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