329, 69-77.	1.2	58
47	Polo-like kinase 3 regulates CtIP during DNA double-strand break repair in G1. Journal of Cell Biology, 2014, 206, 877-894.	2.3	92
48	Robin Holliday 1932â€“2014. Nature Structural and Molecular Biology, 2014, 21, 501-502.	3.6	1
49	The clinical impact of deficiency in DNA non-homologous end-joining. DNA Repair, 2014, 16, 84-96.	1.3	138
50	Reprint of â€œThe clinical impact of deficiency in DNA non-homologous end-joiningâ€• DNA Repair, 2014, 17, 9-20.	1.3	11
51	Statistical analysis of kinetics, distribution and co-localisation of DNA repair foci in irradiated cells: Cell cycle effect and implications for prediction of radiosensitivity. DNA Repair, 2013, 12, 844-855.	1.3	40
52	The many faces of Artemis-deficient combined immunodeficiency â€” Two patients with DCLRE1C mutations and a systematic literature review of genotypeâ€“phenotype correlation. Clinical Immunology, 2013, 149, 464-474.	1.4	39
53	The complexity of DNA double strand breaks is a critical factor enhancing end-resection. DNA Repair, 2013, 12, 936-946.	1.3	71
54	Brief report: A human induced pluripotent stem cell model of cernunnos deficiency reveals an important role for XLF in the survival of the primitive hematopoietic progenitors. Stem Cells, 2013, 31, 2015-2023.	1.4	15

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55	The Repair and Signaling Responses to DNA Double-Strand Breaks. Advances in Genetics, 2013, 82, 1-45.	0.8	186
56	Diminished Origin-Licensing Capacity Specifically Sensitizes Tumor Cells to Replication Stress. Molecular Cancer Research, 2013, 11, 370-380.	1.5	38
57	Deficiency in Origin Licensing Proteins Impairs Cilia Formation: Implications for the Aetiology of Meier-Gorlin Syndrome. PLoS Genetics, 2013, 9, e1003360.	1.5	55
58	A noncatalytic function of the ligation complex during nonhomologous end joining. Journal of Cell Biology, 2013, 200, 173-186.	2.3	81
59	Opposing roles for 53BP1 during homologous recombination. Nucleic Acids Research, 2013, 41, 9719-9731.	6.5	74
60	Co-operation of BRCA1 and POH1 relieves the barriers posed by 53BP1 and RAP80 to resection. Nucleic Acids Research, 2013, 41, 10298-10311.	6.5	99
61	PRKDC mutations in a SCID patient with profound neurological abnormalities. Journal of Clinical Investigation, 2013, 123, 2969-2980.	3.9	121
62	Visualisation of Î³H2AX Foci Caused by Heavy Ion Particle Traversal; Distinction between Core Track versus Non-Track Damage. PLoS ONE, 2013, 8, e70107.	1.1	68
63	Identification of the First ATRIPâ€“Deficient Patient and Novel Mutations in ATR Define a Clinical Spectrum for ATRâ€“ATRIP Seckel Syndrome. PLoS Genetics, 2012, 8, e1002945.	1.5	104
64	Phosphoproteomic analysis reveals that PP4 dephosphorylates KAP-1 impacting the DNA damage response. EMBO Journal, 2012, 31, 2403-2415.	3.5	96
65	The Heterochromatic Barrier to DNA Double Strand Break Repair: How to Get the Entry Visa. International Journal of Molecular Sciences, 2012, 13, 11844-11860.	1.8	92
66	Irradiation induced foci (IRIF) as a biomarker for radiosensitivity. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2012, 736, 39-47.	0.4	74
67	Ionizing radiation biomarkers for potential use in epidemiological studies. Mutation Research - Reviews in Mutation Research, 2012, 751, 258-286.	2.4	181
68	DNA double-strand break repair within heterochromatic regions. Biochemical Society Transactions, 2012, 40, 173-178.	1.6	33
69	ASPM influences DNA double-strand break repair and represents a potential target for radiotherapy. International Journal of Radiation Biology, 2011, 87, 1189-1195.	1.0	42
70	Understanding the limitations of radiation-induced cell cycle checkpoints. Critical Reviews in Biochemistry and Molecular Biology, 2011, 46, 271-283.	2.3	166
71	The role of homologous recombination in radiation-induced double-strand break repair. Radiotherapy and Oncology, 2011, 101, 7-12.	0.3	161
72	Mutations in ORC1, encoding the largest subunit of the origin recognition complex, cause microcephalic primordial dwarfism resembling Meier-Gorlin syndrome. Nature Genetics, 2011, 43, 350-355.	9.4	189

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73	Factors determining DNA double-strand break repair pathway choice in G2 phase. EMBO Journal, 2011, 30, 1079-1092.	3.5	381
74	KAP-1 phosphorylation regulates CHD3 nucleosome remodeling during the DNA double-strand break response. Nature Structural and Molecular Biology, 2011, 18, 831-839.	3.6	205
75	Requirement for DNA Ligase IV during Embryonic Neuronal Development. Journal of Neuroscience, 2011, 31, 10088-10100.	1.7	57
76	Analysis of Human Syndromes with Disordered Chromatin Reveals the Impact of Heterochromatin on the Efficacy of ATM-Dependent G₂/M Checkpoint Arrest. Molecular and Cellular Biology, 2011, 31, 4022-4035.	1.1	32
77	Checkpoint Control Following Radiation Exposure. , 2011, , 53-77.		0
78	An Artemis polymorphic variant reduces Artemis activity and confers cellular radiosensitivity. DNA Repair, 2010, 9, 1003-1010.	1.3	33
79	The influence of heterochromatin on DNA double strand break repair: Getting the strong, silent type to relax. DNA Repair, 2010, 9, 1273-1282.	1.3	269
80	A break is not the End; insight into the damage response to DNA double strand breaks. DNA Repair, 2010, 9, 1217-1218.	1.3	4
81	53BP1 promotes ATM activity through direct interactions with the MRN complex. EMBO Journal, 2010, 29, 574-585.	3.5	105
82	53BP1-dependent robust localized KAP-1 phosphorylation is essential for heterochromatic DNA double-strand break repair. Nature Cell Biology, 2010, 12, 177-184.	4.6	289
83	Sensitization to Radiation and Alkylating Agents by Inhibitors of Poly(ADP-ribose) Polymerase Is Enhanced in Cells Deficient in DNA Double-Strand Break Repair. Molecular Cancer Therapeutics, 2010, 9, 1775-1787.	1.9	118
84	Role of ATM and the Damage Response Mediator Proteins 53BP1 and MDC1 in the Maintenance of G₂/M Checkpoint Arrest. Molecular and Cellular Biology, 2010, 30, 3371-3383.	1.1	97
85	The Limitations of the G1-S Checkpoint. Cancer Research, 2010, 70, 4412-4421.	0.4	70
86	Î³H2AX foci analysis for monitoring DNA double-strand break repair: Strengths, limitations and optimization. Cell Cycle, 2010, 9, 662-669.	1.3	545
87	The Role of the DNA Damage Response Mechanisms after Low-Dose Radiation Exposure and a Consideration of Potentially Sensitive Individuals. Radiation Research, 2010, 174, 825-832.	0.7	35
88	Translesion DNA synthesis-assisted non-homologous end-joining of complex double-strand breaks prevents loss of DNA sequences in mammalian cells. Nucleic Acids Research, 2009, 37, 6737-6745.	6.5	25
89	XLF-Cernunnos promotes DNA ligase IVâ€“XRCC4 re-adenylation following ligation. Nucleic Acids Research, 2009, 37, 482-492.	6.5	98
90	ATM and Artemis promote homologous recombination of radiation-induced DNA double-strand breaks in G2. EMBO Journal, 2009, 28, 3413-3427.	3.5	457

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91	Cellular radiosensitivity: How much better do we understand it?. International Journal of Radiation Biology, 2009, 85, 1061-1081.	1.0	148
92	The impact of heterochromatin on DSB repair. Biochemical Society Transactions, 2009, 37, 569-576.	1.6	138
93	RISKS FROM LOW DOSE/DOSE RATE RADIATION: WHAT AN UNDERSTANDING OF DNA DAMAGE RESPONSE MECHANISMS CAN TELL US. Health Physics, 2009, 97, 416-425.	0.3	23
94	Impaired lymphocyte development and antibody class switching and increased malignancy in a murine model of DNA ligase IV syndrome. Journal of Clinical Investigation, 2009, 119, 1696-1705.	3.9	33
95	'A mover and a shaker': 53BP1 allows DNA doublestrand breaks a chance to dance and unite. F1000 Biology Reports, 2009, 1, 21.	4.0	7
96	Mutations in pericentrin cause Seckel syndrome with defective ATR-dependent DNA damage signaling. Nature Genetics, 2008, 40, 232-236.	9.4	281
97	The role of the DNA damage response pathways in brain development and microcephaly: Insight from human disorders. DNA Repair, 2008, 7, 1039-1050.	1.3	73
98	ATM Signaling Facilitates Repair of DNA Double-Strand Breaks Associated with Heterochromatin. Molecular Cell, 2008, 31, 167-177.	4.5	777
99	Replication independent ATR signalling leads to G2/M arrest requiring Nbs1, 53BP1 and MDC1. Human Molecular Genetics, 2008, 17, 3247-3253.	1.4	33
100	Chromosome breakage after G2 checkpoint release. Journal of Cell Biology, 2007, 176, 749-755.	2.3	220
101	Interaction of the Ku heterodimer with the DNA ligase IV/Xrcc4 complex and its regulation by DNA-PK. DNA Repair, 2007, 6, 712-722.	1.3	112
102	Cellular and Clinical Impact of Haploinsufficiency for Genes Involved in ATR Signaling. American Journal of Human Genetics, 2007, 81, 77-86.	2.6	68
103	An Imperfect G₂M Checkpoint Contributes to Chromosome Instability Following Irradiation of S and G₂Phase Cells. Cell Cycle, 2007, 6, 1682-1686.	1.3	93
104	DNA double-strand breaks: their cellular and clinical impact?. Oncogene, 2007, 26, 7717-7719.	2.6	226
105	The impact of a negligent G2/M checkpoint on genomic instability and cancer induction. Nature Reviews Cancer, 2007, 7, 861-869.	12.8	514
106	DNA repair is limiting for haematopoietic stem cells during ageing. Nature, 2007, 447, 686-690.	13.7	475
107	X-irradiation of cells on glass slides has a dose doubling impact. DNA Repair, 2007, 6, 1692-1697.	1.3	47
108	Identification of a novel motif in DNA ligases exemplified by DNA ligase IV. DNA Repair, 2006, 5, 788-798.	1.3	19

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109	Contribution of DNA repair and cell cycle checkpoint arrest to the maintenance of genomic stability. DNA Repair, 2006, 5, 1192-1198.	1.3	138
110	Radiation-induced DNA damage responses. Radiation Protection Dosimetry, 2006, 122, 124-127.	0.4	118
111	Regulation of mitotic entry by microcephalin and its overlap with ATR signalling. Nature Cell Biology, 2006, 8, 725-733.	4.6	164
112	The role of double-strand break repair â€" insights from human genetics. Nature Reviews Genetics, 2006, 7, 45-54.	7.7	514
113	DNA-PK autophosphorylation facilitates Artemis endonuclease activity. EMBO Journal, 2006, 25, 3880-3889.	3.5	281
114	ATR-dependent phosphorylation and activation of ATM in response to UV treatment or replication fork stalling. EMBO Journal, 2006, 25, 5775-5782.	3.5	319
115	Microcephalin: A Causal Link Between Impaired Damage Response Signalling and Microcephaly. Cell Cycle, 2006, 5, 2339-2344.	1.3	44
116	Radiation-induced delayed cell death in a hypomorphic Artemis cell line. Human Molecular Genetics, 2006, 15, 1303-1311.	1.4	35
117	Nbs1 is required for ATR-dependent phosphorylation events. EMBO Journal, 2005, 24, 199-208.	3.5	160
118	The life and death of DNA-PK. Oncogene, 2005, 24, 949-961.	2.6	400
119	Artemis links ATM to Double Strand Break Rejoining. Cell Cycle, 2005, 4, 359-362.	1.3	95
120	Phosphorylation of linker histones by DNA-dependent protein kinase is required for DNA ligase IV-dependent ligation in the presence of histone H1. Proceedings of the National Academy of Sciences of the United States of America, 2005, 102, 1877-1882.	3.3	49
121	The two edges of the ATM sword: Co-operation between repair and checkpoint functions. Radiotherapy and Oncology, 2005, 76, 112-118.	0.3	72
122	Harmonising the response to DSBs: a new string in the ATM bow. DNA Repair, 2005, 4, 749-759.	1.3	91
123	Genomic Instability in Cancer Development. , 2005, 570, 175-197.		6
124	Analysis of DNA ligase IV mutations found in LIG4 syndrome patients: the impact of two linked polymorphisms. Human Molecular Genetics, 2004, 13, 2369-2376.	1.4	114
125	Seckel syndrome exhibits cellular features demonstrating defects in the ATR-signalling pathway. Human Molecular Genetics, 2004, 13, 3127-3138.	1.4	155
126	A Double-Strand Break Repair Defect in ATM-Deficient Cells Contributes to Radiosensitivity. Cancer Research, 2004, 64, 500-508.	0.4	328

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127	ATM and DNA-PK Function Redundantly to Phosphorylate H2AX after Exposure to Ionizing Radiation. Cancer Research, 2004, 64, 2390-2396.	0.4	896
128	AHNAK interacts with the DNA ligase IVâ€“XRCC4 complex and stimulates DNA ligase IV-mediated double-stranded ligation. DNA Repair, 2004, 3, 245-256.	1.3	16
129	A Pathway of Double-Strand Break Rejoining Dependent upon ATM, Artemis, and Proteins Locating to Î³-H2AX Foci. Molecular Cell, 2004, 16, 715-724.	4.5	790
130	Healing the Wounds Inflicted by Sleeping Beauty Transposition by Double-Strand Break Repair in Mammalian Somatic Cells. Molecular Cell, 2004, 13, 279-290.	4.5	108
131	A splicing mutation affecting expression of ataxiaâ€“telangiectasia and Rad3â€“related protein (ATR) results in Seckel syndrome. Nature Genetics, 2003, 33, 497-501.	9.4	699
132	Ku Stimulation of DNA Ligase IV-dependent Ligation Requires Inward Movement along the DNA Molecule. Journal of Biological Chemistry, 2003, 278, 22466-22474.	1.6	69
133	Chk2 Is a Tumor Suppressor That Regulates Apoptosis in both an Ataxia Telangiectasia Mutated (ATM)-Dependent and an ATM-Independent Manner. Molecular and Cellular Biology, 2002, 22, 6521-6532.	1.1	354
134	The Greek Goddess, Artemis, reveals the secrets of her cleavage. DNA Repair, 2002, 1, 771-777.	1.3	41
135	Immunological disorders and DNA repair. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2002, 509, 109-126.	0.4	25
136	Nbs1 promotes ATM dependent phosphorylation events including those required for G1/S arrest. Oncogene, 2002, 21, 4191-4199.	2.6	113
137	DNA Ligase IV Mutations Identified in Patients Exhibiting Developmental Delay and Immunodeficiency. Molecular Cell, 2001, 8, 1175-1185.	4.5	497
138	Immune diversity and genomic stability: opposite goals but similar paths. Journal of Photochemistry and Photobiology B: Biology, 2001, 65, 88-96.	1.7	14
139	Cellular and Biochemical Impact of a Mutation in DNA Ligase IV Conferring Clinical Radiosensitivity. Journal of Biological Chemistry, 2001, 276, 31124-31132.	1.6	116
140	Splitting the ATM: distinct repair and checkpoint defects in ataxiaâ€“telangiectasia. Trends in Genetics, 1998, 14, 312-316.	2.9	154
141	Targeted Disruption of the Catalytic Subunit of the DNA-PK Gene in Mice Confers Severe Combined Immunodeficiency and Radiosensitivity. Immunity, 1998, 9, 355-366.	6.6	301
142	Menage Ã¡ trois: Double strand break repair, V(D)J recombination and DNA-PK. BioEssays, 1995, 17, 949-957.	1.2	237
143	Nomenclature of human genes involved in ionizing radiation sensitivity. Mutation Research DNA Repair, 1995, 337, 131-134.	3.8	69
144	Defective DNA-dependent protein kinase activity is linked to V(D)J recombination and DNA repair defects associated with the murine scid mutation. Cell, 1995, 80, 813-823.	13.5	809

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145	Studies on mammalian mutants defective in rejoining double-strand breaks in DNA. Mutation Research - Reviews in Genetic Toxicology, 1990, 239, 1-16.	3.0	165