Ali Rashidi-Nezhad

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5841703/publications.pdf

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13	186	1163117	1199594
papers	citations	8	12
		h-index	g-index

16	16	16	344
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	CEP104 gene may involve in the pathogenesis of a new developmental disorder other than joubert syndrome. Molecular Biology Reports, 2022, 49, 7231-7237.	2.3	3
2	Evaluation of Gene Expression Alterations of ATM, TP53, and POLM in Gastric Biopsy Specimens of Patients with Gastritis and its Association with Helicobacter Pylori Infection. Majallah-i DÄ•nishgÄ•h-i â€™UlÅ«m-i PizishkÄ«-i Ä³lÄ•m, 2021, 29, 87-95.	0.0	0
3	A novel case report of spinal muscular atrophy with progressive myoclonic epilepsy from Iran . International Medical Case Reports Journal, 2019, Volume 12, 155-159.	0.8	10
4	Utilization of Whole Exome Sequencing in Lethal Form of Multiple Pterygium Syndrome: Identification of Mutations in Embryonal Subunit of Acetylcholine Receptor. International Journal of Molecular and Cellular Medicine, 2019, 8, 258-269.	1.1	2
5	Association of SP-B gene 9306 A/G polymorphism (rs7316) and risk of RDS. Journal of Maternal-Fetal and Neonatal Medicine, 2018, 31, 2965-2970.	1.5	14
6	Effects of cannabinoid receptor type 2 in respiratory syncytial virus infection in human subjects and mice. Virulence, 2018, 9, 217-230.	4.4	54
7	Newborn screening using <scp>TREC</scp>/<scp>KREC</scp> assay for severe T and B cell lymphopenia in Iran. Scandinavian Journal of Immunology, 2018, 88, e12699.	2.7	27
8	Association of SP-C gene codon 186 polymorphism (rs1124) and risk of RDS. Journal of Maternal-Fetal and Neonatal Medicine, 2017, 30, 2585-2589.	1.5	16
9	Partial trisomy 7q and monosomy 13q in a child with disorder of sex development: Phenotypic and genotypic findings. Gene, 2013, 517, 137-145.	2.2	7
10	2q34-qter duplication and 4q34.2-qter deletion in a patient with developmental delay. European Journal of Medical Genetics, 2012, 55, 203-210.	1.3	7
11	Use of siRNA in knocking down of dopamine receptors, a possible therapeutic option in neuropsychiatric disorders. Molecular Biology Reports, 2012, 39, 2003-2010.	2.3	10
12	TGF-Beta codon 25 polymorphism and the risk of graft-versus-host disease after allogenic hematopoietic stem cell transplantation. Iranian Journal of Allergy, Asthma and Immunology, 2010, 9, 1-6.	0.4	9
13	Transforming growth factor-beta1 codon 10 polymorphism is associated with acute GVHD after allogenic BMT in Iranian population. Annals of Transplantation, 2007, 12, 5-10.	0.9	27