

Kristy Iskandar

List of Publications by Year in descending order

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Version: 2024-02-01

31
papers

524
citations

840119

11
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676716

22
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32
docs citations

32
times ranked

849
citing authors

#	ARTICLE	IF	CITATIONS
1	Forkhead Transcription Factor FoxO1 in Adipose Tissue Regulates Energy Storage and Expenditure. <i>Diabetes</i> , 2008, 57, 563-576.	0.3	174
2	PDK-1/FoxO1 pathway in POMC neurons regulates <i>Pomc</i> expression and food intake. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , 2010, 298, E787-E798.	1.8	59
3	Loss of Pdk1-Foxo1 Signaling in Myeloid Cells Predisposes to Adipose Tissue Inflammation and Insulin Resistance. <i>Diabetes</i> , 2012, 61, 1935-1948.	0.3	54
4	PDK1-Foxo1 in Agouti-Related Peptide Neurons Regulates Energy Homeostasis by Modulating Food Intake and Energy Expenditure. <i>PLoS ONE</i> , 2011, 6, e18324.	1.1	30
5	Aberrant expressions of miRNA-206 target, FN1, in multifactorial Hirschsprung disease. <i>Orphanet Journal of Rare Diseases</i> , 2019, 14, 5.	1.2	19
6	The utility of the hematoxylin and eosin staining in patients with suspected Hirschsprung disease. <i>BMC Surgery</i> , 2017, 17, 71.	0.6	18
7	Full-length genome characterization and phylogenetic analysis of SARS-CoV-2 virus strains from Yogyakarta and Central Java, Indonesia. <i>PeerJ</i> , 2020, 8, e10575.	0.9	17
8	Combined Genetic Effects of RET and NRG1 Susceptibility Variants on Multifactorial Hirschsprung Disease in Indonesia. <i>Journal of Surgical Research</i> , 2019, 233, 96-99.	0.8	16
9	The analysis of DMD gene deletions by multiplex PCR in Indonesian DMD/BMD patients: the era of personalized medicine. <i>BMC Research Notes</i> , 2019, 12, 704.	0.6	15
10	Accuracy of polymerase chain reaction-restriction fragment length polymorphism for RET rs2435357 genotyping as Hirschsprung risk. <i>Journal of Surgical Research</i> , 2016, 203, 91-94.	0.8	14
11	Molecular epidemiology of SARS-CoV-2 isolated from COVID-19 family clusters. <i>BMC Medical Genomics</i> , 2021, 14, 144.	0.7	13
12	Is the Infection of the SARS-CoV-2 Delta Variant Associated With the Outcomes of COVID-19 Patients?. <i>Frontiers in Medicine</i> , 2021, 8, 780611.	1.2	13
13	NRG1 variant effects in patients with Hirschsprung disease. <i>BMC Pediatrics</i> , 2018, 18, 292.	0.7	12
14	Association between prognostic factors and the outcomes of patients infected with SARS-CoV-2 harboring multiple spike protein mutations. <i>Scientific Reports</i> , 2021, 11, 21352.	1.6	10
15	The impact of down-regulated SK3 expressions on Hirschsprung disease. <i>BMC Medical Genetics</i> , 2018, 19, 24.	2.1	8
16	Aberrant UBR4 expressions in Hirschsprung disease patients. <i>BMC Pediatrics</i> , 2019, 19, 493.	0.7	8
17	Effect of FTO rs9939609 variant on insulin resistance in obese female adolescents. <i>BMC Research Notes</i> , 2018, 11, 300.	0.6	7
18	Aberrant Expressions and Variant Screening of SEMA3D in Indonesian Hirschsprung Patients. <i>Frontiers in Pediatrics</i> , 2020, 8, 60.	0.9	7

#	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.	1.8	5
20	Challenge in diagnosis of late onset necrotizing enterocolitis in a term infant: a case report. BMC Pediatrics, 2021, 21, 152.	0.7	4
21	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.	1.7	4
22	Long-term functional outcomes of patients with Hirschsprung disease following pull-through. BMC Pediatrics, 2022, 22, 246.	0.7	4
23	Clinical profile of congenital rubella syndrome in Yogyakarta, Indonesia. Pediatrics International, 2018, 60, 168-172.	0.2	3
24	Effect of semaphorin 3C gene variants in multifactorial Hirschsprung disease. Journal of International Medical Research, 2021, 49, 030006052098778.	0.4	3
25	Congenital Rubella Syndrome Surveillance After Measles Rubella Vaccination Introduction in Yogyakarta, Indonesia. Pediatric Infectious Disease Journal, 2021, 40, 1144-1150.	1.1	2
26	The impact of NRG1 expressions and methylation on multifactorial Hirschsprung disease. BMC Pediatrics, 2022, 22, 216.	0.7	2
27	AB053. NRG1 rare variant effects in Hirschsprung disease patients. Annals of Translational Medicine, 2017, 5, AB053-AB053.	0.7	1
28	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.	0.5	1
29	Is There Any Mosaicism in REarranged During Transfection Variant in Hirschsprung Diseaseâ€™s Patients?. Frontiers in Pediatrics, 2022, 10, 842820.	0.9	1
30	Mutation spectrum analysis of DMD gene in Indonesian Duchenne and Becker muscular dystrophy patients. F1000Research, 0, 11, 148.	0.8	0
31	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.	0.1	0