

Essam Al Ageeli

List of Publications by Year in descending order

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Version: 2024-02-01

18
papers

288
citations

1163117

8
h-index

940533

16
g-index

18
all docs

18
docs citations

18
times ranked

531
citing authors

#	ARTICLE	IF	CITATIONS
1	Duplication of the 15q11-q13 region: Clinical and genetic study of 30 new cases. <i>European Journal of Medical Genetics</i> , 2014, 57, 5-14.	1.3	68
2	Oncogenic long noncoding RNA MALAT1 and HCV-related hepatocellular carcinoma. <i>Biomedicine and Pharmacotherapy</i> , 2018, 102, 653-669.	5.6	40
3	A new 17p13.3 microduplication including the PFAFH1B1 and YWHAE genes resulting from an unbalanced X;17 translocation. <i>European Journal of Medical Genetics</i> , 2011, 54, 287-291.	1.3	27
4	Deciphering the role of circulating lncRNAs: RNCR2, NEAT2, CDKN2B-AS1, and PVT1 and the possible prediction of anti-VEGF treatment outcomes in diabetic retinopathy patients. <i>Graefe's Archive for Clinical and Experimental Ophthalmology</i> , 2019, 257, 1897-1913.	1.9	26
5	Long non-coding RNA MALAT1 and microRNA-499a expression profiles in diabetic ESRD patients undergoing dialysis: a preliminary cross-sectional analysis. <i>Archives of Physiology and Biochemistry</i> , 2020, 126, 172-182.	2.1	24
6	Structure and functional impact of seed region variant in MIR-499 gene family in bronchial asthma. <i>Respiratory Research</i> , 2017, 18, 169.	3.6	17
7	Unleash multifunctional role of long noncoding RNAs biomarker panel in breast cancer: a predictor classification model. <i>Epigenomics</i> , 2020, 12, 1215-1237.	2.1	15
8	MYD88, NFKB1, and IL6 transcripts overexpression are associated with poor outcomes and short survival in neonatal sepsis. <i>Scientific Reports</i> , 2021, 11, 13374.	3.3	13
9	Retinal involvement in two unrelated patients with Myhre syndrome. <i>European Journal of Medical Genetics</i> , 2012, 55, 541-547.	1.3	11
10	Alterations of mitochondria and related metabolic pathways in leukemia: A narrative review. <i>Saudi Journal of Medicine and Medical Sciences</i> , 2020, 8, 3.	0.8	10
11	Association of Angio-LncRNAs MIAT rs1061540/MALAT1 rs3200401 Molecular Variants with Gensini Score in Coronary Artery Disease Patients Undergoing Angiography. <i>Biomolecules</i> , 2022, 12, 137.	4.0	9
12	Long Non-Coding RNAs Gene Variants as Molecular Markers for Diabetic Retinopathy Risk and Response to Anti-VEGF Therapy. <i>Pharmacogenomics and Personalized Medicine</i> , 2021, Volume 14, 997-1014.	0.7	8
13	Noncoding RNAs orchestrate cell growth, death and drug resistance in renal cell carcinoma. <i>Epigenomics</i> , 2020, 12, 199-219.	2.1	5
14	Longevity-Related Gene Transcriptomic Signature in Glioblastoma Multiforme. <i>Oxidative Medicine and Cellular Longevity</i> , 2018, 2018, 1-12.	4.0	4
15	Group-specific component exon 11 haplotypes (D432E and T436K) and risk of albuminuria in type 2 diabetes mellitus patients. <i>Archives of Physiology and Biochemistry</i> , 2022, 128, 111-120.	2.1	3
16	MicroRNA-17-92a-1 Host Gene (MIR17HG) Expression Signature and rs4284505 Variant Association with Alopecia Areata: A Case-Control Study. <i>Genes</i> , 2022, 13, 505.	2.4	3
17	Genotype Triad for HOTAIR rs10783618, LINC-ROR rs1942347, and MALAT1 rs3200401 as Molecular Markers in Systemic Lupus Erythematosus. <i>Diagnostics</i> , 2022, 12, 1197.	2.6	3
18	MicroRNA-499a (rs3746444A/G) gene variant and susceptibility to type 2 diabetes-associated end-stage renal disease. <i>Experimental and Therapeutic Medicine</i> , 2021, 23, 63.	1.8	2