Ruslan Bayramov

List of Publications by Year
in descending order

30	202	1162889 8	1125617 13
papers	citations	h-index	g-index

39	39	39	341
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Clinical and molecular evaluation of MEFV gene variants in the Turkish population: a study by the National Genetics Consortium. Functional and Integrative Genomics, 2022, 22, 291-315.	1.4	7
2	Prenatal Sonographic Detection of Monochorionic Twins with Bipartite Placenta. Fetal and Pediatric Pathology, 2021, 40, 685-690.	0.4	1
3	Gene Expression of Mouse Hippocampal Stem Cells Grown in a Galactose-Derived Molecular Gel Compared to In Vivo and Neurospheres. Processes, 2021, 9, 716.	1.3	3
4	Investigation of the relationship between inherited thrombophilia and novel coronavirus pneumonia. Future Virology, 2021, 16, 341-345.	0.9	6
5	Change in gene expression levels of GABA, glutamate and neurosteroid pathways due to acoustic trauma in the cochlea. Journal of Neurogenetics, 2021, 35, 45-57.	0.6	3
6	PEX10-related autosomal recessive cerebellar ataxia with hearing loss. Acta Neurologica Belgica, 2020, 120, 429-432.	0.5	1
7	Disregulation of Autophagy in the Transgenerational Cc2d1a Mouse Model of Autism. NeuroMolecular Medicine, 2020, 22, 239-249.	1.8	14
8	Comparing expression levels of PERIOD genes PER1, PER2 and PER3 in chronic insomnia patients and medical staff working in the night shift. Sleep Medicine, 2020, 73, 101-105.	0.8	7
9	A heritable profile of six miRNAs in autistic patients and mouse models. Scientific Reports, 2020, 10, 9011.	1.6	32
10	A comprehensive molecular analysis and genotypeâ€“phenotype correlation in patients with familial mediterranean fever. Molecular Biology Reports, 2020, 47, 1835-1843.	1.0	11
11	Propranolol decreases DRD3 and SLC1A2 gene expression in patients with essential tremor. Universa Medicina, 2020, 39, 105-112.	0.1	1
12	The association of endothelin-1 levels with renal survival in polycystic kidney disease patients. Journal of Nephrology, 2019, 32, 83-91.	0.9	11
13	A Novel SGCE Nonsense Variant Associated With Marked Intrafamilial Variability in a Turkish Family With Myoclonusâ€Dystonia. Movement Disorders Clinical Practice, 2019, 6, 479-482.	0.8	3
14	Dysmetabolic markers predict outcomes in autosomal dominant polycystic kidney disease. Clinical and Experimental Nephrology, 2019, 23, 1130-1140.	0.7	6
15	Multiple urinary tract infections are associated with genotype and phenotype in adult polycystic kidney disease. Clinical and Experimental Nephrology, 2019, 23, 1188-1195.	0.7	5
16	The molecular basis and genotypeâ€“phenotype correlations of congenital adrenal hyperplasia (CAH) in Anatolian population. Molecular Biology Reports, 2019, 46, 3677-3690.	1.0	9
17	NPHS2 gene sequencing results in children of the Azerbaijani population with different types of nephrotic syndrome caused by chronic glomerulonephritis. Bratislava Medical Journal, 2019, 120, 102-105.	0.4	2
18	Effects of post-learning REM sleep deprivation on hippocampal plasticity-related genes and microRNA in mice. Behavioural Brain Research, 2019, 361, 7-13.	1.2	23

#	Article	IF	Citations
19	A Rare Cause of Male Infertility: 45,X/46,XY Mosaicism. Urologia Internationalis, 2018, 101, 481-485.	0.6	7
20	Effect of the allopregnanolone and allotetrahydrodeoxycorticosteron on spike-wave discharges in the EEG of absence epilepsy rat models. General Physiology and Biophysics, 2018, 37, 205-211.	0.4	5
21	Increased vitamin D receptor gene expression and rs11568820 and rs4516035 promoter polymorphisms in autistic disorder. Molecular Biology Reports, 2018, 45, 541-546.	1.0	16
22	FP049THE ASSOCIATION OF ENDOTHELIN-1 LEVELS WITH RENAL SURVIVAL IN POLYCYSTIC KIDNEY DISEASE PATIENTS. Nephrology Dialysis Transplantation, 2018, 33, i64-i64.	0.4	0
23	Sialidosis type I presenting with a novel mutation and advanced neuroimaging features. Journal of King Abdulaziz University, Islamic Economics, 2018, 23, 57-61.	0.5	9
24	Prenatal diagnosis of a foetus with partial monosomy 4p and partial trisomy 13q. Journal of Biotechnology, 2017, 256, S76.	1.9	0
25	The effect of CYP2C19 * 2 polymorphism on clopidogrel resistance in COPD patients. Journal of Biotechnology, 2017, 256, S80.	1.9	0
26	Frequency of chromosome variants in families with recurrent pregnancy loss and statistical analysis of infertility. Journal of Biotechnology, 2017, 256, S76.	1.9	0
27	A novel nonsense mutation in GALNS gene in family with MPS4A diagnosed child. Journal of Biotechnology, 2016, 231, S108.	1.9	0
28	A case of XYY male patient with micropenis. Journal of Biotechnology, 2016, 231, S109.	1.9	0
29	Effect of sodium benzoate on DNA breakage, micronucleus formation and mitotic index in peripheral blood of pregnant rats and their newborns. Biotechnology and Biotechnological Equipment, 2016, 30, 1179-1183.	0.5	19
30	A case of SRY positive 46, XX male with speaking disorder. Journal of Biotechnology, 2015, 208, S85.	1.9	1