Nasser A Elhawary

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6767984/publications.pdf

Version: 2024-02-01

20	222	933447 10	1058476 14
papers	citations	h-index	g-index

22	22	22	297
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genetic etiology and clinical challenges of phenylketonuria. Human Genomics, 2022, 16, .	2.9	18
2	Risk of Colorectal Carcinoma May Predispose to the Genetic Variants of the GST, CYP450, and TP53 Genes Among Nonsmokers in the Saudi Community. International Journal of General Medicine, 2021, Volume 14, 1311-1323.	1.8	4
3	Variations in TAP1 and PSMB9 Genes Involved in Antigen Processing and Presentation Increase the Risk of Vitiligo in the Saudi Community. International Journal of General Medicine, 2021, Volume 14, 10031-10044.	1.8	2
4	Genetic biomarkers predict susceptibility to autism spectrum disorder through interactive models of inheritance in a Saudi community. Cogent Biology, 2019, 5, 1606555.	1.7	9
5	Methylenetetrahydrofolate Reductase Gene Variants Confer Potential Vulnerability to Autism Spectrum Disorder in a Saudi Community . Neuropsychiatric Disease and Treatment, 2019, Volume 15, 3569-3581.	2.2	10
6	Molecular characterization of exonic rearrangements and frame shifts in the dystrophin gene in Duchenne muscular dystrophy patients in a Saudi community. Human Genomics, 2018, 12, 18.	2.9	17
7	Combined Genetic Biomarkers Confer Susceptibility to Risk of Urothelial Bladder Carcinoma in a Saudi Population. Disease Markers, 2017, 2017, 1-11.	1.3	13
8	Association betweenANKK1(rs1800497) andLTA(rs909253) Genetic Variants and Risk of Schizophrenia. BioMed Research International, 2015, 2015, 1-8.	1.9	18
9	TransporterTAP1-637G and ImmunoproteasomePSMB9-60H Variants Influence the Risk of Developing Vitiligo in the Saudi Population. Disease Markers, 2014, 2014, 1-8.	1.3	9
10	Azoospermia factor microdeletions: common tag STSs in infertile men with azoospermia and sever oligospermia from Egypt. BMC Genomics, 2014, 15, .	2.8	2
11	TNFâ€238 polymorphism may predict bronchopulmonary dysplasia among preterm infants in the Egyptian population. Pediatric Pulmonology, 2013, 48, 699-706.	2.0	16
12	TheMTHFR677T Allele May Influence the Severity and Biochemical Risk Factors of Alzheimerâ€™s Disease in an Egyptian Population. Disease Markers, 2013, 35, 439-446.	1.3	22
13	Null genetic risk of ACE gene polymorphisms with nephropathy in type 1 diabetes among Egyptian population. Egyptian Journal of Medical Human Genetics, 2011, 12, 187-192.	1.0	2
14	Association between Î²+252 tumour necrosis factor polymorphism and asthma in western Saudi children. Saudi Journal of Biological Sciences, 2011, 18, 107-111.	3.8	4
15	The Impact of Common Tumor Necrosis Factor Haplotypes on the Development of Asthma in Children: An Egyptian Model. Genetic Testing and Molecular Biomarkers, 2011, 15, 293-299.	0.7	16
16	Molecular Updating of Î²-Thalassemia Mutations in the Upper Egyptian Population. Hemoglobin, 2010, 34, 538-547.	0.8	23
17	Common Tag STSs in the AZF Region Associated with Azoospermia and Severe Oligospermia in Infertile Egyptian Men~!2009-12-21~!2010-04-13~!2010-06-18~!. The Open Andrology Journal, 2010, 2, 11-18.	0.2	4
18	High-precision DNA microsatellite genotyping in Duchenne muscular dystrophy families using ion-pair reversed-phase high performance liquid chromatography. Clinical Biochemistry, 2006, 39, 758-761.	1.9	5

#	Article	IF	Citations
19	Mutations in Transglutaminase 1 Gene in Autosomal Recessive Congenital Ichthyosis in Egyptian Families. Disease Markers, 2004, 20, 325-332.	1.3	12
20	Frameshift deletion mechanisms in Egyptian Duchenne and Becker muscular dystrophy families. Molecules and Cells, 2004, 18, 141-9.	2.6	15