Neda M Bogari

List of Publications by Year
in descending order

28	228	1162889 8	1058333 14
papers	citations	h-index	g-index

29	29	29	387
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Strategies for Vaccination: Conventional Vaccine Approaches Versus New-Generation Strategies in Combination with Adjuvants. Pharmaceutics, 2021, 13, 140.	2.0	28
2	Next Generation Exome Sequencing of Pediatric Asthma Identifies Rare and Novel Variants in Candidate Genes. Disease Markers, 2021, 2021, 1-10.	0.6	6
3	Genetic Association of rs10757278 on Chromosome 9p21 and Coronary Artery Disease in a Saudi Population. International Journal of General Medicine, 2021, Volume 14, 1699-1707.	0.8	1
4	Compound A Increases Cell Infiltration in Target Organs of Acute Graft-versus-Host Disease (aGVHD) in a Mouse Model. Molecules, 2021, 26, 4237.	1.7	0
5	Coronary Artery Disease: Association Study of 5 Loci with Angiographic Indices of Disease Severity. Disease Markers, 2021, 2021, 1-8.	0.6	1
6	Targeted next-generation sequencing reveals novel and known variants of thrombophilia associated genes in Saudi patients with venous thromboembolism. Clinica Chimica Acta, 2021, 519, 247-254.	0.5	2
7	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. Saudi Pharmaceutical Journal, 2021, 29, 1458-1465.	1.2	4
8	Whole exome sequencing detects novel variants in Saudi children diagnosed with eczema. Journal of Infection and Public Health, 2020, 13, 27-33.	1.9	3
9	The Co-existence of ADHD With Autism in Saudi Children: An Analysis Using Next-Generation DNA Sequencing. Frontiers in Genetics, 2020, 11, 548559.	1.1	4
10	Genetic construction between polycystic ovarian syndrome and type 2 diabetes. Saudi Journal of Biological Sciences, 2020, 27, 2539-2543.	1.8	13
11	Association between HindIII (rs320) variant in the lipoprotein lipase gene and the presence of coronary artery disease and stroke among the Saudi population. Saudi Journal of Biological Sciences, 2020, 27, 2018-2024.	1.8	8
12	A genetic variant c.553Gâ€‰›â€‰T (rs2075291) in the apolipoprotein A5 gene is associated with altered triglycerides levels in coronary artery disease (CAD) patients with lipid lowering drug. BMC Cardiovascular Disorders, 2019, 19, 2.	0.7	11
13	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. Viruses, 2019, 11, 1063.	1.5	4
14	Whole Exome Sequencing Reveals Multiple Mutations in Uncommon Genes of Familial Hypercholesterolaemia. Journal of Cardiovascular Disease Research (discontinued), 2019, 10, 09-15.	0.1	3
15	The Genetic Variant c.553G>T in the Lipoprotein A5 Effects on Lipid Profile Parameters Levels. World Journal of Cardiovascular Diseases, 2019, 09, 122-131.	0.0	0
16	Identification of six novel factor viii gene variants using next generation sequencing and molecular dynamics simulation. Acta Biochimica Polonica, 2019, 66, 23-31.	0.3	1
17	Pattern of Thyroid Lesions in Western Region of Saudi Arabia: A Retrospective Analysis and Literature Review. Journal of Clinical Medicine Research, 2018, 10, 106-116.	0.6	22
18	Molecular Analysis of Factor VIII and Factor IX Genes in Hemophilia Patients: Identification of Novel Mutations and Molecular Dynamics Studies. Journal of Clinical Medicine Research, 2017, 9, 317-331.	0.6	6

#	Article	IF	Citations
19	Human genome meeting 2016. Human Genomics, 2016, 10, 12.	1.4	18
20	Next generation sequencing (NGS) in glucose-6-phosphate dehydrogenase (G6PD) deficiency studies. Bioinformation, 2016, 12, 41-43.	0.2	6
21	Prediction of Hepatic Fibrosis in Patients with Chronic Hepatitis C Genotype 4: A Non-Invasive Biochemical Analysis. Biosciences, Biotechnology Research Asia, 2016, 13, 287-297.	0.2	0
22	No association of apolipoprotein B gene polymorphism and blood lipids in obese Egyptian subjects. Journal of Negative Results in BioMedicine, 2015, 14, 7.	1.4	6
23	A novel SNP in 3â€² UTR of INS gene: A case report of neonatal diabetes mellitus. Diabetes Research and Clinical Practice, 2015, 109, e14-e17.	1.1	0
24	TransporterTAP1-637G and ImmunoproteasomePSMB9-60H Variants Influence the Risk of Developing Vitiligo in the Saudi Population. Disease Markers, 2014, 2014, 1-8.	0.6	9
25	TheMTHFR677T Allele May Influence the Severity and Biochemical Risk Factors of Alzheimerâ€™s Disease in an Egyptian Population. Disease Markers, 2013, 35, 439-446.	0.6	22
26	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.	0.5	2
27	Molecular Updating of Î²-Thalassemia Mutations in the Upper Egyptian Population. Hemoglobin, 2010, 34, 538-547.	0.4	23
28	Functional polymorphism in ABCA1 influences age of symptom onset in coronary artery disease patients. Human Molecular Genetics, 2007, 16, 1412-1422.	1.4	25