

Abdullah Almutery

List of Publications by Year in descending order

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11
papers

139
citations

1684188

5
h-index

1281871

11
g-index

11
all docs

11
docs citations

11
times ranked

175
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic etiology of hereditary hearing loss in the Gulf Cooperation Council countries. <i>Human Genetics</i> , 2022, 141, 595-605.	3.8	5
2	Assessment of Uptake, Accumulation and Degradation of Paracetamol in Spinach (<i>Spinacia oleracea</i> L.) under Controlled Laboratory Conditions. <i>Plants</i> , 2022, 11, 1626.	3.5	8
3	Whole exome sequencing, in silico and functional studies confirm the association of the GJB2 mutation p.Cys169Tyr with deafness and suggest a role for the TMEM59 gene in the hearing process. <i>Saudi Journal of Biological Sciences</i> , 2021, 28, 4421-4429.	3.8	5
4	Genetic Diversity in Casein Gene Cluster in a Dromedary Camel (<i>C. dromedarius</i>) Population from the United Arab Emirates. <i>Genes</i> , 2021, 12, 1417.	2.4	7
5	Genetic diversity, antifungal evaluation and molecular docking studies of Cu-chitosan nanoparticles as prospective stem rust inhibitor candidates among some Egyptian wheat genotypes. <i>PLoS ONE</i> , 2021, 16, e0257959.	2.5	8
6	Mitochondrial mutations in non-syndromic hearing loss at UAE. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2020, 138, 110286.	1.0	4
7	Significant transcriptomic changes are associated with differentiation of bone marrow-derived mesenchymal stem cells into neural progenitor-like cells in the presence of bFGF and EGF. <i>Cell and Bioscience</i> , 2020, 10, 126.	4.8	22
8	Clinical Exome Sequencing Identifies a Frameshift Mutation Within the <i>STRC</i> Gene in a United Arab Emirates Family with Profound Nonsyndromic Hearing Loss. <i>Genetic Testing and Molecular Biomarkers</i> , 2019, 23, 204-208.	0.7	6
9	A Novel Nonsense Mutation (c.414G>A; p.Trp138*) in CLDN14 Causes Hearing Loss in Yemeni Families: A Case Report. <i>Frontiers in Genetics</i> , 2019, 10, 1087.	2.3	3
10	Genistein Induces Alterations of Epigenetic Modulatory Signatures in Human Cervical Cancer Cells. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2018, 18, 412-421.	1.7	51
11	Prevalence of <i>GJB2</i> Mutations in Affected Individuals from United Arab Emirates with Autosomal Recessive Nonsyndromic Hearing Loss. <i>Genetic Testing and Molecular Biomarkers</i> , 2017, 21, 686-691.	0.7	20