

Li Wang

List of Publications by Year in descending order

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Version: 2024-02-01

16
papers

355
citations

1039406

9
h-index

940134

16
g-index

16
all docs

16
docs citations

16
times ranked

514
citing authors

#	ARTICLE	IF	CITATIONS
1	Relation between hypomethylation of long interspersed nucleotide elements and risk of neural tube defects. <i>American Journal of Clinical Nutrition</i> , 2010, 91, 1359-1367.	2.2	145
2	Elevated H3K79 homocysteinylation causes abnormal gene expression during neural development and subsequent neural tube defects. <i>Nature Communications</i> , 2018, 9, 3436.	5.8	56
3	Association between <i>TCF7L2</i> polymorphisms and gestational diabetes mellitus: A meta-analysis. <i>Journal of Diabetes Investigation</i> , 2017, 8, 560-570.	1.1	23
4	Altered Methylation of the DNA Repair Gene MGMT Is Associated with Neural Tube Defects. <i>Journal of Molecular Neuroscience</i> , 2012, 47, 42-51.	1.1	18
5	High expression levels of microRNA-629, microRNA-525-5p and microRNA-516a-3p in paediatric systemic lupus erythematosus. <i>Clinical Rheumatology</i> , 2014, 33, 807-815.	1.0	18
6	Altered GNAS imprinting due to folic acid deficiency contributes to poor embryo development and may lead to neural tube defects. <i>Oncotarget</i> , 2017, 8, 110797-110810.	0.8	18
7	The effect of folic acid deficiency on FGF pathway via Brachyury regulation in neural tube defects. <i>FASEB Journal</i> , 2019, 33, 4688-4702.	0.2	15
8	Folate deficiency disturbs hsalet-7 g level through methylation regulation in neural tube defects. <i>Journal of Cellular and Molecular Medicine</i> , 2017, 21, 3244-3253.	1.6	14
9	Sonic Hedgehog Signaling Affected by Promoter Hypermethylation Induces Aberrant Gli2 Expression in Spina Bifida. <i>Molecular Neurobiology</i> , 2016, 53, 5413-5424.	1.9	13
10	Imprinting aberrations of <i>SNRPN</i> , <i>ZAC1</i> and <i>INPP5F</i> genes involved in the pathogenesis of congenital heart disease with extracardiac malformations. <i>Journal of Cellular and Molecular Medicine</i> , 2020, 24, 9898-9907.	1.6	11
11	Folate deficiency disturbs PEG10 methylation modifications in human spina bifida. <i>Pediatric Research</i> , 2022, 92, 987-994.	1.1	7
12	The effect of folic acid deficiency on Mest/Peg1 in neural tube defects. <i>International Journal of Neuroscience</i> , 2021, 131, 468-477.	0.8	6
13	Effects of MTHFR A1298C polymorphism on peripheral blood folate concentration in healthy populations: a meta-analysis of observational studies. <i>Asia Pacific Journal of Clinical Nutrition</i> , 2018, 27, 718-727.	0.3	4
14	Polimorfizm rs10830963 w genie receptora melatoniny 1B a cukrzyca ciążowa w populacji chińskiej: metaanaliza badań, asocjacyjnych. <i>Endokrynologia Polska</i> , 2017, 68, 550-560.	0.3	4
15	The association of endothelial nitric oxide synthase gene single nucleotide polymorphisms with paediatric systemic lupus erythematosus. <i>Clinical and Experimental Rheumatology</i> , 2018, 36, 508-512.	0.4	2
16	Relationship between HLA-DPA1 genetic polymorphism and anembryonic pregnancy. <i>Molecular Genetics & Genomic Medicine</i> , 2020, 8, e1046.	0.6	1