

Baojin Ding

List of Publications by Year in descending order

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Version: 2024-02-01

36
papers

948
citations

516710

16
h-index

477307

29
g-index

38
all docs

38
docs citations

38
times ranked

1367
citing authors

#	ARTICLE	IF	CITATIONS
1	Consensus Paper: Cerebellar Development. <i>Cerebellum</i> , 2016, 15, 789-828.	2.5	337
2	IscA Mediates Iron Delivery for Assembly of Iron-Sulfur Clusters in IscU under the Limited Accessible Free Iron Conditions. <i>Journal of Biological Chemistry</i> , 2004, 279, 37499-37504.	3.4	91
3	Mobilization of the iron centre in IscA for the iron-sulphur cluster assembly in IscU. <i>Biochemical Journal</i> , 2005, 389, 797-802.	3.7	53
4	The C-terminal Repeat Domain of Spt5 Plays an Important Role in Suppression of Rad26-independent Transcription Coupled Repair. <i>Journal of Biological Chemistry</i> , 2010, 285, 5317-5326.	3.4	48
5	Lentiviral Vector Production, Titration, and Transduction of Primary Neurons. <i>Methods in Molecular Biology</i> , 2013, 1018, 119-131.	0.9	36
6	Temporal Regulation of Nuclear Factor One Occupancy by Calcineurin/NFAT Governs a Voltage-Sensitive Developmental Switch in Late Maturing Neurons. <i>Journal of Neuroscience</i> , 2013, 33, 2860-2872.	3.6	33
7	Lamin Mutations Accelerate Aging via Defective Export of Mitochondrial mRNAs through Nuclear Envelope Budding. <i>Current Biology</i> , 2016, 26, 2052-2059.	3.9	32
8	Disease Modeling with Human Neurons Reveals LMNB1 Dysregulation Underlying DYT1 Dystonia. <i>Journal of Neuroscience</i> , 2021, 41, 2024-2038.	3.6	32
9	Evidence that the Transcription Elongation Function of Rpb9 Is Involved in Transcription-Coupled DNA Repair in <i>Saccharomyces cerevisiae</i> . <i>Molecular and Cellular Biology</i> , 2006, 26, 9430-9441.	2.3	28
10	Nucleus to Synapse Nesprin1 Railroad Tracks Direct Synapse Maturation through RNA Localization. <i>Neuron</i> , 2015, 86, 1015-1028.	8.1	27
11	The roles of Rad16 and Rad26 in repairing repressed and actively transcribed genes in yeast. <i>DNA Repair</i> , 2007, 6, 1596-1606.	2.8	25
12	Nucleocytoplasmic Transport: Regulatory Mechanisms and the Implications in Neurodegeneration. <i>International Journal of Molecular Sciences</i> , 2021, 22, 4165.	4.1	25
13	Reciprocal autoregulation by NFI occupancy and ETV1 promotes the developmental expression of dendrite-synapse genes in cerebellar granule neurons. <i>Molecular Biology of the Cell</i> , 2016, 27, 1488-1499.	2.1	21
14	Yeast Elc1 plays an important role in global genomic repair but not in transcription coupled repair. <i>DNA Repair</i> , 2009, 8, 40-50.	2.8	20
15	Generation and optimization of highly pure motor neurons from human induced pluripotent stem cells via lentiviral delivery of transcription factors. <i>American Journal of Physiology - Cell Physiology</i> , 2020, 319, C771-C780.	4.6	19
16	Rpb1 Sumoylation in Response to UV Radiation or Transcriptional Impairment in Yeast. <i>PLoS ONE</i> , 2009, 4, e5267.	2.5	18
17	Modulation of Rad26- and Rpb9-mediated DNA Repair by Different Promoter Elements. <i>Journal of Biological Chemistry</i> , 2006, 281, 36643-36651.	3.4	13
18	Differential Influence of Sample Sex and Neuronal Maturation on mRNA and Protein Transport in Induced Human Neurons. <i>Frontiers in Molecular Neuroscience</i> , 2020, 13, 46.	2.9	13

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19	BDNF activates an NFI-dependent neurodevelopmental timing program by sequestering NFATc4. <i>Molecular Biology of the Cell</i> , 2018, 29, 975-987.	2.1	12
20	Evidence that Moderate Eviction of Spt5 and Promotion of Error-Free Transcriptional Bypass by Rad26 Facilitates Transcription Coupled Nucleotide Excision Repair. <i>Journal of Molecular Biology</i> , 2019, 431, 1322-1338.	4.2	11
21	Gene expression in maturing neurons: regulatory mechanisms and related neurodevelopmental disorders. <i>Acta Physiologica Sinica</i> , 2015, 67, 113-33.	0.5	10
22	Generation of highly pure motor neurons from human induced pluripotent stem cells. <i>STAR Protocols</i> , 2022, 3, 101223.	1.2	8
23	Tfb5 is partially dispensable for Rad26 mediated transcription coupled nucleotide excision repair in yeast. <i>DNA Repair</i> , 2007, 6, 1661-1669.	2.8	6
24	Generation of patient-specific motor neurons in modeling movement diseases. <i>Neural Regeneration Research</i> , 2021, 16, 1799.	3.0	6
25	Direct conversion of adult fibroblasts into motor neurons. <i>STAR Protocols</i> , 2021, 2, 100917.	1.2	6
26	Generation of two induced pluripotent stem cell lines with heterozygous and homozygous GAG deletion in TOR1A gene from a healthy hiPSC line. <i>Stem Cell Research</i> , 2021, 56, 102536.	0.7	5
27	Novel insights into the pathogenesis of DYT1 dystonia from induced patient-derived neurons. <i>Neural Regeneration Research</i> , 2022, 17, 561.	3.0	4
28	Generation of gene-corrected isogenic control cell lines from a DYT1 dystonia patient iPSC line carrying a heterozygous GAG mutation in TOR1A gene. <i>Stem Cell Research</i> , 2022, 62, 102807.	0.7	2
29	Chromatin Immunoprecipitation Assay of Brain Tissues Using Percoll Gradient-Purified Nuclei. <i>Methods in Molecular Biology</i> , 2013, 1018, 199-209.	0.9	1
30	How to assist parents of children with autism spectrum disorders in rural area?. <i>Journal of Neurosciences in Rural Practice</i> , 2015, 06, 465-466.	0.8	0
31	Rpb1 Sumoylation in Response to UV Radiation or Transcriptional Impairment in Yeast. <i>FASEB Journal</i> , 2009, 23, 701.1.	0.5	0
32	Yeast Elc1 plays an important role in global genomic repair but not in transcription coupled repair. <i>FASEB Journal</i> , 2009, 23, 836.4.	0.5	0
33	Spt4 and Spt5 cooperatively suppress transcription coupled DNA repair through binding to RNA polymerase II in the absence of Rad26. <i>FASEB Journal</i> , 2009, 23, 836.6.	0.5	0
34	Nuclear factor one controls a voltage-sensitive developmental switch required for late neuronal maturation. <i>FASEB Journal</i> , 2013, 27, 535.3.	0.5	0
35	Auto-regulatory interactions between NFI occupancy and ETV1 direct the timing of gene expression in late maturing neurons (539.10). <i>FASEB Journal</i> , 2014, 28, 539.10.	0.5	0
36	How does a 1.5-Fold Increase in Gene Dosage in Chromosome 21 Causes the Pleiotropic Phenotypes in Down Syndrome?. <i>Journal of Down Syndrome & Chromosome Abnormalities</i> , 2015, 1, .	0.1	0