

Neda M Bogari

List of Publications by Year in descending order

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Version: 2024-02-01

28
papers

228
citations

1162889

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1058333

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citing authors

#	ARTICLE	IF	CITATIONS
1	Strategies for Vaccination: Conventional Vaccine Approaches Versus New-Generation Strategies in Combination with Adjuvants. <i>Pharmaceutics</i> , 2021, 13, 140.	2.0	28
2	Functional polymorphism in ABCA1 influences age of symptom onset in coronary artery disease patients. <i>Human Molecular Genetics</i> , 2007, 16, 1412-1422.	1.4	25
3	Molecular Updating of β^2 -Thalassemia Mutations in the Upper Egyptian Population. <i>Hemoglobin</i> , 2010, 34, 538-547.	0.4	23
4	The <i>MTHFR</i> 677T Allele May Influence the Severity and Biochemical Risk Factors of Alzheimer's Disease in an Egyptian Population. <i>Disease Markers</i> , 2013, 35, 439-446.	0.6	22
5	Pattern of Thyroid Lesions in Western Region of Saudi Arabia: A Retrospective Analysis and Literature Review. <i>Journal of Clinical Medicine Research</i> , 2018, 10, 106-116.	0.6	22
6	Human genome meeting 2016. <i>Human Genomics</i> , 2016, 10, 12.	1.4	18
7	Genetic construction between polycystic ovarian syndrome and type 2 diabetes. <i>Saudi Journal of Biological Sciences</i> , 2020, 27, 2539-2543.	1.8	13
8	A genetic variant c.553G>T (rs2075291) in the apolipoprotein A5 gene is associated with altered triglycerides levels in coronary artery disease (CAD) patients with lipid lowering drug. <i>BMC Cardiovascular Disorders</i> , 2019, 19, 2.	0.7	11
9	Transporter <i>TAP1</i> -637G and Immunoproteasome <i>PSMB9</i> -60H Variants Influence the Risk of Developing Vitiligo in the Saudi Population. <i>Disease Markers</i> , 2014, 2014, 1-8.	0.6	9
10	Association between HindIII (rs320) variant in the lipoprotein lipase gene and the presence of coronary artery disease and stroke among the Saudi population. <i>Saudi Journal of Biological Sciences</i> , 2020, 27, 2018-2024.	1.8	8
11	No association of apolipoprotein B gene polymorphism and blood lipids in obese Egyptian subjects. <i>Journal of Negative Results in BioMedicine</i> , 2015, 14, 7.	1.4	6
12	Molecular Analysis of Factor VIII and Factor IX Genes in Hemophilia Patients: Identification of Novel Mutations and Molecular Dynamics Studies. <i>Journal of Clinical Medicine Research</i> , 2017, 9, 317-331.	0.6	6
13	Next Generation Exome Sequencing of Pediatric Asthma Identifies Rare and Novel Variants in Candidate Genes. <i>Disease Markers</i> , 2021, 2021, 1-10.	0.6	6
14	Next generation sequencing (NGS) in glucose-6-phosphate dehydrogenase (G6PD) deficiency studies. <i>Bioinformatics</i> , 2016, 12, 41-43.	0.2	6
15	Molecular Dynamics Simulation Reveals Exposed Residues in the Ligand-Binding Domain of the Low-Density Lipoprotein Receptor that Interacts with Vesicular Stomatitis Virus-G Envelope. <i>Viruses</i> , 2019, 11, 1063.	1.5	4
16	The Co-existence of ADHD With Autism in Saudi Children: An Analysis Using Next-Generation DNA Sequencing. <i>Frontiers in Genetics</i> , 2020, 11, 548559.	1.1	4
17	Assessment of genetic polymorphism associated with ATP-binding cassette transporter A1 (ABCA1) gene and fluctuations in serum lipid profile levels in patients with coronary artery disease. <i>Saudi Pharmaceutical Journal</i> , 2021, 29, 1458-1465.	1.2	4
18	Whole exome sequencing detects novel variants in Saudi children diagnosed with eczema. <i>Journal of Infection and Public Health</i> , 2020, 13, 27-33.	1.9	3

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19	Whole Exome Sequencing Reveals Multiple Mutations in Uncommon Genes of Familial Hypercholesterolaemia. <i>Journal of Cardiovascular Disease Research (discontinued)</i> , 2019, 10, 09-15.	0.1	3
20	Null genetic risk of ACE gene polymorphisms with nephropathy in type 1 diabetes among Egyptian population. <i>Egyptian Journal of Medical Human Genetics</i> , 2011, 12, 187-192.	0.5	2
21	Targeted next-generation sequencing reveals novel and known variants of thrombophilia associated genes in Saudi patients with venous thromboembolism. <i>Clinica Chimica Acta</i> , 2021, 519, 247-254.	0.5	2
22	Genetic Association of rs10757278 on Chromosome 9p21 and Coronary Artery Disease in a Saudi Population. <i>International Journal of General Medicine</i> , 2021, Volume 14, 1699-1707.	0.8	1
23	Coronary Artery Disease: Association Study of 5 Loci with Angiographic Indices of Disease Severity. <i>Disease Markers</i> , 2021, 2021, 1-8.	0.6	1
24	Identification of six novel factor viii gene variants using next generation sequencing and molecular dynamics simulation. <i>Acta Biochimica Polonica</i> , 2019, 66, 23-31.	0.3	1
25	A novel SNP in 3' UTR of INS gene: A case report of neonatal diabetes mellitus. <i>Diabetes Research and Clinical Practice</i> , 2015, 109, e14-e17.	1.1	0
26	Compound A Increases Cell Infiltration in Target Organs of Acute Graft-versus-Host Disease (aGVHD) in a Mouse Model. <i>Molecules</i> , 2021, 26, 4237.	1.7	0
27	Prediction of Hepatic Fibrosis in Patients with Chronic Hepatitis C Genotype 4: A Non-Invasive Biochemical Analysis. <i>Biosciences, Biotechnology Research Asia</i> , 2016, 13, 287-297.	0.2	0
28	The Genetic Variant c.553G>T in the Lipoprotein A5 Effects on Lipid Profile Parameters Levels. <i>World Journal of Cardiovascular Diseases</i> , 2019, 09, 122-131.	0.0	0