Amein K Al-Ali

List of Publications by Year in descending order

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361413 434195 1,174 67 20 31 citations h-index g-index papers 69 69 69 1803 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Sickle cell disease in <scp>S</scp> audi <scp>A</scp> rabia: the phenotype in adults with the <scp>A</scp> rabia: the phenotype in adults with the <scp>A</scp> rabâ€ <scp>I</scp> ndian haplotype is not benign. British Journal of Haematology, 2014, 164, 597-604.	2.5	72
2	Oral and intraperitoneal LD50 of thymoquinone, an active principle of Nigella sativa, in mice and rats. Journal of Ayub Medical College, Abbottabad: JAMC, 2008, 20, 25-7.	0.1	69
3	Influence of vitamin D levels on bone mineral density and osteoporosis. Annals of Saudi Medicine, 2011, 31, 602-608.	1.1	62
4	Fetal hemoglobin in sickle cell anemia: Genetic studies of the Arab-Indian haplotype. Blood Cells, Molecules, and Diseases, 2013, 51, 22-26.	1.4	50
5	Concept and design of a genome-wide association genotyping array tailored for transplantation-specific studies. Genome Medicine, 2015, 7, 90.	8.2	49
6	25-Hydoxyvitamin D levels among healthy Saudi Arabian women. Journal of King Abdulaziz University, Islamic Economics, 2008, 29, 1765-8.	1.1	49
7	A Comprehensive, Ethnically Diverse Library of Sickle Cell Disease-Specific Induced Pluripotent Stem Cells. Stem Cell Reports, 2017, 8, 1076-1085.	4.8	45
8	Lack of MERS Coronavirus Neutralizing Antibodies in Humans, Eastern Province, Saudi Arabia. Emerging Infectious Diseases, 2013, 19, 2034-2036.	4.3	44
9	Comparative and molecular analysis of MRSA isolates from infection sites and carrier colonization sites. Annals of Clinical Microbiology and Antimicrobials, 2018, 17, 7.	3.8	39
10	Vitamin D deficiency and rickets in the Eastern Province of Saudi Arabia. Annals of Tropical Paediatrics, 2007, 27, 63-67.	1.0	36
11	Detection of Pathogenic Variants With Germline Genetic Testing Using Deep Learning vs Standard Methods in Patients With Prostate Cancer and Melanoma. JAMA - Journal of the American Medical Association, 2020, 324, 1957.	7.4	33
12	Cimetidine enhances the hepatoprotective action of N-acetylcysteine in mice treated with toxic doses of paracetamol. Toxicology, 1997, 121, 223-228.	4.2	32
13	Spectrum ofl±-Thalassemia Mutations in Transfusion-Dependent l²-Thalassemia Patients from the Eastern Province of Saudi Arabia. Hemoglobin, 2013, 37, 65-73.	0.8	32
14	Intronic Polymorphisms in the CDKN2B-AS1 Gene Are Strongly Associated with the Risk of Myocardial Infarction and Coronary Artery Disease in the Saudi Population. International Journal of Molecular Sciences, 2016, 17, 395.	4.1	32
15	A phased SNP-based classification of sickle cell anemia HBB haplotypes. BMC Genomics, 2017, 18, 608.	2.8	31
16	lκÎ'α inhibits apoptosis at the outer mitochondrial membrane independently of <scp>NF</scp> â€₽B retention. EMBO Journal, 2014, 33, 2814-2828.	7.8	29
17	Molecular Bases of \hat{I}^2 -Thalassemia in the Eastern Province of Saudi Arabia. Journal of Biomedicine and Biotechnology, 2005, 2005, 322-325.	3.0	28
18	Vitamin D level among patients with sickle cell anemia and its influence on bone mass. American Journal of Hematology, 2011, 86, 506-507.	4.1	26

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19	Molecular Characterization of Glucose-6-Phosphate Dehydrogenase Deficiency in the Eastern Province of Saudi Arabia. Clinical Chemistry and Laboratory Medicine, 2002, 40, 814-6.	2.3	24
20	A novel HBA2 gene conversion in cis or trans: "α12 allele―in a Saudi population. Blood Cells, Molecules, and Diseases, 2014, 53, 199-203.	1.4	23
21	Spectrum of Î ² -Thalassemia Mutations in the Eastern Province of Saudi Arabia. Hemoglobin, 2011, 35, 125-134.	0.8	21
22	Association of <i>MBL2</i> Gene Polymorphism with Dental Caries in Saudi Children. Caries Research, 2017, 51, 12-16.	2.0	20
23	The impact of common polymorphisms in CETP and ABCA1 genes with the risk of coronary artery disease in Saudi Arabians. Human Genomics, 2016, 10, 8.	2.9	18
24	Assessment of low vitamin D among Saudi Arabians. Did we overshoot the runway?. Journal of King Abdulaziz University, Islamic Economics, 2014, 35, 1243-9.	1.1	18
25	Is there a relationship between body mass index and serum vitamin D levels?. Journal of King Abdulaziz University, Islamic Economics, 2009, 30, 1542-6.	1.1	17
26	A candidate transacting modulator of fetal hemoglobin gene expression in the Arabâ€"Indian haplotype of sickle cell anemia. American Journal of Hematology, 2016, 91, 1118-1122.	4.1	16
27	Hemoglobin A2 (HbA2) has a measure of unreliability in diagnosing \hat{I}^2 -thalassemia trait (\hat{I}^2 -TT). Current Medical Research and Opinion, 2018, 34, 945-951.	1.9	16
28	Prevalence and Diversity of Haplotypes of Sickle Cell Disease in the Eastern Province of Saudi Arabia. Hemoglobin, 2020, 44, 78-81.	0.8	16
29	Co-inheritance of novel ATRX gene mutation and globin ($\hat{l}\pm$ & \hat{l}^2) gene mutations in transfusion dependent beta-thalassemia patients. Blood Cells, Molecules, and Diseases, 2015, 55, 27-29.	1.4	15
30	Identification of on-target mutagenesis during correction of a beta-thalassemia splice mutation in iPS cells with optimised CRISPR/Cas9-double nickase reveals potential safety concerns. APL Bioengineering, 2018, 2, 046103.	6.2	14
31	KLF1 gene and borderline hemoglobin A2 in Saudi population. Archives of Medical Science, 2018, 1, 230-236.	0.9	14
32	Pyridine Nucleotide Redox Potential in Erythrocytes of Saudi Subjects with Sickle Cell Disease. Acta Haematologica, 2002, 108, 19-22.	1.4	13
33	Homozygosity for a haplotype in the <i>HBG2â€OR51B4</i> region is exclusive to Arabâ€Indian haplotype sickle cell anemia. American Journal of Hematology, 2016, 91, E308-11.	4.1	13
34	Tyrosine kinase domain mutations of EGFR gene in head and neck squamous cell carcinoma. OncoTargets and Therapy, 2017, Volume 10, 1527-1533.	2.0	13
35	Bedside testing of CYP2C19 vs. conventional clopidogrel treatment to guide antiplatelet therapy in ST-segment elevation myocardial infarction patients. International Journal of Cardiology, 2021, 343, 15-20.	1.7	12
36	A functional promoter polymorphism of the \hat{l} -globin gene is a specific marker of the Arab-Indian haplotype. American Journal of Hematology, 2012, 87, 824-826.	4.1	11

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37	Variants of ZBTB7A (LRF) and its \hat{l}^2 -globin gene cluster binding motifs in sickle cell anemia. Blood Cells, Molecules, and Diseases, 2016, 59, 49-51.	1.4	11
38	Type 2 diabetes associated variants of KCNQ1 strongly confer the risk of cardiovascular disease among the Saudi Arabian population. Genetics and Molecular Biology, 2017, 40, 586-590.	1.3	11
39	Genetic studies of fetal hemoglobin in the Arabâ€Indian haplotype sickle cellâ€Î² ⁰ thalassemia. American Journal of Hematology, 2013, 88, 531-532.	4.1	8
40	Adeno-Associated Viral Transfer of Glyoxalase-1 Blunts Carbonyl and Oxidative Stresses in Hearts of Type 1 Diabetic Rats. Antioxidants, 2020, 9, 592.	5.1	8
41	TNFSF/TNFRSF cytokine gene expression in sickle cell anemia: Up-regulated TNF-like cytokine 1A (TL1A) and its decoy receptor (DcR3) in peripheral blood mononuclear cells and plasma. Cytokine, 2019, 123, 154744.	3.2	7
42	Exome-Wide Association Analysis of Coronary Artery Disease in the Kingdom of Saudi Arabia Population. PLoS ONE, 2016, 11, e0146502.	2.5	7
43	The â€Î±3.7 deletion in αâ€'globin genes increases the concentration of fetal hemoglobin and hemoglobin A2 in a Saudi Arabian population. Molecular Medicine Reports, 2018, 17, 1879-1884.	2.4	6
44	Exome sequencing of Saudi Arabian patients with ADPKD. Renal Failure, 2019, 41, 842-849.	2.1	6
45	Sickle cell disease in the Eastern Province of Saudi Arabia: Clinical and laboratory features. American Journal of Hematology, 2021, 96, E117-E121.	4.1	6
46	Early detection of SARSâ€CoVâ€⊋ and other infections in solid organ transplant recipients and household members using wearable devices. Transplant International, 2021, 34, 1019-1031.	1.6	6
47	Vitamin D Levels in Healthy Men in Eastern Saudi Arabia. Annals of Saudi Medicine, 2009, 29, 378-382.	1.1	6
48	Frequency of methylenetetrahydrofolate reductase C677T polymorphism in patients with cardiovascular disease in Eastern Saudi Arabia. Journal of King Abdulaziz University, Islamic Economics, 2005, 26, 1886-8.	1.1	6
49	Bedside testing of CYP2C19 gene for treatment of patients with PCI with antiplatelet therapy. BMC Cardiovascular Disorders, 2020, 20, 268.	1.7	5
50	Evaluating the molecular diagnostic yield of joint genotyping–based approach for detecting rare germline pathogenic and putative loss-of-function variants. Genetics in Medicine, 2021, 23, 918-926.	2.4	5
51	Association between hepatitis C virus viremia and the rs12979860, rs2228145 and rs1800795 SNP (CT/AC/GG) genotype in Saudi kidney transplant recipients. Saudi Journal of Medicine and Medical Sciences, 2020, 8, 46.	0.8	3
52	Investigation of KIF6Trp719Arg gene polymorphism in a case-control study of coronary artery disease and non-fatal myocardial infarction in the Eastern Province of Saudi Arabia. Annals of Saudi Medicine, 2016, 36, 105-111.	1.1	3
53	Polymorphism in methylenetetrahydrofolate reductase, plasminogen activator inhibitor-1, and apolipoprotein E in hemodialysis patients. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2008, 19, 937-41.	0.3	3
54	Exome sequencing in high and low fetal haemoglobin Arab–Indian haplotype sickle cell disease. British Journal of Haematology, 2021, 194, e61-e64.	2.5	2

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55	\hat{l}^2 ₂ -adrenergic receptor gene polymorphisms in normal and in patients with myocardial infarction in the eastern province of Saudi Arabia. Saudi Journal of Medicine and Medical Sciences, 2013, 1, 25.	0.8	2
56	Perspective: A novel prognostic for sickle cell disease. Saudi Journal of Medicine and Medical Sciences, 2018, 6, 133.	0.8	2
57	Prevalence of <i>CYP2C19*2</i> carriers in Saudi ischemic stroke patients and the suitability of using genotyping to guide antiplatelet therapy in a university hospital setup. Drug Metabolism and Personalized Therapy, 2022, 37, 35-40.	0.6	2
58	Prevalence of CYP2C19*2 carriers in Saudi ischemic stroke patients and the suitability of using genotyping to guide antiplatelet therapy in a university hospital setup. Drug Metabolism and Personalized Therapy, 2021, .	0.6	1
59	Co-Inheritance of Delta Thalassemia Might Contribute to the High Fetal Hemoglobin in Sickle Cell Anemia Patients with the Saudi-Indian Haplotype. Blood, 2011, 118, 1056-1056.	1.4	1
60	Epsteinâ€'Barr virus infection mediated TP53 and Bclâ€'2Âexpression in nasopharyngeal carcinoma pathogenesis. Molecular and Clinical Oncology, 2021, 15, 260.	1.0	1
61	A concise history of genome-wide association studies. Saudi Journal of Medicine and Medical Sciences, 2013, 1, 4.	0.8	1
62	The Evolutionary Impact Of Malaria On The Saudi Arabian Genome. Blood, 2013, 122, 1001-1001.	1.4	1
63	Increased Prevalence of Glycoprotein IIb/IIIa Leu 33 Pro Polymorphism in End Stage Renal Disease Patients on Hemodialysis. International Journal of Biomedical Science, 2008, 4, 175-8.	0.1	1
64	A Library of Sickle Cell Anemia Induced Pluripotent Stem Cells of Diverse Haplotypes and Ethnicities. Blood, 2015, 126, 2354-2354.	1.4	0
65	Polymorphisms Associated with the Arab-Indian Haplotype of Sickle Cell Anemia Are Candidate Fetal Hemoglobin Gene Modulators. Blood, 2015, 126, 3388-3388.	1.4	0
66	Association of epidermal growth factor receptor protein expression with histopathological and clinical parameters in carcinoma of the larynx. Translational Cancer Research, 2019, 8, 1395-1402.	1.0	0
67	Reply to the Letter to the Editor: "lt is urgent to evaluate the efficacy and safety of genotype guided antiplatelet therapy in patients after percutaneous coronary intervention in East Asian― International Journal of Cardiology, 2022, 348, 57.	1.7	0