Mahsa M Amoli

List of Publications by Year
in descending order

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155 papers	2,179 citations	331259 21 h-index	329751 37 g-index
156 all docs	156 docs citations	156 times ranked	3011 citing authors

#	Article	IF	Citations
1	Competing endogenous RNA (ceRNA) cross talk and language in ceRNA regulatory networks: A new look at hallmarks of breast cancer. Journal of Cellular Physiology, 2019, 234, 10080-10100.	2.0	208
2	Genetic markers of disease susceptibility and severity in giant cell arteritis and polymyalgia rheumatica. Seminars in Arthritis and Rheumatism, 2003, 33, 38-48.	1.6	146
3	Polymorphism in the STAT6 gene encodes risk for nut allergy. Genes and Immunity, 2002, 3, 220-224.	2.2	73
4	VEGF gene polymorphism association with diabetic neuropathy. Molecular Biology Reports, 2010, 37, 3625-3630.	1.0	49
5	Interleukin 1 receptor antagonist gene polymorphism is associated with severe renal involvement and renal sequelae in Henoch-SchÃ¶nlein purpura. Journal of Rheumatology, 2002, 29, 1404-7.	1.0	49
6	Interleukin 8 gene polymorphism is associated with increased risk of nephritis in cutaneous vasculitis. Journal of Rheumatology, 2002, 29, 2367-70.	1.0	49
7	Association between Genetic Variants and Diabetes Mellitus in Iranian Populations: A Systematic Review of Observational Studies. Journal of Diabetes Research, 2015, 2015, 1-21.	1.0	48
8	Macrophage migration inhibitory factor gene polymorphism is associated with sarcoidosis in biopsy proven erythema nodosum. Journal of Rheumatology, 2002, 29, 1671-3.	1.0	44
9	Interleukin 1beta gene polymorphism association with severe renal manifestations and renal sequelae in Henoch-SchÃ¶nlein purpura. Journal of Rheumatology, 2004, 31, 295-8.	1.0	42
10	VEGF gene polymorphism association with diabetic foot ulcer. Diabetes Research and Clinical Practice, 2011, 93, 215-219.	1.1	40
11	Endothelial nitric oxide synthase VNTR (intron 4 a/b) polymorphism association with type 2 diabetes and its chronic complications. Diabetes Research and Clinical Practice, 2011, 91, 348-352.	1.1	39
12	Association between vitamin D receptor gene polymorphisms (Fok1 and Bsm1) and osteoporosis: a systematic review. Journal of Diabetes and Metabolic Disorders, 2014, 13, 98.	0.8	36
13	Survivin gene polymorphism association with papillary thyroid carcinoma. Pathology Research and Practice, 2012, 208, 100-103.	1.0	33
14	HLA-B35 association with nephritis in Henoch-SchÃ¶nlein purpura. Journal of Rheumatology, 2002, 29, 948-9.	1.0	31
15	Methylenetetrahydrofolate reductase gene polymorphism in diabetes and obesity. Molecular Biology Reports, 2010, 37, 105-109.	1.0	30
16	Gender-specific differences in the association of adiponectin gene polymorphisms with body mass index. Review of Diabetic Studies, 2010, 7, 241-6.	0.5	28
17	Replication of TCF7L2 rs7903146 association with type 2 diabetes in an Iranian population. Genetics and Molecular Biology, 2010, 33, 449-451.	0.6	27
18	Intercellular adhesion molecule-1 gene polymorphisms in isolated polymyalgia rheumatica. Journal of Rheumatology, 2002, 29, 502-4.	1.0	26

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19	eNOS gene polymorphism association with retinopathy in type 1 diabetes. Ophthalmic Genetics, 2010, 31, 103-107.	0.5	25
20	Inducible nitric oxide synthase polymorphism is associated with susceptibility to Henoch-SchÃ¶nlein purpura in northwestern Spain. Journal of Rheumatology, 2005, 32, 1081-5.	1.0	25
21	TGF-Î² and IL-23 gene expression in unstimulated PBMCs of patients with diabetes. Endocrine, 2012, 41, 430-434.	1.1	23
22	Vitamin D receptor gene FokI variant in diabetic foot ulcer and its relation with oxidative stress. Gene, 2017, 599, 87-91.	1.0	23
23	Association of Survivin Gene Polymorphism With Endometrial Cancer. International Journal of Gynecological Cancer, 2012, 22, 35-37.	1.2	22
24	The antioxidants dilemma: are they potentially immunosuppressants and carcinogens?. Frontiers in Physiology, 2014, 5, 245.	1.3	21
25	Association of interleukin-6 polymorphisms with obesity: A systematic review and meta-analysis. Cytokine, 2019, 123, 154769.	1.4	21
26	Curcumin Inhibits in Vitro MCP-1 Release From Mouse Pancreatic Islets. Transplantation Proceedings, 2006, 38, 3035-3038.	0.3	20
27	Mutation screening of VHL gene in a family with malignant bilateral pheochromocytoma: from isolated familial pheochromocytoma to von Hippel-Lindau disease. Familial Cancer, 2009, 8, 465-471.	0.9	20
28	VEGF gene mRNA expression in patients with coronary artery disease. Molecular Biology Reports, 2012, 39, 8595-8599.	1.0	20
29	A Genotype-First Approach for Clinical and Genetic Evaluation of Wolcott-Rallison Syndrome in a Large Cohort of Iranian Children With Neonatal Diabetes. Canadian Journal of Diabetes, 2018, 42, 272-275.	0.4	19
30	Apolipoprotein E gene polymorphism and total serum cholesterol level in Iranian population. Journal of Postgraduate Medicine, 2010, 56, 173-175.	0.2	19
31	SLC34A3 Intronic Deletion in a New Kindred with Hereditary Hypophosphatemic Rickets with Hypercalciuria. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2012, 4, 89-93.	0.4	18
32	TNF-Î± and IFN-Î³ gene variation and genetic susceptibility to type 1 diabetes and its microangiopathic complications. Journal of Diabetes and Metabolic Disorders, 2014, 13, 46.	0.8	18
33	Evaluation of the presence of Epstein-Barr virus (EBV) in Iranian patients with thyroid papillary carcinoma. Pathology Research and Practice, 2017, 213, 854-856.	1.0	18
34	MicroRNAâ€Ъinding site polymorphisms and risk of colorectal cancer: A systematic review and metaâ€ѕnalysis. Cancer Medicine, 2019, 8, 7477-7499.	1.3	18
35	Epistatic interactions between HLA-DRB1 and interleukin 4, but not interferon-gamma, increase susceptibility to giant cell arteritis. Journal of Rheumatology, 2004, 31, 2413-7.	1.0	18
36	Acarbose versus trans-chalcone: comparing the effect of two glycosidase inhibitors on obese mice. Archives of Endocrinology and Metabolism, 2015, 59, 202-209.	0.3	17

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37	The p.Arg435His Variation of IgG3 With High Affinity to FcRn Is Associated With Susceptibility for Pemphigus Vulgarisâ€"Analysis of Four Different Ethnic Cohorts. Frontiers in Immunology, 2018, 9, 1788.	2.2	17
38	Over-expression of TGF-Î²1 gene in medication free Schizophrenia. Psychoneuroendocrinology, 2019, 99, 265-270.	1.3	17
39	Prevalence of osteoporosis and vitamin D receptor gene polymorphisms (FokI) in an Iranian general population based study (Kurdistan) (IMOS). Medical Journal of the Islamic Republic of Iran, 2015, 29, 238.	0.9	17
40	A new frameshift MEN1 gene mutation associated with familial malignant insulinomas. Familial Cancer, 2011, 10, 343-348.	0.9	16
41	TGF-Î²1 and IGF-I gene variations in type 1 diabetes microangiopathic complications. Journal of Diabetes and Metabolic Disorders, 2014, 13, 45.	0.8	16
42	Endothelial nitric oxide synthase haplotype associations in biopsy-proven giant cell arteritis. Journal of Rheumatology, 2003, 30, 2019-22.	1.0	16
43	IL-23 Gene Expression in PBMCs of Patients with Coronary Artery Disease. Disease Markers, 2012, 33, 289-293.	0.6	15
44	Epistatic interaction between adiponectin and survivin gene polymorphisms in endometrial carcinoma. Pathology Research and Practice, 2015, 211, 293-297.	1.0	15
45	Potential role of gender specific effect of leptin receptor deficiency in an extended consanguineous family with severe early-onset obesity. European Journal of Medical Genetics, 2018, 61, 465-467.	0.7	15
46	Iranian neonatal diabetes mellitus due to mutation in PDX1 gene: a case report. Journal of Medical Case Reports, 2019, 13, 258.	0.4	15
47	Integrative analyses of triple negative dysregulated transcripts compared with nonâ€triple negative tumors and their functional and molecular interactions. Journal of Cellular Physiology, 2019, 234, 22386-22399.	2.0	15
48	Role of vitamin D and vitamin D receptor gene polymorphisms on residual beta cell function in children with type 1 diabetes mellitus. Pharmacological Reports, 2019, 71, 282-288.	1.5	15
49	Optimizing Conditions for Rat Pancreatic Islets Isolation. Cytotechnology, 2005, 48, 75-78.	0.7	14
50	Would blockage of cytokines improve the outcome of pancreatic islet transplantation?. Medical Hypotheses, 2006, 66, 816-819.	0.8	14
51	Effect of adiponectin gene polymorphisms on waist circumference in patients with diabetes. Journal of Diabetes and Metabolic Disorders, 2012, 11, 14.	0.8	14
52	Adiponectin gene variants and abdominal obesity in an Iranian population. Eating and Weight Disorders, 2017, 22, 85-90.	1.2	14
53	Methylomics of breast cancer: Seeking epimarkers in peripheral blood of young subjects. Tumor Biology, 2017, 39, 101042831769504.	0.8	14
54	Association Between the Polymorphism of Glu298Asp in Exon 7 of the eNOS Gene With Foot Ulcer and Oxidative Stress in Adult Patients With Type 2 Diabetes. Canadian Journal of Diabetes, 2018, 42, 18-22.	0.4	14

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55	Association of vascular endothelial growth factor (VEGF) Gene polymorphisms and expression with the risk of endometriosis: a caseâ€“control study. Molecular Biology Reports, 2019, 46, 3445-3450.	1.0	14
56	Polymorphisms of Antioxidant Genes as a Target for Diabetes Management. International Journal of Molecular and Cellular Medicine, 2017, 6, 135-147.	1.1	14
57	Corticotropin-releasing hormone promoter polymorphisms in patients with rheumatoid arthritis from northwest Spain. Journal of Rheumatology, 2003, 30, 913-7.	1.0	14
58	CXCL5 Gene Polymorphism Association with Diabetes Mellitus. Molecular Diagnosis and Therapy, 2008, 12, 391-394.	1.6	13
59	Associations between HLA-C alleles and papillary thyroid carcinoma. Cancer Biomarkers, 2009, 5, 19-22.	0.8	13
60	HLA-DR Association in Papillary Thyroid Carcinoma. Disease Markers, 2010, 28, 49-53.	0.6	13
61	Association between 318<scp>C</scp>/<scp>T</scp> polymorphism of the <scp>CTLA</scp>â€4 gene and systemic lupus erythematosus in Iranian patients. International Journal of Rheumatic Diseases, 2017, 20, 2040-2044.	0.9	13
62	CDH1 and DDR1 common variants confer risk to vitiligo and autoimmune comorbidities. Gene, 2019, 700, 17-22.	1.0	13
63	IL-23 gene expression in PBMCs of patients with coronary artery disease. Disease Markers, 2012, 33, 289-93.	0.6	13
64	RANTES gene mRNA expression and its â^'403 G/A promoter polymorphism in coronary artery disease. Gene, 2011, 487, 103-106.	1.0	12
65	Establishing a cGMP pancreatic islet processing facility: the first experience in Iran. Cell and Tissue Banking, 2012, 13, 569-575.	0.5	12
66	The Association between Genetic Variation in Wnt Transcription FactorTCF7L2(TCF4) and Alopecia Areata. Immunological Investigations, 2019, 48, 555-562.	1.0	12
67	Inflammation related miRNAs as an important player between obesity and cancers. Journal of Diabetes and Metabolic Disorders, 2019, 18, 675-692.	0.8	12
68	Apolipoprotein E gene polymorphism and its effect on anthropometric measures in normoglycemic subjects and type 2 diabetes. Journal of Diabetes and Metabolic Disorders, 2012, 11, 18.	0.8	11
69	Ectopic Cushing syndrome associated with thymic carcinoid tumor as the first presentation of MEN1 syndrome-report of a family with MEN1 gene mutation. Familial Cancer, 2014, 13, 267-272.	0.9	11
70	Association between Macrophage Migration Inhibitory Factor Gene Variation and Response to Glucocorticoid Treatment in Sudden Sensorineural Hearing Loss. Audiology and Neuro-Otology, 2015, 20, 376-382.	0.6	11
71	Macrophage migration inhibitory factor gene polymorphism is not associated with pemphigus vulgaris in Iranian patients. Journal of the European Academy of Dermatology and Venereology, 2013, 27, 1127-1131.	1.3	10
72	Association of MTHFR C677T polymorphism with elevated homocysteine level and disease development in vitiligo. International Journal of Immunogenetics, 2020, 47, 342-350.	0.8	10

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73	Influence of antioxidants' gene variants on risk of diabetes mellitus and its complications: a systematic review. Minerva Endocrinologica, 2019, 44, 310-325.	1.7	10
74	Association of microRNA gene polymorphisms with Type 2 diabetes mellitus: A systematic review and meta-analysis. Journal of Research in Medical Sciences, 2020, 25, 56.	0.4	10
75	SLC34A3 Intronic Deletion in an Iranian Kindred with Hereditary Hypophosphatemic Rickets with Hypercalciuria and Review of Reported Cases. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2018, 10, 343-349.	0.4	10
76	Detection of KCNJ11 Gene Mutations in a Family with Neonatal Diabetes Mellitus. Molecular Diagnosis and Therapy, 2012, 16, 109-114.	1.6	9
77	MIF 173 G>C variation was associated with depressive disorder in type 2 diabetes in an Iranian population. Psychoneuroendocrinology, 2019, 104, 243-248.	1.3	9
78	Haplotypes in vitamin D receptor gene encode risk in diabetic nephropathy. Gene, 2019, 683, 149-152.	1.0	9
79	HLA-DR association in papillary thyroid carcinoma. Disease Markers, 2010, 28, 49-53.	0.6	9
80	Henoch-SchÃ¶nlein purpura and cutaneous leukocytoclastic angiitis exhibit different HLA-DRB1 associations. Journal of Rheumatology, 2002, 29, 945-7.	1.0	9
81	MCP-1 gene haplotype association in biopsy proven giant cell arteritis. Journal of Rheumatology, 2005, 32, 507-10.	1.0	9
82	In vitro modulation of TCF7L2 gene expression in human pancreatic cells. Molecular Biology Reports, 2009, 36, 2329-2332.	1.0	8
83	Novel mutations of wolframin: a report with a look at the protein structure. Clinical Genetics, 2011, 79, 96-99.	1.0	8
84	eNOS Gene Variant in Patients with Coronary Artery Disease. Journal of Biomarkers, 2013, 2013, 1-6.	1.0	8
85	Effects of extremely low frequency electromagnetic fields on paraoxonase serum activity and lipid peroxidation metabolites in rat. Journal of Diabetes and Metabolic Disorders, 2014, 13, 85.	0.8	8
86	Autoimmune Polyglandular Syndrome Type 1: a case report. BMC Medical Genetics, 2019, 20, 143.	2.1	8
87	MTHFR AND ApoE genetic variants association with sudden sensorineural hearing loss. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2019, 40, 260-264.	0.6	8
88	Lack of association between macrophage migration inhibitory factor gene polymorphism and giant cell arteritis. Journal of Rheumatology, 2005, 32, 74-6.	1.0	8
89	Adenosine deaminase gene polymorphism is associated with obesity in Iranian population. Obesity Research and Clinical Practice, 2007, 1, 173-177.	0.8	7
90	Survivin Gene Polymorphism Association with Tongue Squamous Cell Carcinoma. Genetic Testing and Molecular Biomarkers, 2013, 17, 74-77.	0.3	7

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91	Association between MTHFR variant and diabetic neuropathy. Pharmacological Reports, 2018, 70, 1-5.	1.5	7
92	eNOS gene Glu298Asp variant confer risk in sudden sensorineural hearing loss. Acta Oto-Laryngologica, 2018, 138, 904-908.	0.3	7
93	An in silico approach to identify and prioritize miRNAs target sites polymorphisms in colorectal cancer and obesity. Cancer Medicine, 2020, 9, 9511-9528.	1.3	7
94	Interleukin-1 beta, interferon-gamma, and tumor necrosis factor-alpha gene expression in peripheral blood mononuclear cells of patients with coronary artery disease. ARYA Atherosclerosis, 2015, 11, 267-74.	0.4	7
95	Lack of association between endothelial nitric oxide synthase polymorphisms and Henoch-SchÃ¶nlein purpura. Journal of Rheumatology, 2004, 31, 299-301.	1.0	7
96	Two Polymorphisms in the Epithelial Cell-Derived Neutrophil-Activating Peptide (ENA-78) Gene. Disease Markers, 2005, 21, 75-77.	0.6	6
97	Mutation screening of RET proto-oncogene in a family with medullary thyroid carcinoma, marfanoid habitus and pheochromocytoma; from clinically MEN2B to genetically MEN2A syndrome. Endocrine, 2012, 42, 220-221.	1.1	6
98	A Novel Missense Mutation in Oncostatin M Receptor Beta Causing Primary Localized Cutaneous Amyloidosis. BioMed Research International, 2014, 2014, 1-6.	0.9	6
99	Sex-specific association of RANTES gene â˜’403 variant in Meniereâ€™s disease. European Archives of Oto-Rhino-Laryngology, 2015, 272, 2221-2225.	0.8	6
100	Liver alpha-amylase gene expression as an early obesity biomarker. Pharmacological Reports, 2017, 69, 229-234.	1.5	6
101	Molecular investigation of WFS1 gene exon 8 in Iranian patients with Wolfram syndrome. International Journal of Diabetes in Developing Countries, 2016, 36, 75-80.	0.3	5
102	Letter to the Editor: Comments on â€œAssociation between the ICAM-1 gene polymorphism and coronary heart disease risk: a meta-analysisâ€. Bioscience Reports, 2019, 39, .	1.1	5
103	Investigating Genetic Mutations in a Large Cohort of Iranian Patients with Congenital Hyperinsulinism. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2022, 14, 87-95.	0.4	5
104	Rapid detection of intercellular adhesion molecule 1 (G241R and K469E) polymorphisms by a novel PCR?SSP assay. Tissue Antigens, 2007, 69, 338-341.	1.0	4
105	Overall corrections and assessments of â€œCorrelations between TLR polymorphisms and inflammatory bowel disease: a meta-analysis of 49 case-control studiesâ€. Immunologic Research, 2019, 67, 301-303.	1.3	4
106	Macrophage migration inhibitory factor polymorphism (rs755622) in alopecia areata: a possible role in disease prevention. Archives of Dermatological Research, 2019, 311, 589-594.	1.1	4
107	Association of vitamin D receptor gene polymorphism with the occurrence of low bone density, osteopenia, and osteoporosis in patients with type 2 diabetes. Journal of Diabetes and Metabolic Disorders, 2021, 20, 1375-1383.	0.8	4
108	The Effect of Metformin on Expression of Long Non-coding RNA H19 in Endometrial Cancer. Medical Journal of the Islamic Republic of Iran, 2021, 35, 155.	0.9	4

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109	A New Mitochondrial Mutation in a Patient with Diabetes Mellitus, Deafness, Hydronephrosis and Joint Contractures. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 1185-9.	0.4	3
110	Is DNA methylation responsible for immune system dysfunction in schizophrenia?. Medical Hypotheses, 2011, 77, 573-579.	0.8	3
111	Effect of inulin supplementation in male mice fed with high fat diet on biochemical profile and Î±-amylase gene expression. Tropical Journal of Pharmaceutical Research, 2016, 15, 1197.	0.2	3
112	Investigating the association of rs2346061 (CNDP1), rs7577 (CNDP2), and rs1801133 (MTHFR) variants and homocysteine level with diabetic nephropathy in an Iranian population. Gene Reports, 2019, 16, 100443.	0.4	3
113	Is the +405 G/C single nucleotide polymorphism of the vascular endothelial growth factor (VEGF) gene associated with lateâ€onset vitiligo?. International Journal of Immunogenetics, 2019, 46, 241-246.	0.8	3
114	Investigating the association of matrix metalloproteinase-2 gene variants with endometriosis in an Iranian population. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2021, 258, 353-357.	0.5	3
115	Knowledge discovery in genetics of diabetes in Iran, a roadmap for future researches. Journal of Diabetes and Metabolic Disorders, 2021, 20, 1785-1791.	0.8	3
116	Absence of kl-vs Variant of Klotho Gene in Iranian Cardiac Patients (Comparison to the World) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 462	0.6	3
117	Absence of kl-vs variant of klotho gene in Iranian cardiac patients (comparison to the world) Tj ETQq1 1 0.784314 rgBT /Overlock 10	0.6	3
118	The Role of ERRFI1+808T/G Polymorphism in Diabetic Nephropathy. International Journal of Molecular and Cellular Medicine, 2019, 8, 49-55.	1.1	3
119	Association of <scp>miRNA</scp> targetome variants in <scp>LAMC1</scp> and <scp>GNB3</scp> genes with colorectal cancer and obesity. Cancer Medicine, 2022, 11, 3923-3938.	1.3	3
120	Local Insulin-Derived Amyloidosis Model Confronted with Silymarin: Histological Insights and Gene Expression of MMP, TNF-Î±, and IL-6. International Journal of Molecular Sciences, 2022, 23, 4952.	1.8	3
121	Severe Acanthosis Nigricans in a 17 Year-old Female with Partial Lipodystrophie Syndrome. Journal of Pediatric Endocrinology and Metabolism, 2008, 21, 1027-8.	0.4	2
122	No evidence of association betweenCTLA-4polymorphisms and systemic lupus erythematosus in Iranian patients. International Journal of Rheumatic Diseases, 2013, 16, 681-684.	0.9	2
123	Association between epstein barr virus and tongue squamous cell carcinoma in iranian patients. Pathology Research and Practice, 2018, 214, 130-133.	1.0	2
124	Comments on: â€œMeta-analysis of association between Arg326Gln (rs1503185) and Gln276Pro (rs1566734) polymorphisms of PTPRJ gene and cancer riskâ€• Journal of Applied Genetics, 2019, 60, 431-433.	1.0	2
125	Sex-dependent association of ACE (I/D) polymorphism with Meniere's disease. Meta Gene, 2020, 24, 100659.	0.3	2
126	Non-Muscle Myosin Heavy Chain 9 Gene (MYH9) Polymorphism (rs4821481) is Associated with Urinary Albumin Excretion in Iranian Diabetic Patients. Iranian Red Crescent Medical Journal, 2016, 19, .	0.5	2

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127	Early and delayed puberty among Iranian children with obesity. Minerva Endocrinology, 2020, , .	0.6	2
128	The Association Analysis of Vascular Endothelial Growth Factor -2549 Insertion/ Deletion Variant and Endometriosis Risk. International Journal of Molecular and Cellular Medicine, 2019, 8, 63-68.	1.1	2
129	Adenosine deaminase gene variant in diabetes and obesity. Journal of Diabetes and Metabolic Disorders, 2022, 21, 333-338.	0.8	2
130	Association of Pro-inflammatory Cytokine Gene Polymorphism with Meniereâ€™s Disease in an Iranian Sample. Iranian Journal of Allergy, Asthma and Immunology, 2021, 20, 734-739.	0.3	2
131	A patient with features of albright hereditory osteodystrophy and unusual neuropsychiatric findings without coding Gsalpha mutations. Journal of Diabetes and Metabolic Disorders, 2014, 13, 56.	0.8	1
132	Association between Trp48Arg polymorphism of the CD11c gene and risk for obesity among Iranian population. Journal of Diabetes and Metabolic Disorders, 2018, 17, 197-201.	0.8	1
133	A case of H syndrome with a novel mutation in SLC29A3. Meta Gene, 2019, 21, 100599.	0.3	1
134	Re: â€œAssociation Between the CYP4F2 Gene rs1558139 and rs2108622 Polymorphisms and Hypertension: A Meta-Analysisâ€•by Geng et al. (Genet Test Mol Biomarkers 2019;23:342â€“347; DOI: 10.1089/gtmb.2018.0202). Genetic Testing and Molecular Biomarkers, 2019, 23, 696-697.	0.3	1
135	Comments on and assessments of â€˜Associations between endothelial nitric oxide synthase gene polymorphisms and the risk of coronary artery disease: A systematic review and meta-analysis of 132 caseâ€“control studiesâ€™. European Journal of Preventive Cardiology, 2020, 27, 660-663.	0.8	1
136	Comments on and assessments of â€˜Associations between FCGR polymorphisms and immune thrombocytopenia: A metaâ€analysisâ€™. Scandinavian Journal of Immunology, 2020, 91, e12815.	1.3	1
137	Hydro alcoholic green tea extract effect on high fat diet treated NMRI mice and 3T3L1 cells. Journal of Diabetes and Metabolic Disorders, 2021, 20, 641-648.	0.8	1
138	Investigation of TGF-Î²1 gene variant and expression in a group of Iranian women with endometriosis. Archives of Gynecology and Obstetrics, 2021, 304, 1527-1534.	0.8	1
139	Association of the CTLA-4 1722TC polymorphism and systemic lupus erythematosus: a systematic review and meta analysis. Medical Journal of the Islamic Republic of Iran, 2014, 28, 132.	0.9	1
140	HLA-Cw Allele Frequency in Definite Meniere's Disease Compared to Probable Meniere's Disease and Healthy Controls in an Iranian Sample. Iranian Journal of Otorhinolaryngology, 2016, 28, 262-6.	0.4	1
141	Lack of association between ICAM-1 gene polymorphisms and biopsy-proven erythema nodosum. Journal of Rheumatology, 2004, 31, 403-5.	1.0	1
142	Role of genetic polymorphisms in recurrent aphthous stomatitis: A systematic review and meta-analysis. Cytokine, 2022, 153, 155864.	1.4	1
143	Comments on â€œCorrelation between PNPLA3 rs738409 polymorphism and hepatocellular carcinoma: a meta-analysis of 10,330 subjectsâ€• International Journal of Biological Markers, 2019, 34, 322-324.	0.7	0
144	Comments on â€œAssociation between miR-499 rs3746444 polymorphism and coronary heart disease susceptibility: An evidence-based meta-analysis of 5063 cases and 4603 controlsâ€• Gene, 2019, 707, 100-102.	1.0	0

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145	Comments on: â€œA Meta-Analysis of the Association between Microrna-196A2 and Risk of Ischemic Stroke and Coronary Artery Disease in Asian Populationâ€• Journal of Stroke and Cerebrovascular Diseases, 2019, 28, 1409-1411.	0.7	0
146	Comments on and assessment of â€œThe rs498872 polymorphism is associated with an elevated susceptibility to glioma: a meta-analysis of 36,264 subjectsâ€• Acta Neurologica Belgica, 2020, 120, 1201-1202.	0.5	0
147	Letter to the editor: Association between Î±â€•adducin rs4961 polymorphism and hypertension: A metaâ€•analysis based on 40â€‰432 subjects. Journal of Cellular Biochemistry, 2020, 121, 2728-2729.	1.2	0
148	Comments on â€œEffects of MTNR1B Genetic Variants on Individual Susceptibility to Gestational Diabetes Mellitus: A Meta-Analysisâ€• American Journal of Perinatology, 2021, 38, 310-312.	0.6	0
149	Thyroid Cancer research at endocrinology and metabolism research institute (EMRI): a report of scientific activities between 2005 and 2020. Journal of Diabetes and Metabolic Disorders, 0, , 1.	0.8	0
150	Genetic research in Immunogenetics Group of Endocrinology and Metabolism Research Institute. Journal of Diabetes and Metabolic Disorders, 0, , 1.	0.8	0
151	Evaluation of ERRFI1 +808 T/G variant and its mRNA expression in coronary artery in-stent restenosis. Gene Reports, 2021, 24, 101248.	0.4	0
152	ADA gene haplotype is associated with coronary-in-stent-restenosis. Molecular Biology Reports, 2021, 48, 6665-6671.	1.0	0
153	VEGFA gene haplotypes in Meniere's disease. Gene Reports, 2021, 24, 101244.	0.4	0
154	Meglitinide (repaglinide) therapy in permanent neonatal diabetes mellitus: twoÂcase reports. Journal of Medical Case Reports, 2021, 15, 535.	0.4	0
155	Practical evaluation of late-night salivary cortisol: a real-life approach. Endocrine, 2012, 42, 220-1.	1.1	0