## Yusaku Nakabeppu

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5620826/publications.pdf

Version: 2024-02-01

276 papers 17,963 citations

67 h-index 121 g-index

297 all docs

297 docs citations

times ranked

297

14735 citing authors

#	Article	IF	CITATIONS
1	DNA binding activities of three murine Jun proteins: Stimulation by Fos. Cell, 1988, 55, 907-915.	28.9	841
2	Regulation and Expression of the Adaptive Response to Alkylating Agents. Annual Review of Biochemistry, 1988, 57, 133-157.	11.1	666
3	Differentiation-induced gene expression in 3T3-L1 preadipocytes: CCAAT/enhancer binding protein interacts with and activates the promoters of two adipocyte-specific genes Genes and Development, 1989, 3, 1323-1335.	5.9	561
4	Induction of a long-lasting AP-1 complex composed of altered Fos-like proteins in brain by chronic cocaine and other chronic treatments. Neuron, 1994, 13, 1235-1244.	8.1	535
5	Galectin-1 is essential in tumor angiogenesis and is a target for antiangiogenesis therapy. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 15975-15980.	7.1	424
6	A naturally occurring truncated form of FosB that inhibits Fos/Jun transcriptional activity. Cell, 1991, 64, 751-759.	28.9	415
7	Expression and Differential Intracellular Localization of Two Major Forms of Human 8-Oxoguanine DNA Glycosylase Encoded by Alternatively Spliced OGG1 mRNAs. Molecular Biology of the Cell, 1999, 10, 1637-1652.	2.1	365
8	Altered Expression of Diabetes-Related Genes in Alzheimer's Disease Brains: The Hisayama Study. Cerebral Cortex, 2014, 24, 2476-2488.	2.9	294
9	Chronic Fos-Related Antigens: Stable Variants of ΔFosB Induced in Brain by Chronic Treatments. Journal of Neuroscience, 1997, 17, 4933-4941.	3.6	293
10	Identification of human MutY homolog (hMYH) as a repair enzyme for 2-hydroxyadenine in DNA and detection of multiple forms of hMYH located in nuclei and mitochondria. Nucleic Acids Research, 2000, 28, 1355-1364.	14.5	282
11	Spontaneous tumorigenesis in mice defective in the MTH1 gene encoding 8-oxo-dGTPase. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 11456-11461.	7.1	276
12	JSAP1, a Novel Jun N-Terminal Protein Kinase (JNK)-Binding Protein That Functions as a Scaffold Factor in the JNK Signaling Pathway. Molecular and Cellular Biology, 1999, 19, 7539-7548.	2.3	270
13	Constitutive expression of zif268 in neocortex is regulated by synaptic activity Proceedings of the National Academy of Sciences of the United States of America, 1991, 88, 5106-5110.	7.1	263
14	Oxidative damage in nucleic acids and Parkinson's disease. Journal of Neuroscience Research, 2007, 85, 919-934.	2.9	254
15	Mutagenesis and carcinogenesis caused by the oxidation of nucleic acids. Biological Chemistry, 2006, 387, 373-9.	2.5	212
16	The Oxidized Forms of dATP Are Substrates for the Human MutT Homologue, the hMTH1 Protein. Journal of Biological Chemistry, 1999, 274, 18201-18205.	3.4	204
17	Two distinct pathways of cell death triggered by oxidative damage to nuclear and mitochondrial DNAs. EMBO Journal, 2008, 27, 421-432.	7.8	194
18	Ogg1 knockout-associated lung tumorigenesis and its suppression by Mth1 gene disruption. Cancer Research, 2003, 63, 902-5.	0.9	185

#	Article	IF	CITATIONS
19	DNA Polymerases as Potential Therapeutic Targets for Cancers Deficient in the DNA Mismatch Repair Proteins MSH2 or MLH1. Cancer Cell, 2010, 17, 235-248.	16.8	181
20	Chronic Alterations in Dopaminergic Neurotransmission Produce a Persistent Elevation of ΔFosB-like Protein(s) in both the Rodent and Primate Striatum. European Journal of Neuroscience, 1996, 8, 365-381.	2.6	178
21	Intracellular Localization of 8-Oxo-dGTPase in Human Cells, with Special Reference to the Role of the Enzyme in Mitochondria. Journal of Biological Chemistry, 1995, 270, 14659-14665.	3.4	172
22	Hydrogen in Drinking Water Reduces Dopaminergic Neuronal Loss in the 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine Mouse Model of Parkinson's Disease. PLoS ONE, 2009, 4, e7247.	2.5	170
23	Silencing effect of CpG island hypermethylation and histone modifications on O6-methylguanine-DNA methyltransferase (MGMT) gene expression in human cancer. Oncogene, 2003, 22, 8835-8844.	5.9	164
24	APE1- and APE2-dependent DNA breaks in immunoglobulin class switch recombination. Journal of Experimental Medicine, 2007, 204, 3017-3026.	8.5	156
25	Increased 8-oxo-dGTPase in the mitochondria of substantia nigral neurons in Parkinson's disease. Annals of Neurology, 1999, 46, 920-924.	5.3	155
26	Cellular Levels of 8-Oxoguanine in either DNA or the Nucleotide Pool Play Pivotal Roles in Carcinogenesis and Survival of Cancer Cells. International Journal of Molecular Sciences, 2014, 15, 12543-12557.	4.1	152
27	Genomic Structure and Chromosome Location of the Human mutT Homologue Gene MTH1 Encoding 8-Oxo-dGTPase for Prevention of A:T to C:G Transversion. Genomics, 1994, 24, 485-490.	2.9	149
28	Three-Dimensional Structure of a DNA Repair Enzyme, 3-Methyladenine DNA Glycosylase II, from Escherichia coli. Cell, 1996, 86, 311-319.	28.9	147
29	Human APE2 protein is mostly localized in the nuclei and to some extent in the mitochondria, while nuclear APE2 is partly associated with proliferating cell nuclear antigen. Nucleic Acids Research, 2001, 29, 2349-2360.	14.5	145
30	XRCC1 interactions with multiple DNA glycosylases: A model for its recruitment to base excision repair. DNA Repair, 2005, 4, 826-835.	2.8	145
31	8-oxoguanine causes spontaneous de novo germline mutations in mice. Scientific Reports, 2014, 4, 4689.	3.3	140
32	A single-point mutation in HCF causes temperature-sensitive cell-cycle arrest and disrupts VP16 function Genes and Development, 1997, 11, 726-737.	5.9	139
33	Comparative profiling of cortical gene expression in Alzheimer's disease patients and mouse models demonstrates a link between amyloidosis and neuroinflammation. Scientific Reports, 2017, 7, 17762.	3.3	138
34	Regulation of intracellular localization of human MTH1, OGG1, and MYH proteins for repair of oxidative DNA damage. Progress in Molecular Biology and Translational Science, 2001, 68, 75-94.	1.9	137
35	Metabolic Fate of Oxidized Guanine Ribonucleotides in Mammalian Cellsâ€. Biochemistry, 1999, 38, 3610-3614.	2.5	132
36	The Oxidized Deoxynucleoside Triphosphate Pool Is a Significant Contributor to Genetic Instability in Mismatch Repair-Deficient Cells. Molecular and Cellular Biology, 2004, 24, 465-474.	2.3	126

3

#	Article	IF	CITATIONS
37	MUTYH-Null Mice Are Susceptible to Spontaneous and Oxidative Stress–Induced Intestinal Tumorigenesis. Cancer Research, 2007, 67, 6599-6604.	0.9	125
38	Physical association of pyrimidine dimer DNA glycosylase and apurinic/apyrimidinic DNA endonuclease essential for repair of ultraviolet-damaged DNA Proceedings of the National Academy of Sciences of the United States of America, 1981, 78, 2742-2746.	7.1	123
39	Expression of 8-oxoguanine DNA glycosylase is reduced and associated with neurofibrillary tangles in Alzheimer's disease brain. Acta Neuropathologica, 2002, 103, 20-25.	7.7	122
40	Expression of 8-oxoguanine DNA glycosylase (OGG1) in Parkinson?s disease and related neurodegenerative disorders. Acta Neuropathologica, 2005, 109, 256-262.	7.7	122
41	8-Oxoguanine Formation Induced by Chronic UVB Exposure Makes Ogg1 Knockout Mice Susceptible to Skin Carcinogenesis. Cancer Research, 2005, 65, 6006-6010.	0.9	121
42	An Oxidized Purine Nucleoside Triphosphatase, MTH1, Suppresses Cell Death Caused by Oxidative Stress. Journal of Biological Chemistry, 2003, 278, 37965-37973.	3.4	120
43	Human MTH1 protein hydrolyzes the oxidized ribonucleotide, 2-hydroxy-ATP. Nucleic Acids Research, 2001, 29, 449-454.	14.5	118
44	The dopamine D1 receptor is a critical mediator for cocaineâ€induced gene expression. Journal of Neurochemistry, 2002, 82, 1453-1464.	3.9	113
45	Molecular genetics and structural biology of human MutT homolog, MTH1. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2001, 477, 59-70.	1.0	112
46	8-Oxoguanine causes neurodegeneration during MUTYH-mediated DNA base excision repair. Journal of Clinical Investigation, 2012, 122, 4344-4361.	8.2	110
47	8-Oxoguanine DNA Glycosylase (OGG1) Deficiency Increases Susceptibility to Obesity and Metabolic Dysfunction. PLoS ONE, 2012, 7, e51697.	2.5	108
48	Regulatory mechanisms for induction of synthesis of repair enzymes in response to alkylating agents: ada protein acts as a transcriptional regulator Proceedings of the National Academy of Sciences of the United States of America, 1986, 83, 6297-6301.	7.1	105
49	Molecular pathophysiology of impaired glucose metabolism, mitochondrial dysfunction, and oxidative DNA damage in Alzheimer's disease brain. Mechanisms of Ageing and Development, 2017, 161, 95-104.	4.6	105
50	Stress response gene ATF3 is a target of c-myc in serum-induced cell proliferation. EMBO Journal, 2005, 24, 2590-2601.	7.8	99
51	A genome-wide distribution of 8-oxoguanine correlates with the preferred regions for recombination and single nucleotide polymorphism in the human genome. Genome Research, 2006, 16, 567-575.	5.5	98
52	Differential regulation of fos family genes in the ventrolateral and dorsomedial subdivisions of the rat suprachiasmatic nucleus. Neuroscience, 2000, 98, 535-547.	2.3	93
53	Impairment of mitochondrial DNA repair enzymes against accumulation of 8-oxo-guanine in the spinal motor neurons of amyotrophic lateral sclerosis. Acta Neuropathologica, 2002, 103, 408-414.	7.7	92
54	ReviewThe Defense Mechanisms in Mammalian Cells against Oxidative Damage in Nucleic Acids and their Involvement in the Suppression of Mutagenesis and Cell Death. Free Radical Research, 2004, 38, 423-429.	3.3	92

#	Article	IF	Citations
55	Accumulation of 8-oxoguanine in the cellular DNA and the alteration of the OGG1 expression during ischemia-reperfusion injury in the rat kidney. DNA Repair, 2003, 2, 211-229.	2.8	89
56	Significance of error-avoiding mechanisms for oxidative DNA damage in carcinogenesis. Cancer Science, 2007, 98, 465-470.	3.9	89
57	Synaptic Regulation of Immediate Early Gene Expression in Primary Cultures of Cortical Neurons. Journal of Neurochemistry, 1991, 57, 1862-1872.	3.9	85
58	Suberoylanilide hydroxamic acid (SAHA) induces apoptosis or autophagy-associated cell death in chondrosarcoma cell lines. Anticancer Research, 2008, 28, 1585-91.	1.1	85
59	Replication-Associated Repair of Adenine:8-Oxoguanine Mispairs by MYH. Current Biology, 2002, 12, 335-339.	3.9	81
60	Mutator Phenotype of MUTYH-null Mouse Embryonic Stem Cells. Journal of Biological Chemistry, 2003, 278, 38121-38124.	3.4	80
61	Selective increase in cellular $\hat{A}^2$ 42 is related to apoptosis but not necrosis. NeuroReport, 2000, 11, 167-171.	1.2	78
62	MTH1, an oxidized purine nucleoside triphosphatase, protects the dopamine neurons from oxidative damage in nucleic acids caused by 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine. Cell Death and Differentiation, 2006, 13, 551-563.	11.2	76
63	A Molecular Basis for the Selective Recognition of 2-Hydroxy-dATP and 8-Oxo-dGTP by Human MTH1. Journal of Biological Chemistry, 2002, 277, 8579-8587.	3.4	<b>7</b> 5
64	MTH1, an oxidized purine nucleoside triphosphatase, prevents the cytotoxicity and neurotoxicity of oxidized purine nucleotides. DNA Repair, 2006, 5, 761-772.	2.8	75
65	RNA polymerase II bypasses 8-oxoguanine in the presence of transcription elongation factor TFIIS. DNA Repair, 2007, 6, 841-851.	2.8	<b>7</b> 5
66	Molecular cloning of AtMMH, an Arabidopsis thaliana ortholog of the Escherichia coli mutM gene, and analysis of functional domains of its product. Molecular Genetics and Genomics, 1998, 259, 577-590.	2.4	71
67	Induction of c-Fos-like and FosB-like immunoreactivity reveals forebrain neuronal populations involved differentially in pup-mediated maternal behavior in juvenile and adult rats., 2000, 416, 45-78.		69
68	Biological Significance of the Defense Mechanisms against Oxidative Damage in Nucleic Acids Caused by Reactive Oxygen Species: From Mitochondria to Nuclei. Annals of the New York Academy of Sciences, 2004, 1011, 101-111.	3.8	69
69	DNA glycosylase encoded by <i>MUTYH</i> functions as a molecular switch for programmed cell death under oxidative stress to suppress tumorigenesis. Cancer Science, 2011, 102, 677-682.	3.9	68
70	8-Oxoguanine accumulation in mitochondrial DNA causes mitochondrial dysfunction and impairs neuritogenesis in cultured adult mouse cortical neurons under oxidative conditions. Scientific Reports, 2016, 6, 22086.	3.3	66
71	Synaptic Regulation of Immediate-Early Genes in Brain. Cold Spring Harbor Symposia on Quantitative Biology, 1990, 55, 213-223.	1.1	63
72	Functional sites of the Ada regulatory protein of Escherichia coli. Journal of Molecular Biology, 1988, 201, 261-271.	4.2	62

#	Article	IF	Citations
73	Narrow-Band UVB Induces More Carcinogenic Skin Tumors than Broad-Band UVB through the Formation of Cyclobutane Pyrimidine Dimer. Journal of Investigative Dermatology, 2007, 127, 2865-2871.	0.7	62
74	ITPase-deficient mice show growth retardation and die before weaning. Cell Death and Differentiation, 2009, 16, 1315-1322.	11.2	62
75	Hematopoietic Tissue–Specific Expression of Mouse Neil3 for Endonuclease VIII–Like Protein. Journal of Biochemistry, 2005, 138, 763-772.	1.7	60
76	Novel role of neuronal Ca2+ sensor-1 as a survival factor up-regulated in injured neurons. Journal of Cell Biology, 2006, 172, 1081-1091.	5.2	59
77	Oxidation of mitochondrial deoxynucleotide pools by exposure to sodium nitroprusside induces cell death. DNA Repair, 2008, 7, 418-430.	2.8	58
78	Programmed cell death triggered by nucleotide pool damage and its prevention by MutT homolog-1 (MTH1) with oxidized purine nucleoside triphosphatase. Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2010, 703, 51-58.	1.7	58
79	Oral â€~hydrogen water' induces neuroprotective ghrelin secretion in mice. Scientific Reports, 2013, 3, 3273.	3.3	58
80	MTH1 as a nucleotide pool sanitizing enzyme: Friend or foe?. Free Radical Biology and Medicine, 2017, 107, 151-158.	2.9	58
81	Adaptive response: induced synthesis of DNA repair enzymes by alkylating agents. Trends in Genetics, 1987, 3, 51-54.	6.7	57
82	Regulation of Expression of the Human MTH1 Gene Encoding 8-Oxo-dGTPase. Journal of Biological Chemistry, 1997, 272, 17843-17850.	3.4	57
83	MTH1, an Oxidized Purine Nucleoside Triphosphatase, Suppresses the Accumulation of Oxidative Damage of Nucleic Acids in the Hippocampal Microglia during Kainate-Induced Excitotoxicity. Journal of Neuroscience, 2006, 26, 1688-1698.	3.6	57
84	Genomic Response to Growth Factors. Cold Spring Harbor Symposia on Quantitative Biology, 1988, 53, 893-900.	1.1	57
85	CpG methylation of MGMT and hMLH1 promoter in hepatocellular carcinoma associated with hepatitis viral infection. British Journal of Cancer, 2003, 88, 521-529.	6.4	56
86	Human mitochondrial transcriptional factor A breaks the mitochondria-mediated vicious cycle in Alzheimer's disease. Scientific Reports, 2016, 6, 37889.	3.3	56
87	Deficiency of base excision repair enzyme NEIL3 drives increased predisposition to autoimmunity. Journal of Clinical Investigation, 2016, 126, 4219-4236.	8.2	56
88	Accumulation of 8-oxo-2′-deoxyguanosine and increased expression of hMTH1 protein in brain tumors. Neuro-Oncology, 2001, 3, 73-81.	1.2	55
89	NUDT16 and ITPA play a dual protective role in maintaining chromosome stability and cell growth by eliminating dIDP/IDP and dITP/ITP from nucleotide pools in mammals. Nucleic Acids Research, 2010, 38, 2891-2903.	14.5	55
90	FER-1, an Enhancer of the Ferritin H Gene and a Target of E1A-Mediated Transcriptional Repression. Molecular and Cellular Biology, 1995, 15, 5152-5164.	2.3	54

#	Article	IF	CITATIONS
91	Quantitative Analysis of Oxidized Guanine, 8-Oxoguanine, in Mitochondrial DNA by Immunofluorescence Method. Methods in Molecular Biology, 2009, 554, 199-212.	0.9	54
92	A Role for Oxidized DNA Precursors in Huntington's Disease–Like Striatal Neurodegeneration. PLoS Genetics, 2008, 4, e1000266.	3.5	53
93	DNA-repair methyltransferase as a molecular device for preventing mutation and cancer. Journal of Cancer Research and Clinical Oncology, 1996, 122, 199-206.	2.5	52
94	Deficient expression of O 6-Methylguanine-DNA methyltransferase combined with mismatch-repair proteins hMLH1 and hMSH2 is related to poor prognosis in human biliary tract carcinoma. Annals of Surgical Oncology, 2002, 9, 371-379.	1.5	52
95	<i>Fosb</i> gene products contribute to excitotoxic microglial activation by regulating the expression of complement C5a receptors in microglia. Glia, 2014, 62, 1284-1298.	4.9	52
96	Differential expression of APE1 and APE2 in germinal centers promotes error-prone repair and A:T mutations during somatic hypermutation. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 9217-9222.	7.1	52
97	Celecoxib and 2,5â€dimethylcelecoxib inhibit intestinal cancer growth by suppressing the Wnt∫2â€catenin signaling pathway. Cancer Science, 2017, 108, 108-115.	3.9	52
98	Characterization of human polymorphic DNA repair methyltransferase. Pharmacogenetics and Genomics, 2000, 10, 59-66.	5.7	51
99	Up-regulation of hMUTYH, a DNA repair enzyme, in the mitochondria of substantia nigra in Parkinson's disease. Acta Neuropathologica, 2006, 112, 139-145.	7.7	51
100	Inhibitory Effects of Dietary Spirulina platensis on UVB-Induced Skin Inflammatory Responses and Carcinogenesis. Journal of Investigative Dermatology, 2014, 134, 2610-2619.	0.7	51
101	Futile short-patch DNA base excision repair of adenine:8-oxoguanine mispair. Nucleic Acids Research, 2004, 32, 5928-5934.	14.5	50
102	Targeted disruption of one allele of the Y-box binding protein-1 (YB-1) gene in mouse embryonic stem cells and increased sensitivity to cisplatin and mitomycin C. Cancer Science, 2004, 95, 348-353.	3.9	50
103	Expression of hMTH1 in the hippocampi of control and Alzheimer's disease. NeuroReport, 2001, 12, 2895-2899.	1.2	49
104	Structure of Human MTH1, a Nudix Family Hydrolase That Selectively Degrades Oxidized Purine Nucleoside Triphosphates. Journal of Biological Chemistry, 2004, 279, 33806-33815.	3.4	49
105	Structure and function ofdnaQ andmutD mutators of Escherichia coli. Molecular Genetics and Genomics, 1986, 205, 9-13.	2.4	48
106	Transfer of the E. coli O6-methyltransferase gene into repair-deficient human cells and restoration of cellular resistance to N-methyl-N′-nitro-N-nitrosoguanidine. Mutation Research - DNA Repair Reports, 1986, 166, 135-141.	1.8	47
107	Contrasting Effects of Chronic Clozapine, SeroquelTM(ICI 204,636) and Haloperidol Administration on ΔFosB-like Immunoreactivity in the Rodent Forebrain. European Journal of Neuroscience, 1996, 8, 927-936.	2.6	47
108	Differential Expression of Cytokeratin after Orthotopic Implantation of Newly Established Human Tongue Cancer Cell Lines of Defined Metastatic Ability. American Journal of Pathology, 2000, 156, 1317-1326.	3.8	47

#	Article	IF	Citations
109	Multi-forms of human MTH1 polypeptides produced by alternative translation initiation and single nucleotide polymorphism. Nucleic Acids Research, 1999, 27, 4335-4343.	14.5	45
110	Growth retardation and dyslymphopoiesis accompanied by $G2/M$ arrest in APEX2-null mice. Blood, 2004, 104, 4097-4103.	1.4	45
111	Contrasting Genome-Wide Distribution of 8-Hydroxyguanine and Acrolein-Modified Adenine during Oxidative Stress-Induced Renal Carcinogenesis. American Journal of Pathology, 2006, 169, 1328-1342.	3.8	45
112	Genomic and functional analyses of <i>MUTYH</i> in Japanese patients with adenomatous polyposis. Clinical Genetics, 2008, 73, 545-553.	2.0	45
113	Biochemical and physicochemical characterization of normal and variant forms of human MTH1 protein with antimutagenic activity. Mutation Research DNA Repair, 1997, 384, 181-194.	3.7	44
114	Expression and Prognostic Significance of O 6-Methylguanine-DNA Methyltransferase in Hepatocellular, Gastric, and Breast Cancers. Annals of Surgical Oncology, 2001, 8, 807-816.	1.5	44
115	Galectin-1 promotes basal and kainate-induced proliferation of neural progenitors in the dentate gyrus of adult mouse hippocampus. Cell Death and Differentiation, 2009, 16, 417-427.	11.2	43
116	ITPA protein, an enzyme that eliminates deaminated purine nucleoside triphosphates in cells. Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2010, 703, 43-50.	1.7	43
117	NUDT16 is a (deoxy)inosine diphosphatase, and its deficiency induces accumulation of single-strand breaks in nuclear DNA and growth arrest. Nucleic Acids Research, 2010, 38, 4834-4843.	14.5	42
118	Expression of the ada gene of Escherichia coli in response to alkylating agents. Journal of Molecular Biology, 1988, 202, 483-494.	4.2	41
119	Therapeutic Approach to Neurodegenerative Diseases by Medical Gases: Focusing on Redox Signaling and Related Antioxidant Enzymes. Oxidative Medicine and Cellular Longevity, 2012, 2012, 1-9.	4.0	41
120	MUTYH, an adenine DNA glycosylase, mediates p53 tumor suppression via PARP-dependent cell death. Oncogenesis, 2014, 3, e121-e121.	4.9	41
121	Sensitivity of Escherichia coli (MutT) and human (MTH1) 8-oxo-dGTPases to in vitro inhibition by the carcinogenic metals, nickel(II), copper(II), cobalt(II) and cadmium(II). Carcinogenesis, 1997, 18, 1785-1791.	2.8	40
122	A defect in a single allele of the Mlh1 gene causes dissociation of the killing and tumorigenic actions of an alkylating carcinogen in methyltransferase-deficient mice. Carcinogenesis, 2000, 21, 301-305.	2.8	40
123	fosB-Null Mice Display Impaired Adult Hippocampal Neurogenesis and Spontaneous Epilepsy with Depressive Behavior. Neuropsychopharmacology, 2013, 38, 895-906.	5.4	40
124	Cloning and sequence of the gene encoding the major structural component of mannose-resistant fimbriae of Serratia marcescens. Journal of Bacteriology, 1988, 170, 3567-3574.	2.2	39
125	Synergistic Actions of Ogg1 and Mutyh DNA Glycosylases Modulate Anxiety-like Behavior in Mice. Cell Reports, 2015, 13, 2671-2678.	6.4	39
126	Functional Significance of the Conserved Residues for the 23-Residue Module among MTH1 and MutT Family Proteins. Journal of Biological Chemistry, 1999, 274, 38251-38259.	3.4	38

#	Article	IF	CITATIONS
127	Increased Expression of Versican in the Inflammatory Response to UVB- and Reactive Oxygen Species-Induced Skin Tumorigenesis. American Journal of Pathology, 2011, 179, 3056-3065.	3.8	38
128	A functional analysis of the DNA glycosylase activity of mouse MUTYH protein excising 2-hydroxyadenine opposite guanine in DNA. Nucleic Acids Research, 2005, 33, 672-682.	14.5	37
129	Apex2 is required for efficient somatic hypermutation but not for class switch recombination of immunoglobulin genes. International Immunology, 2009, 21, 947-955.	4.0	37
130	Adenine DNA glycosylase activity of 14 Human MutY homolog (MUTYH) variant proteins found in patients with colorectal polyposis and cancer. Human Mutation, 2010, 31, E1861-E1874.	2.5	37
131	GDNF promotes neurite outgrowth and upregulates galectin-1 through the RET/PI3K signaling in cultured adult rat dorsal root ganglion neurons. Neurochemistry International, 2013, 62, 330-339.	3.8	37
132	Identification and characterization of two forms of mouse MUTYH proteins encoded by alternatively spliced transcripts. Nucleic Acids Research, 2004, 32, 477-487.	14.5	36
133	Galectin- $\hat{1}^2$ , a natural monomeric form of galectin-1 lacking its six amino-terminal residues promotes axonal regeneration but not cell death. Cell Death and Differentiation, 2004, 11, 1076-1083.	11.2	36
134	FosB Is Essential for the Enhancement of Stress Tolerance and Antagonizes Locomotor Sensitization by î"FosB. Biological Psychiatry, 2011, 70, 487-495.	1.3	36
135	Deficient Expression of O6-Methylguanine-DNA Methyltransferase Combined With Mismatch-Repair Proteins hMLH1 and hMSH2 Is Related to Poor Prognosis in Human Biliary Tract Carcinoma. Annals of Surgical Oncology, 2002, 9, 371-379.	1.5	36
136	MutT Homolog-1 Attenuates Oxidative DNA Damage and Delays Photoreceptor Cell Death in Inherited Retinal Degeneration. American Journal of Pathology, 2012, 181, 1378-1386.	3.8	35
137	Polymorphism in the human O6-methylguanine-DNA methyltransferase gene detected by PCR-SSCP analysis. Pharmacogenetics and Genomics, 1996, 6, 361-363.	5.7	33
138	MUTYH prevents OGG1 or APEX1 from inappropriately processing its substrate or reaction product with its C-terminal domain. Nucleic Acids Research, 2004, 32, 3198-3211.	14.5	33
139	Chronic atypical antipsychotics, but not haloperidol, increase neurogenesis in the hippocampus of adult mouse. Brain Research, 2017, 1676, 77-82.	2.2	33
140	Accumulation of 8-oxo-2'-deoxyguanosine and increased expression of hMTH1 protein in brain tumors. Neuro-Oncology, 2001, 3, 73-81.	1.2	33
141	The conjunctive stimuli that cause long-term desensitization also predominantly induce c-Fos and Jun-B in cerebellar Purkinje cells. NeuroReport, 1993, 4, 1275-1278.	1.2	32
142	A novel Nudix hydrolase for oxidized purine nucleoside triphosphates encoded by ORFYLR151c (PCD1) Tj ETQqC	0 0 ggBT	/Overlock 10 1
143	Mice Heterozygous for the Xanthine Oxidoreductase Gene Facilitate Lipid Accumulation in Adipocytes. Arteriosclerosis, Thrombosis, and Vascular Biology, 2014, 34, 44-51.	2.4	32
144	Ability of various alkylating agents to induce adaptive and SOS responses: A study with lacZ fusion. Mutation Research - DNA Repair Reports, 1985, 146, 149-154.	1.8	31

#	Article	IF	CITATIONS
145	Isoforms of JSAP1 scaffold protein generated through alternative splicing. Gene, 2000, 255, 229-234.	2.2	31
146	Selective induction of î"FosB in the brain after transient forebrain ischemia accompanied by an increased expression of galectin-1, and the implication of î"FosB and galectin-1 in neuroprotection and neurogenesis. Cell Death and Differentiation, 2005, 12, 1078-1096.	11.2	31
147	Regulation of expression of the cloned ada gene in Escherichia coli. Mutation Research - DNA Repair Reports, 1985, 146, 155-167.	1.8	30
148	Contribution of Estrogen Receptor $\hat{l}_{\pm}$ to Oncogenic K-Ras-mediated NIH3T3 Cell Transformation and Its Implication for Escape from Senescence by Modulating the p53 Pathway. Journal of Biological Chemistry, 2002, 277, 11217-11224.	3.4	30
149	DIF-1 inhibits tumor growth in vivo reducing phosphorylation of GSK- $3\hat{l}^2$ and expressions of cyclin D1 and TCF7L2 in cancer model mice. Biochemical Pharmacology, 2014, 89, 340-348.	4.4	30
150	Characterization of the genomic structure and expression of the mouse Apex2 gene. Genomics, 2003, 81, 47-57.	2.9	30
151	Intracellular localization and function of DNA repair methyltransferase in human cells. Mutation Research DNA Repair, 1994, 315, 199-212.	3.7	29
152	Functional significance of conserved residues in the phosphohydrolase module of Escherichia coli MutT protein. Nucleic Acids Research, 2000, 28, 3240-3249.	14.5	29
153	Altered expression of MUTYH and an increase in 8â€hydroxydeoxyguanosine are early events in ulcerative colitisâ€associated carcinogenesis. Journal of Pathology, 2009, 219, 77-86.	4.5	29
154	Nucleotides function as endogenous chemical sensors for oxidative stress signaling. Journal of Clinical Biochemistry and Nutrition, 2010, 48, 33-39.	1.4	29
155	Prognostic impact of MutT homologâ€1 expression on esophageal squamous cell carcinoma. Cancer Medicine, 2017, 6, 258-266.	2.8	29
156	Oxidative Stress and Microglial Response in Retinitis Pigmentosa. International Journal of Molecular Sciences, 2020, 21, 7170.	4.1	29
157	In Vitro Development of Mouse Embryonic Stem Cells Lacking JNK/Stress-activated Protein Kinase-associated Protein 1 (JSAP1) Scaffold Protein Revealed Its Requirement during Early Embryonic Neurogenesis. Journal of Biological Chemistry, 2003, 278, 48422-48433.	3.4	28
158	Structural and Kinetic Studies of the Human Nudix Hydrolase MTH1 Reveal the Mechanism for Its Broad Substrate Specificity. Journal of Biological Chemistry, 2017, 292, 2785-2794.	3.4	28
159	8-oxoguanine DNA glycosylase (OGG1) deficiency elicits coordinated changes in lipid and mitochondrial metabolism in muscle. PLoS ONE, 2017, 12, e0181687.	2.5	28
160	Association of adipocyte enhancerâ€binding protein 1 with <scp>A</scp> lzheimer's disease pathology in human hippocampi. Brain Pathology, 2018, 28, 58-71.	4.1	28
161	D1-receptor-related priming is attenuated by antisense-meditated `knockdown' of fosB expression. Molecular Brain Research, 1998, 53, 69-77.	2.3	27
162	Probing the Substrate Recognition Mechanism of the Human MTH1 Protein by Nucleotide Analogs. Journal of Molecular Biology, 2004, 336, 843-850.	4.2	27

#	Article	IF	CITATIONS
163	Expression of Adipose Differentiation-Related Protein (ADRP) Is Conjointly Regulated by PU.1 and AP-1 in Macrophages. Journal of Biochemistry, 2005, 138, 399-412.	1.7	27
164	Biological Significance of the Defense Mechanisms against Oxidative Damage in Nucleic Acids Caused by Reactive Oxygen Species., 2004, 1011, 101-111.		27
165	Ischemia-induced CA1 neuronal death is preceded by elevated FosB and Jun expression and reduced NGFI-A and JunB levels. Molecular Brain Research, 1998, 56, 146-161.	2.3	26
166	Apurinic/Apyrimidinic Endonuclease 2 Is Necessary for Normal B Cell Development and Recovery of Lymphoid Progenitors after Chemotherapeutic Challenge. Journal of Immunology, 2011, 186, 1943-1950.	0.8	26
167	Nucleotide excision repair of oxidised genomic DNA is not a source of urinary 8-oxo-7,8-dihydro-2′-deoxyguanosine. Free Radical Biology and Medicine, 2016, 99, 385-391.	2.9	26
168	MUTYH promotes oxidative microglial activation and inherited retinal degeneration. JCI Insight, 2016, $1$ , e87781.	5.0	26
169	Identification and Biochemical Characterization of a Novel Transcription Elongation Factor, Elongin A3. Journal of Biological Chemistry, 2002, 277, 26444-26451.	3.4	25
170	fosB Gene Products Trigger Cell Proliferation and Morphological Alteration with an Increased Expression of a Novel Processed Form of Galectin-1 in the Rat 3Y1 Embroyo Cell Line. Journal of Biochemistry, 2002, 131, 653-661.	1.7	24
171	Clozapine-, but not haloperidol-, induced increases in î"FosB-like immunoreactivity are completely blocked in the striatum of mice lacking D3 dopamine receptors. European Journal of Neuroscience, 2004, 20, 3189-3194.	2.6	24
172	Accumulation of 8-oxo-deoxyguanosine in cardiovascular tissues with the development of hypertension. DNA Repair, 2007, 6, 760-769.	2.8	24
173	Cisplatin-Mediated Upregulation of APE2 Binding to MYH9 Provokes Mitochondrial Fragmentation and Acute Kidney Injury. Cancer Research, 2021, 81, 713-723.	0.9	24
174	Cloning and expression of the Bacillus subtilis methyltransferase gene in Escherichia coli adaâ^' cells. Mutation Research DNA Repair, 1989, 218, 153-163.	3.7	23
175	Polymorphisms and Probable Lack of Mutation in a Human mutT Homolog, hMTH1, in Hereditary Nonpoliposis Colorectal Cancer. Biochemical and Biophysical Research Communications, 1995, 214, 1239-1245.	2.1	23
176	Significance of the conserved amino acid sequence for human MTH1 protein with antimutator activity. Nucleic Acids Research, 1997, 25, 1170-1176.	14.5	23
177	FosB in rat striatum: Normal regional distribution and enhanced expression after 6-month haloperidol administration. Synapse, 2001, 39, 122-132.	1.2	23
178	î"FosB, but not FosB, induces delayed apoptosis independent of cell proliferation in the Rat1a embryo cell line. Cell Death and Differentiation, 2003, 10, 496-507.	11.2	23
179	Defense mechanism to oxidative DNA damage in glial cells. Neuropathology, 2004, 24, 125-130.	1.2	23
180	Mutagenic target for hydroxyl radicals generated in Escherichia coli mutant deficient in Mn- and Fe-superoxide dismutases and Fur, a repressor for iron-uptake systems. DNA Repair, 2002, 1, 411-418.	2.8	22

#	Article	IF	CITATIONS
181	Downâ€regulation of <scp>MET</scp> in hippocampal neurons of <scp>A</scp> lzheimer's disease brains. Neuropathology, 2014, 34, 284-290.	1.2	22
182	The Disease-modifying Drug Candidate, SAK3 Improves Cognitive Impairment and Inhibits Amyloid beta Deposition in App Knock-in Mice. Neuroscience, 2018, 377, 87-97.	2.3	22
183	Expression of the truncated E. coliO6-methylguanine methyltransferase gene in repair-deficient human cells and restoration of cellular resistance to alkylating agents. Mutation Research - DNA Repair Reports, 1987, 184, 121-128.	1.8	21
184	A comparative immunohistochemistry of O6-methylguanine-DNA methyltransferase and p53 in diffusely infiltrating astrocytomas. Neuropathology, 2003, 23, 203-209.	1.2	21
185	Mouse RS21â€C6 is a mammalian 2′â€deoxycytidine 5′â€triphosphate pyrophosphohydrolase that prefers 5â€iodocytosine. FEBS Journal, 2009, 276, 1654-1666.	4.7	21
186	Association of MTH1 expression with the tumor malignant potential and poor prognosis in patients with resected lung cancer. Lung Cancer, 2017, 109, 52-57.	2.0	21
187	Characterization of the Structure and Expression of Mouse Itpa Gene and its Related Sequences in the Mouse Genome. DNA Research, 2005, 12, 39-51.	3.4	20
188	Impaired spermatogenesis and elevated spontaneous tumorigenesis in xeroderma pigmentosum group A gene (Xpa)-deficient mice. DNA Repair, 2008, 7, 1938-1950.	2.8	20
189	Cancer-Related PRUNE2 Protein Is Associated with Nucleotides and Is Highly Expressed in Mature Nerve Tissues. Journal of Molecular Neuroscience, 2011, 44, 103-114.	2.3	20
190	Silencing of SNX1 by siRNA stimulates the ligand-induced endocytosis of EGFR and increases EGFR phosphorylation in gefitinib-resistant human lung cancer cell lines. International Journal of Oncology, 2012, 41, 1520-1530.	3.3	20
191	Serum galectin-3, but not galectin-1, levels are elevated in schizophrenia: implications for the role of inflammation. Psychopharmacology, 2017, 234, 2919-2927.	3.1	20
192	A Novel Autoantibody against Plexin <scp>D</scp> 1 in Patients with Neuropathic Pain. Annals of Neurology, 2018, 84, 208-224.	5.3	20
193	A variant form of hMTH1, a human homologue of the E coli mutT gene, correlates with somatic mutation in the p53 tumour suppressor gene in gastric cancer patients. Journal of Medical Genetics, 2004, 41, e57-e57.	3.2	19
194	The human HYMAI/PLAGL1 differentially methylated region acts as an imprint control region in mice. Genomics, 2006, 88, 650-658.	2.9	19
195	Neonatal Ventral Hippocampal Lesions Produce an Elevation of ΔFosB-Like Protein(s) in the Rodent Neocortex. Neuropsychopharmacology, 2006, 31, 700-711.	5.4	19
196	Skin tumours induced by narrowband UVB have higher frequency of p53 mutations than tumours induced by broadband UVB independent of Ogg1 genotype. Mutagenesis, 2012, 27, 637-643.	2.6	19
197	Complexity of Stomach–Brain Interaction Induced by Molecular Hydrogen in Parkinson's Disease Model Mice. Neurochemical Research, 2017, 42, 2658-2665.	3.3	19
198	PCBP1 and PCBP2 both bind heavily oxidized RNA but cause opposing outcomes, suppressing or increasing apoptosis under oxidative conditions. Journal of Biological Chemistry, 2020, 295, 12247-12261.	3.4	19

#	Article	IF	CITATIONS
199	A highâ€fat diet exacerbates the Alzheimer's disease pathology in the hippocampus of the <i>App<i><i><i><i><i><i><i><i><i><i><i><i< td=""><td>6.7</td><td>19</td></i<></i></i></i></i></i></i></i></i></i></i></i></i>	6.7	19
200	Immediate-early gene expression in the brain of the thiamine-deficient rat. Journal of Molecular Neuroscience, 1998, 10, 1-15.	2.3	18
201	Galectinâ€1 deficiency improves axonal swelling of motor neurones in <scp>SOD</scp> 1 <sup>G93A</sup> transgenic mice. Neuropathology and Applied Neurobiology, 2015, 41, 227-244.	3.2	18
202	Differentiation-inducing factor-3 inhibits intestinal tumor growth inÂvitro and inÂvivo. Journal of Pharmacological Sciences, 2015, 127, 446-455.	2.5	18
203	Hyperactive mTOR signals in the proopiomelanocortin-expressing hippocampal neurons cause age-dependent epilepsy and premature death in mice. Scientific Reports, 2016, 6, 22991.	3.3	18
204	Molecular Pathophysiology of Insulin Depletion, Mitochondrial Dysfunction, and Oxidative Stress in Alzheimer's Disease Brain. Advances in Experimental Medicine and Biology, 2019, 1128, 27-44.	1.6	18
205	OGG1 deficiency alters the intestinal microbiome and increases intestinal inflammation in a mouse model. PLoS ONE, 2020, 15, e0227501.	2.5	18
206	MTH1 and OGG1 maintain a low level of 8-oxoguanine in Alzheimer's brain, and prevent the progression of Alzheimer's pathogenesis. Scientific Reports, 2021, 11, 5819.	3.3	18
207	Role ofc-junin the Inhibition of Erythropoietin Receptor-Mediated Apoptosis. Biochemical and Biophysical Research Communications, 1996, 222, 1-6.	2.1	17
208	Mammalian Elongin A Is Not Essential for Cell Viability but Is Required for Proper Cell Cycle Progression with Limited Alteration of Gene Expression. Journal of Biological Chemistry, 2003, 278, 13585-13589.	3.4	17
209	Regulation of the Neuronal Fate by ΔFosB and its Downstream Target, Galectin-1. Current Drug Targets, 2005, 6, 437-444.	2.1	17
210	Abnormality in Wnt Signaling is Causatively Associated with Oxidative Stress-Induced Intestinal Tumorigenesis in MUTYH-Null Mice. International Journal of Biological Sciences, 2014, 10, 940-947.	6.4	17
211	MUTYH Actively Contributes to Microglial Activation and Impaired Neurogenesis in the Pathogenesis of Alzheimer's Disease. Oxidative Medicine and Cellular Longevity, 2021, 2021, 1-30.	4.0	17
212	A unique structural feature of rabbit DNA repair methyltransferase as revealed by cDNA cloning. Carcinogenesis, 1994, 15, 627-633.	2.8	16
213	8-Chloro-dGTP, a hypochlorous acid-modified nucleotide, is hydrolyzed by hMTH1, the human MutT homolog. FEBS Letters, 2002, 512, 149-151.	2.8	15
214	Some flavonoids and DHEA-S prevent the cis-effect of expanded CTG repeats in a stable PC12 cell transformant. Biochemical Pharmacology, 2005, 69, 503-516.	4.4	15
215	Induction of apoptosis and cellular senescence in mice lacking transcription elongation factor, Elongin A. Cell Death and Differentiation, 2007, 14, 716-726.	11.2	15
216	Antagonistic Regulation of Cell-Matrix Adhesion by FosB and Î"FosB Î"2Î"FosB Encoded by Alternatively Spliced Forms of <i>fosB</i> Transcripts. Molecular Biology of the Cell, 2008, 19, 4717-4729.	2.1	15

#	Article	IF	Citations
217	Apurinic/Apyrimidinic Endonuclease 2 Regulates the Expansion of Germinal Centers by Protecting against Activation-Induced Cytidine Deaminase–Independent DNA Damage in B Cells. Journal of Immunology, 2014, 193, 931-939.	0.8	15
218	Deoxyinosine triphosphate induces MLH1/PMS2- and p53-dependent cell growth arrest and DNA instability in mammalian cells. Scientific Reports, 2016, 6, 32849.	3.3	15
219	Fenton reactionâ€induced renal carcinogenesis in <i>Mutyh</i> â€deficient mice exhibits less chromosomal aberrations than the rat model. Pathology International, 2017, 67, 564-574.	1.3	14
220	<i>GNAO1</i> organizes the cytoskeletal remodeling and firing of developing neurons. FASEB Journal, 2020, 34, 16601-16621.	0.5	14
221	Structure of the mammalian adenine DNA glycosylase MUTYH: insights into the base excision repair pathway and cancer. Nucleic Acids Research, 2021, 49, 7154-7163.	14.5	14
222	Therapeutic Effects of Hydrogen in Animal Models of Parkinson's Disease. Parkinson's Disease, 2011, 2011, 1-9.	1.1	13
223	Environmental aircraft noise aggravates oxidative DNA damage, granulocyte oxidative burst and nitrate resistance in <i>Ogg1</i> aê"/–mice. Free Radical Research, 2020, 54, 280-292.	3.3	12
224	Accumulation of Adenine DNA Glycosylase-sensitive Sites in Human Mitochondrial DNA. Journal of Biological Chemistry, 2000, 275, 12326-12330.	3.4	11
225	The GT to GC single nucleotide polymorphism at the beginning of an alternative exon 2C of human MTH1 gene confers an amino terminal extension that functions as a mitochondrial targeting signal. Journal of Molecular Medicine, 2006, 84, 660-670.	3.9	11
226	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
227	Role of the DNA repair glycosylase OGG1 in the activation of murine splenocytes. DNA Repair, 2017, 58, 13-20.	2.8	11
228	Oxidative stress induces different tissue dependent effects on Mutyh-deficient mice. Free Radical Biology and Medicine, 2019, 143, 482-493.	2.9	11
229	Origins of Brain Insulin and Its Function. Advances in Experimental Medicine and Biology, 2019, 1128, 1-11.	1.6	11
230	Crystallization and preliminary X-ray diffraction studies of 3-methyladenine—DNA glycosylase II from Escherichia coli. Journal of Molecular Biology, 1988, 204, 1055-1056.	4.2	10
231	Intracellular distribution of the antimutagenic enzyme MTH1 in the liver, kidney and testis of F344 ratsand its modulation by cadmium. Experimental and Toxicologic Pathology, 2001, 53, 325-335.	2.1	10
232	Role of Tryptophan Residues in the Recognition of Mutagenic Oxidized Nucleotides by Human Antimutator MTH1 Protein. Journal of Molecular Biology, 2002, 319, 129-139.	4.2	10
233	Impairment of cardiomyogenesis in embryonic stem cells lacking scaffold protein JSAP1. Biochemical and Biophysical Research Communications, 2005, 338, 1152-1157.	2.1	10
234	Characterization of galectin-1-positive cells in the mouse hippocampus. NeuroReport, 2014, 25, 171-176.	1.2	10

#	Article	IF	CITATIONS
235	8-Oxoguanine accumulation in aged female brain impairs neurogenesis in the dentate gyrus and major island of Calleja, causing sexually dimorphic phenotypes. Progress in Neurobiology, 2019, 180, 101613.	5.7	10
236	Changes in expression of $\hat{l}$ FosB and the Fos family proteins following NMDA receptor activation in the rat striatum. Molecular Brain Research, 1997, 47, 31-43.	2.3	9
237	Enhancement of Laminar FosB Expression in Frontal Cortex of Rats Receiving Long Chronic Clozapine Administration. Experimental Neurology, 2001, 168, 392-401.	4.1	9
238	Angiotensin I-converting enzyme gene polymorphism modifies the smoking–cancer association: the Hisayama Study. European Journal of Cancer Prevention, 2006, 15, 196-201.	1.3	9
239	Altered gene expression profiles and higher frequency of spontaneous DNA strand breaks in APEX2-null thymus. DNA Repair, 2008, 7, 1437-1454.	2.8	9
240	A comprehensive screening system for damaged nucleotide-binding proteins. Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2010, 703, 37-42.	1.7	9
241	An intronic single nucleotide polymorphism in the MUTYH gene is associated with increased risk for HCV-induced hepatocellular carcinoma. Free Radical Biology and Medicine, 2018, 129, 88-96.	2.9	9
242	Action of T4 Endonuclease V on Polydeoxyribonucleotides with Apyrimidinic or Apurinic Sites1. Journal of Biochemistry, 1982, 91, 2057-2065.	1.7	8
243	Recognition of Nucleotide Analogs Containing the 7,8-Dihydro-8-oxo Structure by the Human MTH1 Protein. Journal of Biochemistry, 2006, 140, 843-849.	1.7	8
244	APE2 Is a General Regulator of the ATR-Chk1 DNA Damage Response Pathway to Maintain Genome Integrity in Pancreatic Cancer Cells. Frontiers in Cell and Developmental Biology, 2021, 9, 738502.	3.7	8
245	Crystallization and preliminary X-ray analysis of human MTH1 complexed with two oxidized nucleotides, 8-oxo-dGMP and 2-oxo-dATP. Acta Crystallographica Section F: Structural Biology Communications, 2006, 62, 1283-1285.	0.7	7
246	A Role for SNX1 in the Regulation of EGF-Dependent Phosphorylated EGFR Endocytosis Via the Early/Late Endocytic Pathway in a Gefitinib-Sensitive Human Lung Cancer Cells. Current Signal Transduction Therapy, 2011, 6, 383-395.	0.5	7
247	Expression of CRYM in different rat organs during development and its decreased expression in degenerating pyramidal tracts in amyotrophic lateral sclerosis. Neuropathology, 2018, 38, 247-259.	1.2	7
248	Neuroendocrine phenotypes in a boy with 5q14 deletion syndrome implicate the regulatory roles of myocyte-specific enhancer factor 2C in the postnatal hypothalamus. European Journal of Medical Genetics, 2013, 56, 475-483.	1.3	6
249	Prevention of the Mutagenicity and Cytotoxicity of Oxidized Purine Nucleotides., 2007,, 40-53.		6
250	Detection of Angiotensin-Converting Enzyme Gene Insertion/Deletion Polymorphism From Paraffin-Embedded Tissues. The Hisayama Study. Circulation Journal, 2002, 66, 1034-1036.	1.6	5
251	<scp>PKC</scp> η deficiency improves lipid metabolism and atherosclerosis in apolipoprotein <scp>E</scp> â€deficient mice. Genes To Cells, 2016, 21, 1030-1048.	1.2	5
252	Co-regulation of Cxcl1 and versican in the inflammatory response to UVB induced reactive oxygen species in skin photo-tumorigenesis. Journal of Dermatological Science, 2017, 85, 140-143.	1.9	5

#	Article	IF	Citations
253	<i>Mth1</i> deficiency provides longer survival upon intraperitoneal crocidolite injection in female mice. Free Radical Research, 2020, 54, 195-205.	3.3	5
254	MUTYH is associated with hepatocarcinogenesis in a non-alcoholic steatohepatitis mouse model. Scientific Reports, 2021, 11, 3599.	3.3	5
255	Neurodegeneration Caused by Accumulation of an Oxidized Base Lesion, 8-oxoguanine, in Nuclear and Mitochondrial DNA: From Animal Models to Human Diseases. , 2017, , 523-556.		5
256	Neural stem cell–specific ITPA deficiency causes neural depolarization and epilepsy. JCI Insight, 2020, 5,	5.0	5
257	Crystallization and preliminary X-ray analysis of human MTH1 with a homogeneous N-terminus. Acta Crystallographica Section F: Structural Biology Communications, 2013, 69, 45-48.	0.7	4
258	Serum Anti-oligodendrocyte Autoantibodies in Patients With Multiple Sclerosis Detected by a Tissue-Based Immunofluorescence Assay. Frontiers in Neurology, 2021, 12, 681980.	2.4	3
259	Construction and Characterization of a Cell Line Deficient in Repair of Mitochondrial, but Not Nuclear, Oxidative DNA Damage. Methods in Molecular Biology, 2009, 554, 251-264.	0.9	3
260	Deficiency of MTH1 and/or OGG1 increases the accumulation of 8-oxoguanine in the brain of the AppNL-G-F/NL-G-F knock-in mouse model of Alzheimer's disease, accompanied by accelerated microgliosis and reduced anxiety-like behavior. Neuroscience Research, 2022, 177, 118-134.	1.9	3
261	APE1- and APE2-dependent DNA breaks in immunoglobulin class switch recombination. Journal of Experimental Medicine, 2007, 204, 3295-3295.	8.5	2
262	OXIDATIVE STRESS-INDUCED TUMORIGENESIS IN THE SMALL INTESTINES OF DNA REPAIR-DEFICIENT MICE. Health Physics, 2011, 100, 293-294.	0.5	2
263	Nature of nontargeted radiation effects observed during fractionated irradiation-induced thymic lymphomagenesis in mice. Journal of Radiation Research, 2013, 54, 453-466.	1.6	2
264	2-Oxoadenosine induces cytotoxicity through intracellular accumulation of 2-oxo-ATP and depletion of ATP but not via the p38 MAPK pathway. Scientific Reports, 2017, 7, 6528.	3.3	2
265	MUTYH Deficiency Is Associated with Attenuated Pulmonary Fibrosis in a Bleomycin-Induced Model. Oxidative Medicine and Cellular Longevity, 2020, 2020, 1-14.	4.0	2
266	Galectin-1 and galectin-3 as key molecules for peripheral nerve degeneration and regeneration. AIMS Molecular Science, 2016, 3, 325-337.	0.5	2
267	Transcriptome Analysis in Hippocampus of Rats Prenatally Exposed to Valproic Acid and Effects of Intranasal Treatment of Oxytocin. Frontiers in Psychiatry, 2022, 13, 859198.	2.6	2
268	Expression and Prognostic Significance of O6-Methylguanine-DNA Methyltransferase in Hepatocellular, Gastric, and Breast Cancers. Annals of Surgical Oncology, 2001, 8, 807-816.	1.5	1
269	Mutyh deficiency downregulates mitochondrial fusion proteins and causes cardiac dysfunction via $\hat{l}\pm$ -ketoglutaric acid reduction with oxidative stress. Free Radical Research, 2022, , 1-16.	3.3	1
270	8-Oxoguanine DNA Glycosylase (OGG1) Deficiency Exacerbates Doxorubicin-Induced Cardiac Dysfunction. Oxidative Medicine and Cellular Longevity, 2022, 2022, 1-11.	4.0	1

#	Article	IF	CITATIONS
271	Possible target gene abnormality in 06-methyl-guanine-DNA methyltransferase (MGMT) deficient phenotype in gall bladder carcinoma. Gastroenterology, 2000, 118, A1458.	1.3	O
272	Galectin-1 promotes neurogenesis in the dentate gyrus of mouse hippocampus after brain damage caused by excitotoxicity. Neuroscience Research, 2007, 58, S210.	1.9	0
273	fosB-null mice exhibit impaired adult hippocampal neurogenesis and spontaneous epileptic seizures. Neuroscience Research, 2010, 68, e419.	1.9	O
274	î"FosB and/or î"2î"FosB regulate proliferation of adult hippocampal neural progenitor cells and suppress spontaneous epileptic seizures. Neuroscience Research, 2011, 71, e295.	1.9	0
275	International Symposium on "Germline Mutagenesis and Biodiversification― Genes and Genetic Systems, 2014, 89, 93-95.	0.7	0
276	Requirement for two conserved cysteine residues in the Ada protein of. Molecular Genetics and Genomics, 1996, 250, 523.	2.4	0