Kristy Iskandar

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4845129/publications.pdf

Version: 2024-02-01

31 papers	524 citations	840776 11 h-index	677142 22 g-index
32 all docs	32 docs citations	32 times ranked	849 citing authors

#	Article	IF	Citations
1	The role of whole exome sequencing in the UBE3A point mutation of Angelman Syndrome: A case report. Annals of Medicine and Surgery, 2022, 73, 103170.	1.1	1
2	Is There Any Mosaicism in REarranged During Transfection Variant in Hirschsprung Diseaseâ€™s Patients?. Frontiers in Pediatrics, 2022, 10, 842820.	1.9	1
3	The impact of NRG1 expressions and methylation on multifactorial Hirschsprung disease. BMC Pediatrics, 2022, 22, 216.	1.7	2
4	Long-term functional outcomes of patients with Hirschsprung disease following pull-through. BMC Pediatrics, 2022, 22, 246.	1.7	4
5	Effect of semaphorin 3C gene variants in multifactorial Hirschsprung disease. Journal of International Medical Research, 2021, 49, 030006052098778.	1.0	3
6	Challenge in diagnosis of late onset necrotizing enterocolitis in a term infant: a case report. BMC Pediatrics, 2021, 21, 152.	1.7	4
7	Molecular epidemiology of SARS-CoV-2 isolated from COVID-19 family clusters. BMC Medical Genomics, 2021, 14, 144.	1.5	13
8	Safety and immunogenicity of human neonatal RV3 rotavirus vaccine (Bio Farma) in adults, children, and neonates in Indonesia: Phase I Trial. Vaccine, 2021, 39, 4651-4658.	3.8	4
9	Congenital Rubella Syndrome Surveillance After Measles Rubella Vaccination Introduction in Yogyakarta, Indonesia. Pediatric Infectious Disease Journal, 2021, 40, 1144-1150.	2.0	2
10	Association between prognostic factors and the outcomes of patients infected with SARS-CoV-2 harboring multiple spike protein mutations. Scientific Reports, 2021, 11, 21352.	3.3	10
11	Diagnostic Value of Dystrophin Immunostaining in the Diagnosis of Duchenne and Becker Muscular Dystrophy Patients. Open Access Macedonian Journal of Medical Sciences, 2021, 9, 1137-1141.	0.2	0
12	Is the Infection of the SARS-CoV-2 Delta Variant Associated With the Outcomes of COVID-19 Patients?. Frontiers in Medicine, 2021, 8, 780611.	2.6	13
13	Aberrant Expressions and Variant Screening of SEMA3D in Indonesian Hirschsprung Patients. Frontiers in Pediatrics, 2020, 8, 60.	1.9	7
14	Full-length genome characterization and phylogenetic analysis of SARS-CoV-2 virus strains from Yogyakarta and Central Java, Indonesia. PeerJ, 2020, 8, e10575.	2.0	17
15	Combined Genetic Effects of RET and NRG1 Susceptibility Variants on Multifactorial Hirschsprung Disease in Indonesia. Journal of Surgical Research, 2019, 233, 96-99.	1.6	16
16	Aberrant expressions of miRNA-206 target, FN1, in multifactorial Hirschsprung disease. Orphanet Journal of Rare Diseases, 2019, 14, 5.	2.7	19
17	The analysis of DMD gene deletions by multiplex PCR in Indonesian DMD/BMD patients: the era of personalized medicine. BMC Research Notes, 2019, 12, 704.	1.4	15
18	Aberrant UBR4 expressions in Hirschsprung disease patients. BMC Pediatrics, 2019, 19, 493.	1.7	8

#	Article	IF	Citations
19	Use of air stacking to improve pulmonary function in Indonesian DuchenneÂmuscular dystrophy patients: bridging the standard of care gap in low middle income country setting. BMC Proceedings, 2019, 13, 21.	1.6	5
20	The impact of down-regulated SK3 expressions on Hirschsprung disease. BMC Medical Genetics, 2018, 19, 24.	2.1	8
21	Clinical profile of congenital rubella syndrome in Yogyakarta, Indonesia. Pediatrics International, 2018, 60, 168-172.	0.5	3
22	NRG1 variant effects in patients with Hirschsprung disease. BMC Pediatrics, 2018, 18, 292.	1.7	12
23	Effect of FTO rs9939609 variant on insulin resistance in obese female adolescents. BMC Research Notes, 2018, 11, 300.	1.4	7
24	The utility of the hematoxylin and eosin staining in patients with suspected Hirschsprung disease. BMC Surgery, 2017, 17, 71.	1.3	18
25	AB053. NRG1 rare variant effects in Hirschsprung disease patients. Annals of Translational Medicine, 2017, 5, AB053-AB053.	1.7	1
26	Accuracy of polymerase chain reaction-restriction fragment length polymorphism for RET rs2435357 genotyping as Hirschsprung risk. Journal of Surgical Research, 2016, 203, 91-94.	1.6	14
27	Loss of Pdk1-Foxo1 Signaling in Myeloid Cells Predisposes to Adipose Tissue Inflammation and Insulin Resistance. Diabetes, 2012, 61, 1935-1948.	0.6	54
28	PDK1-Foxo1 in Agouti-Related Peptide Neurons Regulates Energy Homeostasis by Modulating Food Intake and Energy Expenditure. PLoS ONE, 2011, 6, e18324.	2.5	30
29	PDK-1/FoxO1 pathway in POMC neurons regulatesPomcexpression and food intake. American Journal of Physiology - Endocrinology and Metabolism, 2010, 298, E787-E798.	3.5	59
30	Forkhead Transcription Factor FoxO1 in Adipose Tissue Regulates Energy Storage and Expenditure. Diabetes, 2008, 57, 563-576.	0.6	174
31	Mutation spectrum analysis of DMD gene in Indonesian Duchenne and Becker muscular dystrophy patients. F1000Research, 0, 11, 148.	1.6	0