Lili Liang

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/861840/publications.pdf

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17	220	1040056	1058476
papers	citations	9	14
		h-index	g-index

17	17	17	329
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Long noncoding RNA ZFAS1 promotes tumorigenesis through regulation of miR-150-5p/RAB9A in melanoma. Melanoma Research, 2019, 29, 569-581.	1.2	35
2	Genetic Evaluation of 114 Chinese Short Stature Children in the Next Generation Era: a Single Center Study. Cellular Physiology and Biochemistry, 2018, 49, 295-305.	1.6	28
3	Mutation spectrum of hyperphenylalaninemia candidate genes and the genotype-phenotype correlation in the Chinese population. Clinica Chimica Acta, 2018, 481, 132-138.	1.1	24
4	Gambogic Acid Inhibits Melanoma through Regulation of miRâ€199aâ€3p/ZEB1 Signalling. Basic and Clinical Pharmacology and Toxicology, 2018, 123, 692-703.	2.5	24
5	High Detection Rate of Copy Number Variations Using Capture Sequencing Data: A Retrospective Study. Clinical Chemistry, 2020, 66, 455-462.	3.2	16
6	Prenatal diagnosis of methylmalonic aciduria from amniotic fluid using genetic and biochemical approaches. Prenatal Diagnosis, 2019, 39, 993-997.	2.3	14
7	TWEAK/Fn14 Interaction Confers AggressiveÂProperties to Cutaneous SquamousÂCell Carcinoma. Journal of Investigative Dermatology, 2019, 139, 796-806.	0.7	14
8	Clinical, genetic profile and therapy evaluation of 55 children and 5 adults with sitosterolemia. Journal of Clinical Lipidology, 2022, 16, 40-51.	1.5	10
9	Prenatal Diagnosis of Glutaric Acidemia I Based on Amniotic Fluid Samples in 42 Families Using Genetic and Biochemical Approaches. Frontiers in Genetics, 2020, 11, 496.	2.3	9
10	Clinical, Biochemical, and Molecular Analyses of Medium-Chain Acyl-CoA Dehydrogenase Deficiency in Chinese Patients. Frontiers in Genetics, 2021, 12, 577046.	2.3	9
11	Tumor Necrosis Factor (TNF) Receptor Expression Determines Keratinocyte Fate upon Stimulation with TNF-Like Weak Inducer of Apoptosis. Mediators of Inflammation, 2019, 2019, 1-13.	3.0	7
12	Biochemical and genetic approaches to the prenatal diagnosis of propionic acidemia in 78 pregnancies. Orphanet Journal of Rare Diseases, 2020, 15, 276.	2.7	7
13	The Follow-Up of Chinese Patients in cblC Type Methylmalonic Acidemia Identified Through Expanded Newborn Screening. Frontiers in Genetics, 2022, 13, 805599.	2.3	7
14	Noninvasive Prenatal Testing of Methylmalonic Acidemia cblC Type Using the cSMART Assay for MMACHC Gene Mutations. Frontiers in Genetics, 2021, 12, 750719.	2.3	6
15	A rare mutation c.1663Gâ€‰>â€‰A (p.A555T) in the MMUT gene associated with mild clinical and biochemical phenotypes of methylmalonic acidemia in 30 Chinese patients. Orphanet Journal of Rare Diseases, 2021, 16, 22.	2.7	5
16	TWEAK Promotes the Proliferation of Squamous Cell Carcinoma Cells Through Activating cIAP1 Signals. Frontiers in Oncology, 2020, 10, 439.	2.8	3
17	Diagnosis and followâ€up of glycogen storage disease (GSD) type VI from the largest GSD center in China. Human Mutation, 2022, 43, 557-567.	2.5	2