

Chunhua Zeng

List of Publications by Year in descending order

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papers

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840776

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33
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580
citing authors

#	ARTICLE	IF	CITATIONS
1	Two De Novo Mosaic Variants Within the Same Site of PHEX Gene in a Girl with X-Linked Hypophosphatemic Rickets. <i>Calcified Tissue International</i> , 2022, 110, 266-271.	3.1	0
2	Assessment of the diagnostic value of serum ceruloplasmin for Wilson's disease in children. <i>BMC Gastroenterology</i> , 2022, 22, 124.	2.0	3
3	A novel variant in SLC16A2 associated with typical Allan-Herndon-Dudley syndrome: a case report. <i>BMC Pediatrics</i> , 2022, 22, 180.	1.7	6
4	Thyroid function in children with Prader-Willi syndrome in Southern China: a single-center retrospective case series. <i>BMC Pediatrics</i> , 2022, 22, 234.	1.7	0
5	The EGFR-p38 MAPK axis up-regulates PD-L1 through miR-675-5p and down-regulates HLA-ABC via hexokinase-2 in hepatocellular carcinoma cells. <i>Cancer Communications</i> , 2021, 41, 62-78.	9.2	28
6	A GM1 gangliosidosis mutant mouse model exhibits activated microglia and disturbed autophagy. <i>Experimental Biology and Medicine</i> , 2021, 246, 1330-1341.	2.4	16
7	Molecular and clinical characteristics of monogenic diabetes mellitus in southern Chinese children with onset before 3 years of age. <i>BMJ Open Diabetes Research and Care</i> , 2020, 8, e001345.	2.8	6
8	Chinese family with Blau syndrome: Mutated <i>NOD2</i> allele transmitted from the father with de novo somatic and germ line mosaicism. <i>Journal of Dermatology</i> , 2020, 47, e395.	1.2	2
9	Distinct severity of phenotype in Hajdu-Cheney syndrome: a case report and literature review. <i>BMC Musculoskeletal Disorders</i> , 2020, 21, 154.	1.9	9
10	High risk screening for Gaucher disease in patients with splenomegaly and/or thrombocytopenia in China: 55 cases identified. <i>Clinica Chimica Acta</i> , 2020, 506, 22-27.	1.1	16
11	Molecular diagnosis of maturity-onset diabetes of the young in a cohort of Chinese children. <i>Pediatric Diabetes</i> , 2020, 21, 431-440.	2.9	11
12	Isolated germline mosaicism in the phenotypically normal father of a girl with X-linked hypophosphatemic rickets. <i>European Journal of Endocrinology</i> , 2020, 182, K1-K6.	3.7	9
13	Clinical features and mutational analysis in 114 young children with Wilson disease from South China. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 1451-1458.	1.2	9
14	A novel homozygous splice-site variant of <i>NCAPD2</i> gene identified in two siblings with primary microcephaly: The second case report. <i>Clinical Genetics</i> , 2019, 96, 98-101.	2.0	2
15	The Enamel Phenotype in Homozygous <i>Fam83h</i> Truncation Mice. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e724.	1.2	16
16	Clinical features, molecular characteristics, and treatments of a Chinese girl with sitosterolemia: A case report and literature review. <i>Journal of Clinical Lipidology</i> , 2019, 13, 246-250.	1.5	9
17	A novel GTPCH deficiency mouse model exhibiting tetrahydrobiopterin-related metabolic disturbance and infancy-onset motor impairments. <i>Metabolism: Clinical and Experimental</i> , 2019, 94, 96-104.	3.4	14
18	Two novel mutations in the <i>ALPL</i> gene of unrelated Chinese children with Hypophosphatasia: case reports and literature review. <i>BMC Pediatrics</i> , 2019, 19, 456.	1.7	10

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19	Clinical Management and Gene Mutation Analysis of Children with Congenital Hyperinsulinism in South China. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2019, 11, 400-409.	0.9	7
20	Somatic and germline <i>FOXP3</i> mosaicism in the mother of a boy with IPEX syndrome. <i>European Journal of Immunology</i> , 2018, 48, 885-887.	2.9	6
21	Clinical features, <i>BTD</i> gene mutations, and their functional studies of eight symptomatic patients with biotinidase deficiency from Southern China. <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 589-596.	1.2	11
22	Somatosensory Projections to Cochlear Nucleus Are Upregulated after Unilateral Deafness. <i>Journal of Neuroscience</i> , 2012, 32, 15791-15801.	3.6	84
23	Cuneate and spinal trigeminal nucleus projections to the cochlear nucleus are differentially associated with vesicular glutamate transporter-2. <i>Neuroscience</i> , 2011, 176, 142-151.	2.3	34
24	Vesicular Glutamate Transporter 2 Is Associated with the Cochlear Nucleus Commissural Pathway. <i>JARO - Journal of the Association for Research in Otolaryngology</i> , 2010, 11, 675-687.	1.8	9
25	Cochlear Damage Changes the Distribution of Vesicular Glutamate Transporters Associated with Auditory and Nonauditory Inputs to the Cochlear Nucleus. <i>Journal of Neuroscience</i> , 2009, 29, 4210-4217.	3.6	103
26	Tetrathiomolybdate Is Partially Protective Against Hyperglycemia in Rodent Models of Diabetes. <i>Experimental Biology and Medicine</i> , 2008, 233, 1021-1025.	2.4	8
27	Antitumor and antiinflammatory effects of tetrathiomolybdate in comparison with tetrathiomolybdate. <i>Translational Research</i> , 2007, 149, 260-264.	5.0	11
28	Comparison of lowering copper levels with tetrathiomolybdate and zinc on mouse tumor and doxorubicin models. <i>Translational Research</i> , 2006, 148, 309-314.	5.0	7
29	The use of tetrathiomolybdate in treating fibrotic, inflammatory, and autoimmune diseases, including the non-obese diabetic mouse model. <i>Journal of Inorganic Biochemistry</i> , 2006, 100, 927-930.	3.5	28