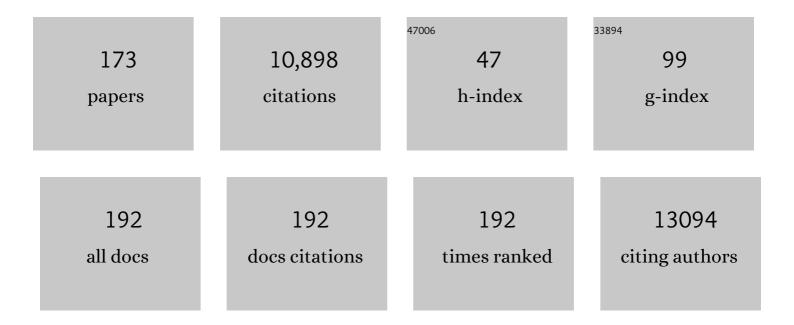
List of Publications by Year in descending order

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ΜΑΟΝΑΡ ΒΙΑ ΡΑΥς

#	Article	IF	CITATIONS
1	Cloning and expression of a rat brain L-glutamate transporter. Nature, 1992, 360, 464-467.	27.8	1,197
2	Base Excision Repair. Cold Spring Harbor Perspectives in Biology, 2013, 5, a012583-a012583.	5.5	908
3	Human and bacterial oxidative demethylases repair alkylation damage in both RNA and DNA. Nature, 2003, 421, 859-863.	27.8	558
4	The base excision repair pathway. Trends in Biochemical Sciences, 1995, 20, 391-397.	7.5	492
5	Broad histone H3K4me3 domains in mouse oocytes modulate maternal-to-zygotic transition. Nature, 2016, 537, 548-552.	27.8	484
6	OGG1 initiates age-dependent CAG trinucleotide expansion in somatic cells. Nature, 2007, 447, 447-452.	27.8	392
7	Exercise induces cerebral VEGF and angiogenesis via the lactate receptor HCAR1. Nature Communications, 2017, 8, 15557.	12.8	321
8	Human DNA glycosylases of the bacterial Fpg/MutM superfamily: an alternative pathway for the repair of 8-oxoguanine and other oxidation products in DNA. Nucleic Acids Research, 2002, 30, 4926-4936.	14.5	245
9	Mitochondrial DNA Integrity Is Essential For Mitochondrial Maturation During Differentiation of Neural Stem Cells. Stem Cells, 2010, 28, 2195-2204.	3.2	228
10	New functions of XPC in the protection of human skin cells from oxidative damage. EMBO Journal, 2006, 25, 4305-4315.	7.8	227
11	Ythdf2-mediated m6A mRNA clearance modulates neural development in mice. Genome Biology, 2018, 19, 69.	8.8	216
12	Discovery and development of safe-in-man broad-spectrum antiviral agents. International Journal of Infectious Diseases, 2020, 93, 268-276.	3.3	169
13	Structure-based mutagenesis reveals the albumin-binding site of the neonatal Fc receptor. Nature Communications, 2012, 3, 610.	12.8	160
14	Human ABH3 structure and key residues for oxidative demethylation to reverse DNA/RNA damage. EMBO Journal, 2006, 25, 3389-3397.	7.8	157
15	Epitope-resolved profiling of the SARS-CoV-2 antibody response identifies cross-reactivity with endemic human coronaviruses. Cell Reports Medicine, 2021, 2, 100189.	6.5	149
16	PGM3 Mutations Cause a Congenital Disorder of Glycosylation with Severe Immunodeficiency and Skeletal Dysplasia. American Journal of Human Genetics, 2014, 95, 96-107.	6.2	148
17	The <i>Saccharomyces cerevisiae</i> Homologues of Endonuclease III from <i>Escherichia coli</i> , Ntg1 and Ntg2, Are Both Required for Efficient Repair of Spontaneous and Induced Oxidative DNA Damage in Yeast. Molecular and Cellular Biology, 1999, 19, 3779-3787.	2.3	144
18	DNA base repair – recognition and initiation of catalysis. FEMS Microbiology Reviews, 2009, 33, 1044-1078.	8.6	140

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19	Extending Serum Half-life of Albumin by Engineering Neonatal Fc Receptor (FcRn) Binding. Journal of Biological Chemistry, 2014, 289, 13492-13502.	3.4	132
20	Inosine in DNA and RNA. Current Opinion in Genetics and Development, 2014, 26, 116-123.	3.3	117
21	WRN Interacts Physically and Functionally with the Recombination Mediator Protein RAD52. Journal of Biological Chemistry, 2003, 278, 36476-36486.	3.4	105
22	Substrate specificities of bacterial and human AlkB proteins. Nucleic Acids Research, 2004, 32, 3456-3461.	14.5	104
23	Reciprocal "flipping―underlies substrate recognition and catalytic activation by the human 8-oxo-guanine DNA glycosylase. Journal of Molecular Biology, 2002, 317, 171-177.	4.2	101
24	Potential Antiviral Options against SARS-CoV-2 Infection. Viruses, 2020, 12, 642.	3.3	92
25	Excision of 3-methylguanine from alkylated DNA by 3-methyladenine DNA glycosylase I ofEscherichia coli. Nucleic Acids Research, 1993, 21, 2045-2049.	14.5	86
26	Hippocampal Adult Neurogenesis Is Maintained by Neil3-Dependent Repair of Oxidative DNA Lesions in Neural Progenitor Cells. Cell Reports, 2012, 2, 503-510.	6.4	84
27	Dynamic relocalization of hOGG1 during the cell cycle is disrupted in cells harbouring the hOGG1-Cys326 polymorphic variant. Nucleic Acids Research, 2005, 33, 1813-1824.	14.5	83
28	Structures of endonuclease V with DNA reveal initiation of deaminated adenine repair. Nature Structural and Molecular Biology, 2009, 16, 138-143.	8.2	83
29	Endonuclease VIII-like 3 (Neil3) DNA glycosylase promotes neurogenesis induced by hypoxia-ischemia. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 18802-18807.	7.1	83
30	Human NEIL3 is mainly a monofunctional DNA glycosylase removing spiroimindiohydantoin and guanidinohydantoin. DNA Repair, 2013, 12, 1159-1164.	2.8	80
31	Dissection of the Molecular Defects Caused by Pathogenic Mutations in the DNA Repair Factor XPC. Molecular and Cellular Biology, 2008, 28, 7225-7235.	2.3	79
32	Human OGG1 undergoes serine phosphorylation and associates with the nuclear matrix and mitotic chromatin in vivo. Nucleic Acids Research, 2002, 30, 2349-2357.	14.5	75
33	Endonuclease V cleaves at inosines in RNA. Nature Communications, 2013, 4, 2271.	12.8	71
34	Separation-of-Function Mutants Unravel the Dual-Reaction Mode of Human 8-Oxoguanine DNA Glycosylase. Structure, 2011, 19, 117-127.	3.3	70
35	Oxygenation of the Newborn: A Molecular Approach. Neonatology, 2012, 101, 315-325.	2.0	70
36	Cell-cycle regulation, intracellular sorting and induced overexpression of the human NTH1 DNA glycosylase involved in removal of formamidopyrimidine residues from DNA. Mutation Research DNA Repair, 2000, 460, 95-104.	3.7	68

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37	Novel Antiviral Activities of Obatoclax, Emetine, Niclosamide, Brequinar, and Homoharringtonine. Viruses, 2019, 11, 964.	3.3	68
38	Ginsenoside Rd promotes glutamate clearance by up-regulating glial glutamate transporter GLT-1 via PI3K/AKT and ERK1/2 pathways. Frontiers in Pharmacology, 2013, 4, 152.	3.5	67
39	Common Nodes of Virus–Host Interaction Revealed Through an Integrated Network Analysis. Frontiers in Immunology, 2019, 10, 2186.	4.8	67
40	Expression patterns of Neil3 during embryonic brain development and neoplasia. BMC Neuroscience, 2009, 10, 45.	1.9	66
41	Neil1 is a genetic modifier of somatic and germline CAG trinucleotide repeat instability in R6/1 mice. Human Molecular Genetics, 2012, 21, 4939-4947.	2.9	66
42	Widespread distribution of DNA glycosylases removing oxidative DNA lesions in human and rodent brains. DNA Repair, 2008, 7, 1578-1588.	2.8	65
43	Human OXR1 maintains mitochondrial DNA integrity and counteracts hydrogen peroxide-induced oxidative stress by regulating antioxidant pathways involving p21. Free Radical Biology and Medicine, 2014, 77, 41-48.	2.9	64
44	Novel activities of safe-in-human broad-spectrum antiviral agents. Antiviral Research, 2018, 154, 174-182.	4.1	64
45	Incision at hypoxanthine residues in DNA by a mammalian homologue of the Escherichia coli antimutator enzyme endonuclease V. Nucleic Acids Research, 2003, 31, 3893-3900.	14.5	58
46	A new protein superfamily includes two novel 3-methyladenine DNA glycosylases fromBacillus cereus, AlkC and AlkD. Molecular Microbiology, 2006, 59, 1602-1609.	2.5	57
47	Limited repair of 8-hydroxy-7,8-dihydroguanine residues in human testicular cells. Nucleic Acids Research, 2003, 31, 1351-1363.	14.5	56
48	The DNA modification N6-methyl-2'-deoxyadenosine (m6dA) drives activity-induced gene expression and is required for fear extinction. Nature Neuroscience, 2019, 22, 534-544.	14.8	51
49	Screening of FDA-Approved Drugs Using a MERS-CoV Clinical Isolate from South Korea Identifies Potential Therapeutic Options for COVID-19. Viruses, 2021, 13, 651.	3.3	50
50	Impaired base excision repair and accumulation of oxidative base lesions in CD4+ T cells of HIV-infected patients. Blood, 2005, 105, 4730-4735.	1.4	49
51	Identification and Tracking of Antiviral Drug Combinations. Viruses, 2020, 12, 1178.	3.3	48
52	Release from quiescence stimulates the expression of human NEIL3 under the control of the Ras dependent ERK–MAP kinase pathway. DNA Repair, 2012, 11, 401-409.	2.8	45
53	Secretoneurin Is a Novel Prognostic Cardiovascular Biomarker Associated With Cardiomyocyte Calcium Handling. Journal of the American College of Cardiology, 2015, 65, 339-351.	2.8	45
54	NEIL3-Dependent Regulation of Cardiac Fibroblast Proliferation Prevents Myocardial Rupture. Cell Reports, 2017, 18, 82-92.	6.4	45

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55	Catalytically impaired hMYH and NEIL1 mutant proteins identified in patients with primary sclerosing cholangitis and cholangiocarcinoma. Carcinogenesis, 2009, 30, 1147-1154.	2.8	43
56	Silver nanoparticles induce premutagenic DNA oxidation that can be prevented by phytochemicals from Gentiana asclepiadea. Mutagenesis, 2012, 27, 759-769.	2.6	43
57	Transcriptome analysis of human OXR1 depleted cells reveals its role in regulating the p53 signaling pathway. Scientific Reports, 2015, 5, 17409.	3.3	43
58	Base excision repair activities required for yeast to attain a full chronological life span. Aging Cell, 2003, 2, 93-104.	6.7	42
59	Increased nuclear DNA damage precedes mitochondrial dysfunction in peripheral blood mononuclear cells from Huntington's disease patients. Scientific Reports, 2018, 8, 9817.	3.3	40
60	A general role of the DNA glycosylase Nth1 in the abasic sites cleavage step of base excision repair in Schizosaccharomyces pombe. Nucleic Acids Research, 2004, 32, 5119-5125.	14.5	39
61	Product inhibition and magnesium modulate the dual reaction mode of hOgg1. DNA Repair, 2005, 4, 381-387.	2.8	39
62	Loss of Neil3, the major DNA glycosylase activity for removal of hydantoins in single stranded DNA, reduces cellular proliferation and sensitizes cells to genotoxic stress. Biochimica Et Biophysica Acta - Molecular Cell Research, 2013, 1833, 1157-1164.	4.1	39
63	Cockayne Syndrome group B protein stimulates NEIL2 DNA glycosylase activity. Mechanisms of Ageing and Development, 2014, 135, 1-14.	4.6	39
64	Synergistic Actions of Ogg1 and Mutyh DNA Glycosylases Modulate Anxiety-like Behavior in Mice. Cell Reports, 2015, 13, 2671-2678.	6.4	39
65	Dissection of the Neonatal Fc Receptor (FcRn)-Albumin Interface Using Mutagenesis and Anti-FcRn Albumin-blocking Antibodies. Journal of Biological Chemistry, 2014, 289, 17228-17239.	3.4	38
66	Biochemical mapping of human NEIL1 DNA glycosylase and AP lyase activities. DNA Repair, 2012, 11, 766-773.	2.8	37
67	No cancer predisposition or increased spontaneous mutation frequencies in NEIL DNA glycosylases-deficient mice. Scientific Reports, 2017, 7, 4384.	3.3	37
68	Lack of the DNA glycosylases MYH and OGG1 in the cancer prone double mutant mouse does not increase mitochondrial DNA mutagenesis. DNA Repair, 2012, 11, 278-285.	2.8	36
69	A new Schizosaccharomyces pombe base excision repair mutant, nth1, reveals overlapping pathways for repair of DNA base damage. Molecular Microbiology, 2003, 48, 465-480.	2.5	34
70	Genetic Interactions of DNA Repair Pathways in the Pathogen Neisseria meningitidis. Journal of Bacteriology, 2007, 189, 5728-5737.	2.2	34
71	Antimicrobial resistance: A challenge awaiting the post-COVID-19 era. International Journal of Infectious Diseases, 2021, 111, 322-325.	3.3	34
72	N6-methyladenosine in RNA of atherosclerotic plaques: An epitranscriptomic signature of human carotid atherosclerosis. Biochemical and Biophysical Research Communications, 2020, 533, 631-637.	2.1	33

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73	Expanding the activity spectrum of antiviral agents. Drug Discovery Today, 2019, 24, 1224-1228.	6.4	31
74	The ada operon of Mycobacterium tuberculosis encodes two DNA methyltransferases for inducible repair of DNA alkylation damage. DNA Repair, 2011, 10, 595-602.	2.8	29
75	AP endonuclease independent repair of abasic sites in Schizosaccharomyces pombe. Nucleic Acids Research, 2012, 40, 2000-2009.	14.5	29
76	Uracil-DNA Glycosylase UNG Promotes Tet-mediated DNA Demethylation. Journal of Biological Chemistry, 2016, 291, 731-738.	3.4	29
77	Susceptibility to infections, without concomitant hyper-IgE, reported in 1976, is caused by hypomorphic mutation in the phosphoglucomutase 3 (PGM3) gene. Clinical Immunology, 2015, 161, 366-372.	3.2	28
78	Glycosylated Chromogranin A in Heart Failure. Circulation: Heart Failure, 2017, 10, .	3.9	28
79	Alkyladenine DNA glycosylase associates with transcription elongation to coordinate DNA repair with gene expression. Nature Communications, 2019, 10, 5460.	12.8	28
80	Structural insight into repair of alkylated DNA by a new superfamily of DNA glycosylases comprising HEAT-like repeats. Nucleic Acids Research, 2007, 35, 2451-2459.	14.5	27
81	Neil3-dependent base excision repair regulates lipid metabolism and prevents atherosclerosis in Apoe-deficient mice. Scientific Reports, 2016, 6, 28337.	3.3	26
82	Monitoring of the spatial and temporal dynamics of BER/SSBR pathway proteins, including MYH, UNG2, MPG, NTH1 and NEIL1-3, during DNA replication. Nucleic Acids Research, 2017, 45, 8291-8301.	14.5	25
83	The Bacillus subtilis Counterpart of the Mammalian 3-Methyladenine DNA Glycosylase Has Hypoxanthine and 1,N6-Ethenoadenine as Preferred Substrates. Journal of Biological Chemistry, 2004, 279, 13601-13606.	3.4	24
84	Neil3 induced neurogenesis protects against prion disease during the clinical phase. Scientific Reports, 2016, 6, 37844.	3.3	24
85	Neuromodulatory Effect of NLRP3 and ASC in Neonatal Hypoxic Ischemic Encephalopathy. Neonatology, 2019, 115, 355-362.	2.0	24
86	Expression and purification of NEIL3, a human DNA glycosylase homolog. Protein Expression and Purification, 2009, 65, 160-164.	1.3	23
87	OXR1A, a Coactivator of PRMT5 Regulating Histone Arginine Methylation. Cell Reports, 2020, 30, 4165-4178.e7.	6.4	23
88	Novel <i>UCHL1</i> mutations reveal new insights into ubiquitin processing. Human Molecular Genetics, 2017, 26, ddw391.	2.9	22
89	Inducible expression of the GLT-1 glutamate transporter in a CHO cell line selected for low endogenous glutamate uptake. FEBS Letters, 1998, 422, 339-342.	2.8	21
90	Mutually exclusive RNA secondary structures regulate translation initiation of DinQ in <i>Escherichia coli</i> . Rna, 2016, 22, 1739-1749.	3.5	21

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91	Synthetic lethality between murine DNA repair factors XLF and DNA-PKcs is rescued by inactivation of Ku70. DNA Repair, 2017, 57, 133-138.	2.8	21
92	Impaired oxidative stress response characterizes HUWE1-promoted X-linked intellectual disability. Scientific Reports, 2017, 7, 15050.	3.3	21
93	The capacity to remove 8-oxoG is enhanced in newborn neural stem/progenitor cells and decreases in juvenile mice and upon cell differentiation. DNA Repair, 2007, 6, 723-732.	2.8	20
94	Genetic variants in the DNA repair gene NEIL3 and the risk of myocardial infarction in a nested case–control study. The HUNT Study. DNA Repair, 2015, 28, 21-27.	2.8	20
95	Peptides containing the PCNA interacting motif APIM bind to the β-clamp and inhibit bacterial growth and mutagenesis. Nucleic Acids Research, 2020, 48, 5540-5554.	14.5	20
96	Synergistic Interferon-Alpha-Based Combinations for Treatment of SARS-CoV-2 and Other Viral Infections. Viruses, 2021, 13, 2489.	3.3	20
97	Absence of NLRP3 Inflammasome in Hematopoietic Cells Reduces Adverse Remodeling After Experimental Myocardial Infarction. JACC Basic To Translational Science, 2020, 5, 1210-1224.	4.1	19
98	Mono- and combinational drug therapies for global viral pandemic preparedness. IScience, 2022, 25, 104112.	4.1	19
99	Fumarylacetoacetate inhibits the initial step of the base excision repair pathway: implication for the pathogenesis of tyrosinemia type I. Journal of Inherited Metabolic Disease, 2013, 36, 773-778.	3.6	18
100	3CAPS – a structural AP–site analogue as a tool to investigate DNA base excision repair. Nucleic Acids Research, 2016, 44, 2187-2198.	14.5	18
101	Integrative whole-genome sequence analysis reveals roles of regulatory mutations in BCL6 and BCL2 in follicular lymphoma. Scientific Reports, 2017, 7, 7040.	3.3	18
102	Comparative analysis of 8-oxoG:C, 8-oxoG:A, A:C and C:C DNA repair in extracts from wild type or 8-oxoG DNA glycosylase deficient mammalian and bacterial cells. DNA Repair, 2003, 2, 707-718.	2.8	17
103	Parp3 promotes astrocytic differentiation through a tight regulation of Nox4-induced ROS and mTorc2 activation. Cell Death and Disease, 2020, 11, 954.	6.3	17
104	The Schizosaccharomyces pombe AlkB homolog Abh1 exhibits AP lyase activity but no demethylase activity. DNA Repair, 2012, 11, 453-462.	2.8	16
105	Genome instability in Maple Syrup Urine Disease correlates with impaired mitochondrial biogenesis. Metabolism: Clinical and Experimental, 2014, 63, 1063-1070.	3.4	16
106	A transgenic minipig model of Huntington's disease shows early signs of behavioral and molecular pathologies. DMM Disease Models and Mechanisms, 2018, 11, .	2.4	15
107	Nafamostat–Interferon-α Combination Suppresses SARS-CoV-2 Infection In Vitro and In Vivo by Cooperatively Targeting Host TMPRSS2. Viruses, 2021, 13, 1768.	3.3	15
108	DrugVirus.info 2.0: an integrative data portal for broad-spectrum antivirals (BSA) and BSA-containing drug combinations (BCCs). Nucleic Acids Research, 2022, 50, W272-W275.	14.5	15

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109	Breaking the speed limit with multimode fast scanning of DNA by Endonuclease V. Nature Communications, 2018, 9, 5381.	12.8	14
110	Generation of a Mouse Model Lacking the Non-Homologous End-Joining Factor Mri/Cyren. Biomolecules, 2019, 9, 798.	4.0	14
111	Antibiotic-induced DNA damage results in a controlled loss of pH homeostasis and genome instability. Scientific Reports, 2020, 10, 19422.	3.3	14
112	The Chromatin Remodeling Factor SMARCB1 Forms a Complex with Human Cytomegalovirus Proteins UL114 and UL44. PLoS ONE, 2012, 7, e34119.	2.5	14
113	Human cytomegalovirus infection modulates DNA base excision repair in fibroblast cells. Virology, 2006, 348, 389-397.	2.4	13
114	The distribution of DNA damage is defined by region-specific susceptibility to DNA damage formation rather than repair differences. DNA Repair, 2014, 18, 44-51.	2.8	13
115	Chemical, Physical and Biological Triggers of Evolutionary Conserved Bcl-xL-Mediated Apoptosis. Cancers, 2020, 12, 1694.	3.7	13
116	ALKBH3 partner ASCC3 mediates P-body formation and selective clearance of MMS-induced 1-methyladenosine and 3-methylcytosine from mRNA. Journal of Translational Medicine, 2021, 19, 287.	4.4	13
117	Up-regulation of myocardial DNA base excision repair activities in experimental heart failure. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2009, 666, 32-38.	1.0	12
118	8-oxoguanine DNA glycosylase (Ogg1) controls hepatic gluconeogenesis. DNA Repair, 2018, 61, 56-62.	2.8	12
119	Base excision repair activities in organotypic hippocampal slice cultures exposed to oxygen and glucose deprivation. DNA Repair, 2008, 7, 869-878.	2.8	11
120	Accelerated clinical course of prion disease in mice compromised in repair of oxidative DNA damage. Free Radical Biology and Medicine, 2014, 68, 1-7.	2.9	11
121	Genetic diversity and drug resistance pattern of Mycobacterium tuberculosis strains isolated from pulmonary tuberculosis patients in the Benishangul Gumuz region and its surroundings, Northwest Ethiopia. PLoS ONE, 2020, 15, e0231320.	2.5	11
122	DNA glycosylase Neil3 regulates vascular smooth muscle cell biology during atherosclerosis development. Atherosclerosis, 2021, 324, 123-132.	0.8	11
123	A Systems Approach to Study Immuno- and Neuro-Modulatory Properties of Antiviral Agents. Viruses, 2018, 10, 423.	3.3	10
124	Diverse functions of DNA glycosylases processing oxidative base lesions in brain. DNA Repair, 2019, 81, 102665.	2.8	10
125	Histone Methylations Define Neural Stem/Progenitor Cell Subtypes in the Mouse Subventricular Zone. Molecular Neurobiology, 2020, 57, 997-1008.	4.0	10
126	PML regulates neuroprotective innate immunity and neuroblast commitment in a hypoxic–ischemic encephalopathy model. Cell Death and Disease, 2016, 7, e2320-e2320.	6.3	9

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127	Crystal structure and MD simulation of mouse EndoV reveal wedge motif plasticity in this inosine-specific endonuclease. Scientific Reports, 2016, 6, 24979.	3.3	9
128	Metabolism and DNA repair shape a specific modification pattern in mitochondrial DNA. Mitochondrion, 2018, 40, 16-28.	3.4	9
129	Deletion of Endonuclease V suppresses chemically induced hepatocellular carcinoma. Nucleic Acids Research, 2020, 48, 4463-4479.	14.5	9
130	Structural basis for substrate and product recognition in human phosphoglucomutase-1 (PGM1) isoform 2, a member of the α-d-phosphohexomutase superfamily. Scientific Reports, 2020, 10, 5656.	3.3	9
131	Sculpting of DNA at Abasic Sites by DNA Glycosylase Homolog Mag2. Structure, 2013, 21, 154-166.	3.3	8
132	A new family of proteins related to the HEAT-like repeat DNA glycosylases with affinity for branched DNA structures. Journal of Structural Biology, 2013, 183, 66-75.	2.8	8
133	Structural basis for incision at deaminated adenines in DNA and RNA by endonuclease V. Progress in Biophysics and Molecular Biology, 2015, 117, 134-142.	2.9	8
134	Regulation of Human Endonuclease V Activity and Relocalization to Cytoplasmic Stress Granules. Journal of Biological Chemistry, 2016, 291, 21786-21801.	3.4	8
135	Excision of the doubly methylated base <i>N</i> ⁴ ,5-dimethylcytosine from DNA by <i>Escherichia coli</i> Nei and Fpg proteins. Philosophical Transactions of the Royal Society B: Biological Sciences, 2018, 373, 20170337.	4.0	8
136	Synthesis and antimicrobial activities of chitosan/polypropylene carbonate-based nanoparticles. RSC Advances, 2021, 11, 10121-10129.	3.6	8
137	NEIL1 and NEIL2 DNA glycosylases modulate anxiety and learning in a cooperative manner in mice. Communications Biology, 2021, 4, 1354.	4.4	8
138	Reduced expression of DNA glycosylases in post-hypoxic newborn pigs undergoing therapeutic hypothermia. Brain Research, 2010, 1363, 198-205.	2.2	7
139	Insight into ALKBH8-related intellectual developmental disability based on the first pathogenic missense variant. Human Genetics, 2022, 141, 209-215.	3.8	7
140	Production, isolation and purification of bacteriocins expressed by two strains of <i>Neisseria meningitidis</i> . Apmis, 1998, 106, 1181-1187.	2.0	6
141	Modulation of DNA glycosylase activities in mesenchymal stem cells. Experimental Cell Research, 2009, 315, 2558-2567.	2.6	6
142	DNA repair enzyme NEIL3 enables a stable neural representation of space by shaping transcription in hippocampal neurons. IScience, 2021, 24, 103470.	4.1	6
143	Human NEIL1 localizes with the centrosomes and condensed chromosomes during mitosis. DNA Repair, 2007, 6, 1425-1433.	2.8	5
144	Addressing RNA Integrity to Determine the Impact of Mitochondrial DNA Mutations on Brain Mitochondrial Function with Age. PLoS ONE, 2014, 9, e96940.	2.5	5

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145	Increased p53 signaling impairs neural differentiation in HUWE1-promoted intellectual disabilities. Cell Reports Medicine, 2021, 2, 100240.	6.5	5
146	Impact of Oxidative DNA Damage and the Role of DNA Glycosylases in Neurological Dysfunction. International Journal of Molecular Sciences, 2021, 22, 12924.	4.1	5
147	Neonatal Ogg1/Mutyh knockout mice have altered inflammatory gene response compared to wildtype mice in the brain and lung after hypoxia-reoxygenation. Journal of Perinatal Medicine, 2018, 47, 114-124.	1.4	4
148	Additive manufacturing of laminar flow cells for single-molecule experiments. Scientific Reports, 2019, 9, 16784.	3.3	4
149	HMST-Seq-Analyzer: A new python tool for differential methylation and hydroxymethylation analysis in various DNA methylation sequencing data. Computational and Structural Biotechnology Journal, 2020, 18, 2877-2889.	4.1	4
150	DNA glycosylase Neil2 contributes to genomic responses in the spleen during clinical prion disease. Free Radical Biology and Medicine, 2020, 152, 348-354.	2.9	4
151	NEIL3-deficiency increases gut permeability and contributes to a pro-atherogenic metabolic phenotype. Scientific Reports, 2021, 11, 19749.	3.3	4
152	Development of DinQ from Escherichia coli as an anti-cell-envelope antibiotic. International Journal of Antimicrobial Agents, 2015, 45, 196-197.	2.5	3
153	Enhanced base excision repair capacity in carotid atherosclerosis may protect nuclear DNA but not mitochondrial DNA. Free Radical Biology and Medicine, 2016, 97, 386-397.	2.9	3
154	Complex alternative splicing of human Endonuclease V mRNA, but evidence for only a single protein isoform. PLoS ONE, 2019, 14, e0225081.	2.5	3
155	Active Components of Commonly Prescribed Medicines Affect Influenza A Virus–Host Cell Interaction: A Pilot Study. Viruses, 2021, 13, 1537.	3.3	3
156	Efficient and Reliable Production of Vectors for the Study of the Repair, Mutagenesis, and Phenotypic Consequences of Defined DNA Damage Lesions in Mammalian Cells. PLoS ONE, 2016, 11, e0158581.	2.5	2
157	NEIL3-deficient bone marrow displays decreased hematopoietic capacity and reduced telomere length. Biochemistry and Biophysics Reports, 2022, 29, 101211.	1.3	2
158	IGAP-integrative genome analysis pipeline reveals new gene regulatory model associated with nonspecific TF-DNA binding affinity. Computational and Structural Biotechnology Journal, 2020, 18, 1270-1286.	4.1	1
159	Loss of Mediator complex subunit 13 (MED13) promotes resistance to alkylation through cyclin D1 upregulation. Nucleic Acids Research, 2021, 49, 1470-1484.	14.5	1
160	abc4pwm: affinity based clustering for position weight matrices in applications of DNA sequence analysis. BMC Bioinformatics, 2022, 23, 83.	2.6	1
161	Epitranscriptome in Ischemic Cardiovascular Disease: Potential Target for Therapies. Stroke, 2022, 53, 2114-2122.	2.0	1
162	PML Regulates the Epidermal Differentiation Complex and Skin Morphogenesis during Mouse Embryogenesis. Genes, 2020, 11, 1130.	2.4	0

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163	Non-flipping DNA glycosylase AlkD scans DNA without formation of a stable interrogation complex. Communications Biology, 2021, 4, 876.	4.4	0
164	Cause and consequences of genome instability in Huntington's Disease. FASEB Journal, 2010, 24, 411.3.	0.5	0
165	A chemicalâ€genetic screen to unravel the genetic network of CDC28/CDK1 links ubiquitin and Rad6–Bre1 to cell cycle progression. FASEB Journal, 2012, 26, 590.1.	0.5	0
166	An in vitro Study of Protein Adsorption to Biocompatible Coatings. Studies in Health Technology and Informatics, 2015, 211, 166-71.	0.3	0
167	Title is missing!. , 2020, 15, e0231320.		0
168	Title is missing!. , 2020, 15, e0231320.		0
169	Title is missing!. , 2020, 15, e0231320.		0
170	Title is missing!. , 2020, 15, e0231320.		0
171	Title is missing!. , 2020, 15, e0231320.		0
172	Title is missing!. , 2020, 15, e0231320.		0
173	Title is missing!. , 2020, 15, e0231320.		0