Zohreh Rahimi

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1239574/publications.pdf

Version: 2024-02-01

163 papers	3,006 citations	185998 28 h-index	264894 42 g-index
171 all docs	171 docs citations	171 times ranked	4193 citing authors

#	Article	IF	Citations
1	COVID-19 and psoriasis: biologic treatment and challenges. Journal of Dermatological Treatment, 2022, 33, 699-703.	1.1	30
2	Oxidative stress parameters and keap 1 variants in T2DM: Association with T2DM, diabetic neuropathy, diabetic retinopathy, and obesity. Journal of Clinical Laboratory Analysis, 2022, 36, e24163.	0.9	11
3	Demographics, clinical characteristics, and outcomes of 27,256 hospitalized COVID-19 patients in Kermanshah Province, Iran: a retrospective one-year cohort study. BMC Infectious Diseases, 2022, 22, 319.	1.3	19
4	Association of Keap1 (rs11085735) polymorphism and lncRNA MEG3 hypermethylation status with the risk of preeclampsia. Egyptian Journal of Medical Human Genetics, 2022, 23, .	0.5	0
5	New inflammatory biomarkers (lymphocyte and monocyte percentage to high-density lipoprotein) Tj ETQq1 1 0.784314 rgBT /Overlock cardiometabolic diseases. Wiener Klinische Wochenschrift, 2022, 134, 626-635.	1.0	6
6	Gene variants and haplotypes of Vitamin D biosynthesis, transport, and function in preeclampsia. Hypertension in Pregnancy, 2021, 40, 1-8.	0.5	8
7	Establishing hematological reference intervals in healthy adults: Ravansar nonâ€communicable disease cohort study, Iran. International Journal of Laboratory Hematology, 2021, 43, 199-209.	0.7	3
8	Variants of Genes Involved in Metabolism of Folate among Patients with Breast Cancer: Association of TYMS 3R Allele with Susceptibility to Breast Cancer and Metastasis. Iranian Journal of Pathology, 2021, 16, 62-68.	0.2	5
9	Chitosan/tripolyphosphate nanoparticles in active and passive microchannels. Research in Pharmaceutical Sciences, 2021, 16, 79.	0.6	4
10	Efficacy and safety of sofosbuvir/velpatasvir versus the standard of care in adults hospitalized with COVID-19: a single-centre, randomized controlled trial. Journal of Antimicrobial Chemotherapy, 2021, 76, 2158-2167.	1.3	15
11	Co-encapsulation of tertinoin and resveratrol by solid lipid nanocarrier (SLN) improves mice inÂvitro matured oocyte/ morula-compact stage embryo development. Theriogenology, 2021, 171, 1-13.	0.9	4
12	The role of caveolinâ€1 and endothelial nitric oxide synthase polymorphisms in susceptibility to prostate cancer. International Journal of Experimental Pathology, 2021, 102, 260-267.	0.6	6
13	A systematic review and meta-analysis of the DNA methylation in colorectal cancer among Iranian population. Gene Reports, 2021, 23, 101080.	0.4	0
14	Sickle cell disease and COVIDâ€19: Susceptibility and severity. Pediatric Blood and Cancer, 2021, 68, e29075.	0.8	25
15	Liver Enzymes and Their Association with Some Cardiometabolic Diseases: Evidence from a Large Kurdish Cohort. BioMed Research International, 2021, 2021, 1-8.	0.9	14
16	Vitamin D level, lipid profile, and vitamin D receptor and transporter gene variants in sickle cell disease patients from Kurdistan of Iraq. Journal of Clinical Laboratory Analysis, 2021, 35, e23908.	0.9	4
17	Association between RBC Indices, Anemia, and Obesity-Related Diseases Affected by Body Mass Index in Iranian Kurdish Population: Results from a Cohort Study in Western Iran. International Journal of Endocrinology, 2021, 2021, 1-13.	0.6	5
18	Aberrant expression profile of miR-32, miR-98 and miR-374 in chronic lymphocytic leukemia. Leukemia Research, 2021, 111, 106691.	0.4	7

#	Article	IF	Citations
19	Evaluation of The Relationship among The Levels of SIRT1 and SIRT3 with Oxidative Stress and DNA Fragmentation in Asthenoteratozoospermic Men. International Journal of Fertility & Sterility, 2021, 15, 135-140.	0.2	5
20	Modulation of Fibroblast Growth Factor-21 and Î²klotho Proteins Expression in Type 2 Diabetic Women with Non-alcoholic Fatty Liver Disease Following Endurance and Strength Training. Hepatitis Monthly, 2021, 21, .	0.1	5
21	Analysis of glucocerebrosidase (GBA) gene mutations in Iranian patients with Gaucher disease. Iranian Journal of Child Neurology, 2021, 15, 139-166.	0.2	0
22	<scp>COVID</scp>â€19 and renin angiotensin aldosterone system: Pathogenesis and therapy. Health Science Reports, 2021, 4, e440.	0.6	13
23	Thalassemia and COVID-19: Susceptibility and Severity. Iranian Journal of Pediatrics, 2021, 31, .	0.1	3
24	Oxidative Stress Parameters, Trace Elements, and Lipid Profile in Iranian Patients with Gaucher Disease. Biological Trace Element Research, 2020, 193, 130-137.	1.9	3
25	CYP24A1 genetic variants in the vitamin D metabolic pathway are involved in the outcomes of hepatitis C virus infection among high-risk Chinese population. International Journal of Infectious Diseases, 2020, 91, 270.	1.5	0
26	Leukocytosis and alteration of hemoglobin level in patients with severe <scp>COVID</scp>â€19: Association of leukocytosis with mortality. Health Science Reports, 2020, 3, e194.	0.6	12
27	The clinical significance of circulating DSCAM-AS1 in patients with ER-positive breast cancer and construction of its competitive endogenous RNA network. Molecular Biology Reports, 2020, 47, 7685-7697.	1.0	5
28	A caseâ€control study on the SNP309T â†' G and 40â€bp Del1518 of the MDM2 gene and a systematic review for MDM2 polymorphisms in the patients with breast cancer. Journal of Clinical Laboratory Analysis, 2020, 34, e23529.	0.9	2
29	Activities and polymorphisms of MMP-2 and MMP-9, smoking, diabetes and risk of prostate cancer. Molecular Biology Reports, 2020, 47, 9373-9383.	1.0	16
30	Blood coagulation parameters in patients with severe COVID-19 from Kermanshah Province, Islamic Republic of Iran. Eastern Mediterranean Health Journal, 2020, 26, 999-1004.	0.3	17
31	The Insulin-like Growth Factor-1 (G>A) and 5,10-methylenetetrahydrofolate Reductase (C677T) Gene Variants and the Serum Levels of Insulin-like Growth Factor-1, Insulin, and Homeostasis Model Assessment in Patients with Acne Vulgaris. Iranian Journal of Pathology, 2020, 15, 23-29.	0.2	4
32	Comment on â€œAssociation between Interleukin-32 and Interleukin-17A Single Nucleotide Polymorphisms and Serum Levels with Polycystic Ovary Syndromeâ€. Iranian Journal of Allergy, Asthma and Immunology, 2020, 19, 318-319.	0.3	0
33	Hemoglobinopathies in Iran: An Updated Review. International Journal of Hematology-Oncology and Stem Cell Research, 2020, 14, 140-150.	0.3	2
34	Circulating CYTOR as a Potential Biomarker in Breast Cancer. International Journal of Molecular and Cellular Medicine, 2020, 9, 83-90.	1.1	6
35	Antioxidant activities of Î±-lipoic acid free and nano-capsule inhibit the growth of Ehrlich carcinoma. Molecular Biology Reports, 2019, 46, 6685-6686.	1.0	0
36	Vitamin D-binding protein and vitamin D receptor genotypes and 25-hydroxyvitamin D levels are associated with development of aortic and mitral valve calcification and coronary artery diseases. Molecular Biology Reports, 2019, 46, 5225-5236.	1.0	13

#	Article	IF	Citations
37	Mapping 123 million neonatal, infant and child deaths between 2000 and 2017. Nature, 2019, 574, 353-358.	13.7	161
38	Promoter Methylation Status of the Retinoic Acid Receptor-Beta 2 Gene in Breast Cancer Patients: A Case Control Study and Systematic Review. Breast Care, 2019, 14, 117-123.	0.8	9
39	New insight into the role of long non-coding RNAs in the pathogenesis of preeclampsia. Hypertension in Pregnancy, 2019, 38, 41-51.	0.5	23
40	NOTCH1, SF3B1, MDM2 and MYD88 mutations in patients with chronic lymphocytic leukemia. Oncology Letters, 2019, 17, 4016-4023.	0.8	11
41	Interaction of long noncoding RNA MEG3 with miRNAs: A reciprocal regulation. Journal of Cellular Biochemistry, 2019, 120, 3339-3352.	1.2	35
42	Association between the â˜11377 C/G and â˜11391 G/A polymorphisms of adiponectin gene and adiponectin levels with susceptibility to type 1 and type 2 diabetes mellitus in population from the west of Iran, correlation with lipid profile. Journal of Cellular Biochemistry, 2019, 120, 3574-3582.	1.2	8
43	Association between activity and genotypes of paraoxonase1 L55M (rs854560) increases the disease activity of rheumatoid arthritis through oxidative stress. Molecular Biology Reports, 2019, 46, 741-749.	1.0	8
44	The effect of VDR gene polymorphisms and vitamin D level on blood pressure, risk of preeclampsia, gestational age, and body mass index. Journal of Cellular Biochemistry, 2019, 120, 6441-6448.	1.2	30
45	Angiotensinâ€converting enzyme insertion/deletion (rs106180) and angiotensin type 1 receptor A 1166 C (rs106165) genotypes and psoriasis: Correlation with cellular immunity, lipid profile, and oxidative stress markers. Journal of Cellular Biochemistry, 2019, 120, 2627-2633.	1.2	6
46	MTHFR C677T Polymorphism Is Associated with the Risk of Breast Cancer among Kurdish Population from Western Iran. International Journal of Cancer Management, 2019, In Press, .	0.2	5
47	The CYP17 MSP AI (T-34C) and CYP19A1 (Trp39Arg) variants in polycystic ovary syndrome: A case-control study. International Journal of Reproductive BioMedicine, 2019, 17, .	0.5	8
48	The Prevalence of Hemoglobinopathies in Reference Laboratory of Kermanshah, Western Iran. Iranian Journal of Public Health, 2019, 48, 359-361.	0.3	1
49	Association between CYP19A<G rs700518 Polymorphism with Acne Vulgaris and its Severity: Influence on Sex Hormones Level. International Journal of Molecular and Cellular Medicine, 2019, 8, 162-168.	1.1	5
50	Chemerin rs17173608 and vaspin rs2236242 gene variants on the risk of end stage renal disease (ESRD) and correlation with plasma malondialdehyde (MDA) level. Renal Failure, 2018, 40, 350-356.	0.8	10
51	Synergism between apolipoprotein E Æ4 allele and paraoxonase (PON1) 55-M allele is associated with risk of systemic lupus erythematosus. Clinical Rheumatology, 2018, 37, 971-977.	1.0	11
52	Association of the CYP17 MSP AI (T-34C) and CYP19 codon 39 (Trp/Arg) polymorphisms with susceptibility to acne vulgaris. Clinical and Experimental Dermatology, 2018, 43, 183-186.	0.6	13
53	MMP-8 C-799T and MMP-8 C+17G polymorphisms in mild and severe preeclampsia: Association between MMP-8 C-799T with susceptibility to severe preeclampsia. Clinical and Experimental Hypertension, 2018, 40, 175-178.	0.5	6
54	Matrix metalloproteinase-2 C-735T and its interaction with matrix metalloproteinase-7 A-181G polymorphism are associated with the risk of preeclampsia: influence on total antioxidant capacity and blood pressure. Journal of Obstetrics and Gynaecology, 2018, 38, 327-332.	0.4	10

#	Article	IF	Citations
55	PPARÎ³ Pro12Ala and C161T polymorphisms in patients with acne vulgaris: Contribution to lipid and lipoprotein profile. Advances in Medical Sciences, 2018, 63, 147-151.	0.9	11
56	Sleep Architecture and Hypothalamic-Pituitary-Adrenal Activity in Paradoxical and Psychophysiological Insomnia. Basic and Clinical Neuroscience, 2018, 9, 397-407.	0.3	8
57	Relationship between serum homovanillic acid, DRD2 C957T (rs6277), and hDAT A559V (rs28364997) polymorphisms and developmental stuttering. Journal of Communication Disorders, 2018, 76, 37-46.	0.8	8
58	The Relationship Between Polymorphisms of Glutathione S-Transferase (GSTM1, GSTT1) Gene and Type 2 Diabetes Mellitus in Western Iran?. Journal of Kermanshah University of Medical Sciences, 2018, In Press, .	0.1	0
59	The Association of PPARÎ³ Pro12Ala and C161T Polymorphisms with Polycystic Ovary Syndrome and Their Influence on Lipid and Lipoprotein Profiles. International Journal of Fertility & Sterility, 2018, 12, 147-151.	0.2	4
60	Effects of Resveratrol on and a Genes Expression in Adipose Tissue, Serum Insulin, Insulin Resistance and Serum SOD Activity in Type 2 Diabetic Rats. International Journal of Molecular and Cellular Medicine, 2018, 7, 176-184.	1.1	5
61	Cytochrome P450 (CYP450,2D6*A), N-Acetyltransferase-2 (NAT2*7, A) and Multidrug Resistance 1 (MDR1) Tj ETQq1 1 0.784314 rgBT /O 530-535.	0.2	3
62	Angiotensin type 1 receptor A1166C polymorphism and systemic lupus erythematosus: correlation with cellular immunity and oxidative stress markers. Lupus, 2017, 26, 1534-1539.	0.8	8
63	Sex steroid hormones and sex hormone binding globulin levels, CYP17 MSP AI (â^'34 T:C) and CYP19 codon 39 (Trp:Arg) variants in children with developmental stuttering. Brain and Language, 2017, 175, 47-56.	0.8	12
64	The GPX1 Pro198Leu polymorphism in gastric cancer patients with and without Helicobacter pylori infection. Genes and Genomics, 2017, 39, 1265-1269.	0.5	2
65	Modulation of oxidative and glycolytic skeletal muscle fibers Na+/H+ exchanger1 (NHE1) and Na+/HCO3- co-transporter1 (NBC1) genes and proteins expression in type 2 diabetic rat (Streptozotocin) Tj ETQq1 1 0.784314 rgBT /O 11-18.	0.3	9
66	Cytotoxic T-lymphocyte Associated Antigen-4 (CTLA-4) Polymorphism, Cancer, and Autoimmune Diseases. AIMS Medical Science, 2017, 4, 395-412.	0.2	6
67	Genetic Variants of Pre-microRNAs A-499G(rs3746444) and T-196a2C(rs11614913) with Ulcerative Colitis (UC) and Investigated with Thiopurine-S-Methyltransferase (TPMT) Activity. Clinical Laboratory, 2017, 63, 1683-1690.	0.2	3
68	Methylenetetrahydrofolate Reductase (MTHFR) C677T and A1298C Variants, Folate Intake, and Susceptibility to Breast Cancer. International Journal of Cancer Management, 2017, 10, .	0.2	1
69	I/D and A-181G variants and the risk of end stage renal disease. Molecular Biology Research Communications, 2017, 6, 41-44.	0.2	2
70	Association between GSTM1, GSTT1, and GSTP1 variants and the risk of end stage renal disease. Renal Failure, 2016, 38, 1455-1461.	0.8	29
71	The T Allele of MTHFR c.C677T and Its Synergism with G (Val 158) Allele of COMT c.G472A Polymorphism Are Associated with the Risk of Bipolar I Disorder. Genetic Testing and Molecular Biomarkers, 2016, 20, 510-515.	0.3	3
72	Functional Promoter Polymorphisms of MMP-2 C-735T and MMP-9 C-1562T and Their Synergism with MMP-7 A-181G in Multiple Sclerosis. Immunological Investigations, 2016, 45, 543-552.	1.0	19

#	Article	IF	Citations
73	Association of the hypermethylation status of PTEN tumor suppressor gene with the risk of breast cancer among Kurdish population from Western Iran. Tumor Biology, 2016, 37, 8145-8152.	0.8	29
74	The Role of Renin Angiotensin Aldosterone System Genes in Diabetic Nephropathy. Canadian Journal of Diabetes, 2016, 40, 178-183.	0.4	44
75	Brain-Derived Neurotrophic Factor Val66Met Polymorphism and Its Synergism with L/S Polymorphism in the Promoter Region of Serotonin Transporter in Bipolar I Disorder Patients in Western Iran. Iranian Journal of Psychiatry and Behavioral Sciences, 2016, In Press, .	0.1	0
76	Effect of Acetylcholinesterase and Butyrylcholinesterase on Intrauterine Insemination, Contribution to Inflammations, Oxidative Stress and Antioxidant Status; A Preliminary Report. Journal of Reproduction and Infertility, 2016, 17, 157-62.	1.0	1
77	Chitotriosidase Activity and Gene Polymorphism in Iranian Patients with Gaucher Disease and Sibling Carriers. Iranian Journal of Child Neurology, 2016, 10, 62-70.	0.2	3
78	Association between the cytotoxic T-lymphocyte antigen-4 mutations and the susceptibility to systemic lupus erythematosus; Contribution markers of inflammation and oxidative stress. Cellular and Molecular Biology, 2016, 62, 56-61.	0.3	11
79	Allele specific-PCR and melting curve analysis showed relatively high frequency of Î²-casein gene A1 allele in Iranian Holstein, Simmental and native cows. Cellular and Molecular Biology, 2016, 62, 138-143.	0.3	4
80	MMP-7 A-181G Polymorphism in Breast Cancer Patients from Western Iran. Breast Care, 2015, 10, 398-402.	0.8	12
81	The serotonin transporter (5-HTTLPR) but not serotonin receptor (5-HT2C Cys23Ser) variant is associated with bipolar I disorder in Kurdish population from Western Iran. Neuroscience Letters, 2015, 590, 91-95.	1.0	7
82	Matrix metalloproteinase 9 polymorphisms and systemic lupus erythematosus: correlation with systemic inflammatory markers and oxidative stress. Lupus, 2015, 24, 597-605.	0.8	38
83	Association between butyrylcholinesterase activity and phenotypes, paraoxonase192 rs662 gene polymorphism and their enzymatic activity with severity of rheumatoid arthritis: Correlation with systemic inflammatory markers and oxidative stress, preliminary report. Clinical Biochemistry, 2015, 48, 63-69.	0.8	35
84	Matrix metalloproteinase-7 A-181G and its interaction with matrix metalloproteinase-9 C-1562T polymorphism in preeclamptic patients: association with malondialdehyde level and severe preeclampsia. Archives of Gynecology and Obstetrics, 2015, 291, 45-51.	0.8	11
85	Angiotensin converting enzyme insertion/deletion (I/D) (rs4646994) and Vegf polymorphism (+405G/C;) Tj ETQq1 1 0.784314 rgBT /Ov Journal of the Renin-Angiotensin-Aldosterone System, 2015, 16, 672-680.	1.0	41
86	AT1R A1166C variants in patients with type 2 diabetes mellitus and diabetic nephropathy. Journal of Nephropathology, 2015, 4, 69-76.	0.1	18
87	The Association Between Matrix Metalloproteinase-7 A-181G Polymorphism and the Risk of Relapsing-Remitting Multiple Sclerosis in Iranian Kurdish Patients from Kermanshah. Avicenna Journal of Medical Biochemistry, 2015, 3, .	0.5	1
88	Cancer Notification at a Referral Hospital of Kermanshah, Western Iran (2006-2009). Asian Pacific Journal of Cancer Prevention, 2015, 16, 133-137.	0.5	7
89	Manganese Superoxide Dismutase (MnSOD Val-9Ala) Gene Polymorphism and Susceptibility to Gastric Cancer. Asian Pacific Journal of Cancer Prevention, 2015, 16, 485-488.	0.5	13
90	Matrix Metalloproteinase-9 -1562T Allele and its Combination with MMP-2 -735 C Allele are Risk Factors for Breast Cancer. Asian Pacific Journal of Cancer Prevention, 2015, 16, 1175-1179.	0.5	26

#	Article	IF	Citations
91	Methylentetrahydrofolatereductase (rs1801133) polymorphism and psoriasis: contribution to oxidative stress, lipid peroxidation and correlation with vascular adhesion protein 1, preliminary report. Journal of the European Academy of Dermatology and Venereology, 2014, 28, 1192-1198.	1.3	27
92	Association of matrix metalloproteinase-7A-181G variants with the risk of multiple sclerosis. Personalized Medicine, 2014, 11, 727-733.	0.8	1
93	Synergistic effects of angiotensinogen â^'217 Gâ†'A and T704C (M235T) variants on the risk of severe preeclampsia. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2014, 15, 156-161.	1.0	11
94	Synergistic effects of BuChE non-UU phenotype and paraoxonase (PON1) 55â€‰M allele on the risk of systemic lupus erythematosus: influence on lipid and lipoprotein metabolism and oxidative stress, preliminary report. Lupus, 2014, 23, 263-272.	0.8	22
95	The green synthesis, characterization and antimicrobial activities of silver nanoparticles synthesized from green alga Enteromorpha flexuosa (wulfen) J. Agardh. Materials Letters, 2014, 137, 1-4.	1.3	113
96	Hemospermia: long-term outcome in 165 patients. International Journal of Impotence Research, 2014, 26, 83-86.	1.0	22
97	AT2R â^'1332 G:A polymorphism and its interaction with AT1R 1166 A:C, ACE I/D and MMP-9 â^'1562 C:T polymorphisms: Risk factors for susceptibility to preeclampsia. Gene, 2014, 538, 176-181.	1.0	44
98	The MMP-2 -735 C Allele is a Risk Factor for Susceptibility to Breast Cancer. Asian Pacific Journal of Cancer Prevention, 2014, 15, 6199-6203.	0.5	28
99	The prevalence of anemia and hemoglobinopathies in the hematologic clinics of the kermanshah province, Western iran. International Journal of Hematology-Oncology and Stem Cell Research, 2014, 8, 33-7.	0.3	4
100	A systematic review of the role of renin angiotensin aldosterone system genes in diabetes mellitus, diabetic retinopathy and diabetic neuropathy. Journal of Research in Medical Sciences, 2014, 19, 1090-8.	0.4	37
101	Butyrylcholinesterase (BChE) activity is associated with the risk of preeclampsia: influence on lipid and lipoprotein metabolism and oxidative stress. Journal of Maternal-Fetal and Neonatal Medicine, 2013, 26, 1590-1594.	0.7	16
102	MTHFR C677T and eNOS G894T variants in preeclamptic women: Contribution to lipid peroxidation and oxidative stress. Clinical Biochemistry, 2013, 46, 143-147.	0.8	28
103	Genetic Epidemiology, Hematological and Clinical Features of Hemoglobinopathies in Iran. BioMed Research International, 2013, 2013, 1-10.	0.9	34
104	Mental health problems of young refugees: Duration of settlement, risk factors and community-based interventions. Clinical Child Psychology and Psychiatry, 2013, 18, 604-623.	0.8	131
105	MMP-9 (-1562 C:T) polymorphism as a biomarker of susceptibility to severe pre-eclampsia. Biomarkers in Medicine, 2013, 7, 93-98.	0.6	32
106	Preeclampsia and angiotensin converting enzyme (ACE) I/D and angiotensin II type-1 receptor (AT1R) A1166C polymorphisms: association with ACE I/D polymorphism. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2013, 14, 174-180.	1.0	35
107	eNOS 4a/b Polymorphism and Its Interaction with eNOS G894T Variants in Type 2 Diabetes Mellitus: Modifying the Risk of Diabetic Nephropathy. Disease Markers, 2013, 34, 437-443.	0.6	18
108	AT2R -1332 G:A polymorphism and diabetic nephropathy in type 2 diabetes mellitus patients. Journal of Renal Injury Prevention, 2013, 2, 97-101.	0.6	18

#	Article	IF	Citations
109	eNOS 4a/b polymorphism and its interaction with eNOS G894T variants in type 2 diabetes mellitus: modifying the risk of diabetic nephropathy. Disease Markers, 2013, 34, 437-43.	0.6	15
110	Endothelial Nitric Oxide Synthase (eNOS) 4a/b and G894T Polymorphisms and Susceptibility to Preeclampsia. Journal of Reproduction and Infertility, 2013, 14, 184-9.	1.0	12
111	Paraoxonase (PON1) 55 polymorphism and association with systemic lupus erythematosus. Iranian Journal of Allergy, Asthma and Immunology, 2013, 12, 211-9.	0.3	10
112	NPHP4 Variants Are Associated With Pleiotropic Heart Malformations. Circulation Research, 2012, 110, 1564-1574.	2.0	46
113	Interaction ofMTHFR1298C withACED Allele Augments the Risk of Diabetic Nephropathy in Western Iran. DNA and Cell Biology, 2012, 31, 553-559.	0.9	20
114	Apolipoprotein E Genotypes, Lipid Peroxidation, and Antioxidant Status among Mild and Severe Preeclamptic Women from Western Iran: Protective Role of Apolipoprotein Ïµ2 Allele in Severe Preeclampsia. Hypertension in Pregnancy, 2012, 31, 405-418.	0.5	17
115	Association of Endothelial Nitric Oxide Synthase Gene Variant (G894T) With Coronary Artery Disease in Western Iran. Angiology, 2012, 63, 131-137.	0.8	13
116	Lack of Association BetweenMTHFRC677T and A1298C Polymorphisms and Risk of Childhood Acute Lymphoblastic Leukemia in the Kurdish Population from Western Iran. Genetic Testing and Molecular Biomarkers, 2012, 16, 198-202.	0.3	17
117	Strong interaction between T allele of endothelial nitric oxide synthase with B1 allele of cholesteryl ester transfer protein TaqIB highly elevates the risk of coronary artery disease and type 2 diabetes mellitus. Human Genomics, 2012, 6, 20.	1.4	12
118	ACE insertion/deletion (I/D) polymorphism and diabetic nephropathy. Journal of Nephropathology, 2012, 1, 143-151.	0.1	67
119	Concomitant presence of endothelial nitric oxide 894T and angiotensin IIâ€converting enzyme D alleles are associated with diabetic nephropathy in a Kurdish population from Western Iran. Nephrology, 2012, 17, 175-181.	0.7	26
120	Thrombophilic mutations and susceptibility to preeclapmsia in Western Iran. Journal of Thrombosis and Thrombolysis, 2012, 33, 109-115.	1.0	18
121	Matrix metalloproteinas-9 functional promoter polymorphism 1562C>T increased risk of early-onset coronary artery disease. Molecular Biology Reports, 2012, 39, 555-562.	1.0	26
122	Thymidylate synthase and methionine synthase polymorphisms are not associated with susceptibility to childhood acute lymphoblastic leukemia in Kurdish population from Western Iran. Molecular Biology Reports, 2012, 39, 2195-2200.	1.0	14
123	Synergism between paraoxonase Arg 192 and the angiotensin converting enzyme D allele is associated with severity of coronary artery disease. Molecular Biology Reports, 2012, 39, 2723-2731.	1.0	3
124	ACE gene polymorphism and serum ACE activity in Iranians type II diabetic patients with macroalbuminuria. Molecular and Cellular Biochemistry, 2011, 346, 23-30.	1.4	33
125	Interaction of eNOS polymorphism with MTHFR variants increase the risk of diabetic nephropathy and its progression in type 2 diabetes mellitus patients. Molecular and Cellular Biochemistry, 2011, 353, 23-34.	1.4	35
126	The association between GSTT1, M1, and P1 polymorphisms with coronary artery disease in Western Iran. Molecular and Cellular Biochemistry, 2011, 354, 181-187.	1.4	29

#	Article	IF	Citations
127	Association between apolipoprotein Îµ4 allele, factor V Leiden, and plasma lipid and lipoprotein levels with sickle cell disease in southern Iran. Molecular Biology Reports, 2011, 38, 703-710.	1.0	14
128	The frequency of factor V Leiden mutation, ACE gene polymorphism, serum ACE activity and response to ACE inhibitor and angiotensin II receptor antagonist drugs in Iranians type II diabetic patients with microalbuminuria. Molecular Biology Reports, 2011, 38, 2117-2123.	1.0	34
129	Plasma lipids and lipoproteins in children and young adults with major Î²-thalassemia from western Iran: influence of genotype. Molecular Biology Reports, 2011, 38, 2573-2578.	1.0	11
130	Rapid separation of human globin chains in normal and thalassemia patients by RP-HPLC. Molecular Biology Reports, 2011, 38, 3213-3218.	1.0	15
131	Paraoxonase Arg 192 allele is an independent risk factor for three-vessel stenosis of coronary artery disease. Molecular Biology Reports, 2011, 38, 5421-5428.	1.0	24
132	Association Between Cholesteryl Ester Transfer Protein TaqIB Variants and Risk of Coronary Artery Disease and Diabetes Mellitus in the Population of Western Iran. Genetic Testing and Molecular Biomarkers, 2011, 15, 813-819.	0.3	14
133	SICKLE CELL DISEASE AND VENOUS THROMBOEMBOLISM. Mediterranean Journal of Hematology and Infectious Diseases, 2011, 3, e2011024.	0.5	19
134	Deep venous thrombosis and thrombophilic mutations in western Iran: association with factor V Leiden. Blood Coagulation and Fibrinolysis, 2010, 21, 385-388.	0.5	24
135	Abnormal hemoglobins among Kurdish population of Western Iran: hematological and molecular features. Molecular Biology Reports, 2010, 37, 51-57.	1.0	12
136	Detection of responsible mutations for beta thalassemia in the Kermanshah Province of Iran using PCR-based techniques. Molecular Biology Reports, 2010, 37, 149-154.	1.0	29
137	The Xmn1 polymorphic site 5â€² to the GÎ³ gene and its correlation to the GÎ³:AÎ³ ratio, age at first blood transfusion and clinical features in Î²-Thalassemia patients from Western Iran. Molecular Biology Reports, 2010, 37, 159-164.	1.0	36
138	Butyrylcholinesterase K variant and the APOE-Îµ4 allele work in synergy to increase the risk of coronary artery disease especially in diabetic patients. Molecular Biology Reports, 2010, 37, 2083-2091.	1.0	42
139	The angiotensin converting enzyme D allele is an independent risk factor for early onset coronary artery disease. Clinical Biochemistry, 2010, 43, 1189-1194.	0.8	46
140	Synergistic effects of the MTHFR C677T and A1298C polymorphisms on the increased risk of micro- and macro-albuminuria and progression of diabetic nephropathy among Iranians with type 2 diabetes mellitus. Clinical Biochemistry, 2010, 43, 1333-1339.	0.8	39
141	Cerebral Venous and Sinus Thrombosis and Thrombophilic Mutations in Western Iran: Association With Factor V Leiden. Clinical and Applied Thrombosis/Hemostasis, 2010, 16, 430-434.	0.7	15
142	The prevalence of factor V Leiden, prothrombin G20210A and methylenetetrahydrofolate reductase polymorphism C677T among G6PD deficient individuals from Western Iran. Molecular Biology Reports, 2009, 36, 2361-2364.	1.0	9
143	Serum butyrylcholinesterase activity and phenotype associations with lipid profile in stroke patients. Clinical Biochemistry, 2009, 42, 210-214.	0.8	21
144	Haplotype analysis of beta thalassemia patients in Western Iran. Blood Cells, Molecules, and Diseases, 2009, 42, 140-143.	0.6	24

#	Article	IF	Citations
145	Factor V G1691A, prothrombin G20210A and methylenetetrahydrofolate reductase polymorphism C677T are not associated with coronary artery disease and type 2 diabetes mellitus in western Iran. Blood Coagulation and Fibrinolysis, 2009, 20, 252-256.	0.5	22
146	Infant With Concomitant Presence of Hernia/Hydrocele and Primary Paratesticular Neuroblastoma. Journal of Pediatric Hematology/Oncology, 2009, 31, 349.	0.3	6
147	Association between enzymatic and non-enzymatic antioxidant defense mechanism with apolipoprotein E genotypes in Alzheimer disease. Clinical Biochemistry, 2008, 41, 932-936.	0.8	71
148	Prevalence of factor V Leiden (G1691A) and prothrombin (G20210A) among Kurdish population from Western Iran. Journal of Thrombosis and Thrombolysis, 2008, 25, 280-283.	1.0	28
149	Thrombophilic mutations among Southern Iranian patients with sickle cell disease: high prevalence of factor V Leiden. Journal of Thrombosis and Thrombolysis, 2008, 25, 288-292.	1.0	20
150	Prevalence of thrombotic risk factors among Î²-thalassemia patients from Western Iran. Journal of Thrombosis and Thrombolysis, 2008, 26, 229-233.	1.0	18
151	Prevalence of iron deficiency anemia among adolescent schoolgirls from Kermanshah, Western Iran. Hematology, 2008, 13, 352-355.	0.7	40
152	Molecular and hematologic analysis of hemoglobin Q-Iran and hemoglobin Setif in Iranian families. Archives of Iranian Medicine, 2008, 11, 382-6.	0.2	8
153	An Iranian Child With HbQ-Iran [Î±75 (EF4) Aspâ†'His]/â^'Î±3.7â€‰kb/IVSII.1 Gâ†'A. Journal of Pediatric Hematology/Oncology, 2007, 29, 649-651.	0.3	5
154	Determination of butyrylcholinesterase (BChE) phenotypes to predict the risk of prolonged apnea in persons receiving succinylcholine in the healthy population of western Iran. Clinical Biochemistry, 2007, 40, 629-633.	0.8	20
155	The presence of apolipoprotein Îµ4 and Îµ2 alleles augments the risk of coronary artery disease in type 2 diabetic patients. Clinical Biochemistry, 2007, 40, 1150-1156.	0.8	42
156	The Î²-GLobin Gene Haplotypes Associated With Hb D-Los Angeles [Î²121(GH4)Gluâ†'Gln] in Western Iran. Hemoglobin, 2006, 30, 39-44.	0.4	24
157	Plasma lipids in Iranians with sickle cell disease: Hypocholesterolemia in sickle cell anemia and increase of HDL-cholesterol in sickle cell trait. Clinica Chimica Acta, 2006, 365, 217-220.	0.5	40
158	Molecular characterization of glucose-6-phosphate dehydrogenase deficiency in the Kurdish population of Western Iran. Blood Cells, Molecules, and Diseases, 2006, 37, 91-94.	0.6	24
159	Association between apolipoprotein E polymorphism and serum lipid and apolipoprotein levels with Alzheimer's disease. Neuroscience Letters, 2006, 408, 68-72.	1.0	62
160	Hb D-Punjab [Î² 121 (GH4) Gluâ†'Gln]/Î²0-thalassemia [IVSII.1(Gâ†'A)] in two cases from an Iranian family: First report. American Journal of Hematology, 2006, 81, 302-303.	2.0	7
161	Implications of the Genetic Epidemiology of Globin Haplotypes Linked to the Sickle Cell Gene in Southern Iran. Human Biology, 2006, 78, 719-731.	0.4	30
162	Hemoglobinopathies in Iran: An Updated Review. International Journal of Hematology-Oncology and Stem Cell Research, 0, , .	0.3	4

#	Article	IF	Citations
163	The Prevalence of Hemoglobinopathies in Reference Laboratory of Kermanshah, Western Iran. Iranian Journal of Public Health, 0, , .	0.3	1