Mahsa M Amoli

List of Publications by Year in Descending Order

150 papers	1,697 citations	20 h-index	34 g-index
156 ext. papers	1,949 ext. citations	2.9 avg, IF	4.61 L-index

#	Paper	IF	Citations
150	Role of genetic polymorphisms in recurrent aphthous stomatitis: A systematic review and meta-analysis.. *Cytokine*, **2022**, 153, 155864	4	
149	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	1.1	0
148	Meglitinide (repaglinide) therapy in permanent neonatal diabetes mellitus: two case reports. *Journal of Medical Case Reports*, **2021**, 15, 535	1.2	
147	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	2.5	0
146	Knowledge discovery in genetics of diabetes in Iran, a roadmap for future researches.. *Journal of Diabetes and Metabolic Disorders*, **2021**, 20, 1785-1791	2.5	1
145	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	2.4	1
144	Comments on "Effects of MTNR1B Genetic Variants on Individual Susceptibility to Gestational Diabetes Mellitus: A Meta-Analysis". *American Journal of Perinatology*, **2021**, 38, 310-312	3.3	
143	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	2.5	0
142	Investigation of TGF-β1 gene variant and expression in a group of Iranian women with endometriosis. *Archives of Gynecology and Obstetrics*, **2021**, 304, 1527-1534	2.5	
141	Evaluation of ERRFI1 +808 T/G variant and its mRNA expression in coronary artery in-stent restenosis. *Gene Reports*, **2021**, 24, 101248	1.4	
140	ADA gene haplotype is associated with coronary-in-stent-restenosis. *Molecular Biology Reports*, **2021**, 48, 6665-6671	2.8	
139	VEGFA gene haplotypes in Meniere's disease. *Gene Reports*, **2021**, 24, 101244	1.4	
138	Association of Pro-inflammatory Cytokine Gene Polymorphism with Meniere's Disease in an Iranian Sample.. *Iranian Journal of Allergy, Asthma and Immunology*, **2021**, 20, 734-739	1.1	0
137	Association of MTHFR C677T polymorphism with elevated homocysteine level and disease development in vitiligo. *International Journal of Immunogenetics*, **2020**, 47, 342-350	2.3	7
136	Letter to the editor: Association between α-adducin rs4961 polymorphism and hypertension: A meta-analysis based on 40 432 subjects. *Journal of Cellular Biochemistry*, **2020**, 121, 2728-2729	4.7	
135	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	1.6	6
134	An in silico approach to identify and prioritize miRNAs target sites polymorphisms in colorectal cancer and obesity. *Cancer Medicine*, **2020**, 9, 9511-9528	4.8	3

133	Sex-dependent association of ACE (I/D) polymorphism with Meniere's disease. *Meta Gene*, **2020**, 24, 100659	0.7	1
132	Comments on and assessment of "The rs498872 polymorphism is associated with an elevated susceptibility to glioma: a meta-analysis of 36,264 subjects". *Acta Neurologica Belgica*, **2020**, 120, 1201-1202	1.5	
131	Comments on and assessments of "Associations between endothelial nitric oxide synthase gene polymorphisms and the risk of coronary artery disease: A systematic review and meta-analysis of 132 case-control studies". *European Journal of Preventive Cardiology*, **2020**, 27, 660-663	3.9	0
130	Comments on and assessments of "Associations between FCGR polymorphisms and immune thrombocytopenia: A meta-analysis". *Scandinavian Journal of Immunology*, **2020**, 91, e12815	3.4	1
129	Autoimmune Polyglandular Syndrome Type 1: a case report. *BMC Medical Genetics*, **2019**, 20, 143	2.1	6
128	Association of interleukin-6 polymorphisms with obesity: A systematic review and meta-analysis. *Cytokine*, **2019**, 123, 154769	4	11
127	"Association Between the Gene rs1558139 and rs2108622 Polymorphisms and Hypertension: A Meta-Analysis" by Geng (Genet Test Mol Biomarkers 2019;23:342-347; DOI: 10.1089/gtmb.2018.0202). *Genetic Testing and Molecular Biomarkers*, **2019**, 23, 696-697	1.6	1
126	Is the +405 G/C single nucleotide polymorphism of the vascular endothelial growth factor (VEGF) gene associated with late-onset vitiligo?. *International Journal of Immunogenetics*, **2019**, 46, 241-246	2.3	3
125	Macrophage migration inhibitory factor polymorphism (rs755622) in alopecia areata: a possible role in disease prevention. *Archives of Dermatological Research*, **2019**, 311, 589-594	3.3	3
124	Comments on "Association between miR-499 rs3746444 polymorphism and coronary heart disease susceptibility: An evidence-based meta-analysis of 5063 cases and 4603 controls". *Gene*, **2019**, 707, 100-102	3.8	
123	Integrative analyses of triple negative dysregulated transcripts compared with non-triple negative tumors and their functional and molecular interactions. *Journal of Cellular Physiology*, **2019**, 234, 22386-22399	4	10
122	Letter to the Editor: Comments on "Association between the ICAM-1 gene polymorphism and coronary heart disease risk: a meta-analysis". *Bioscience Reports*, **2019**, 39,	4.1	3
121	Association of vascular endothelial growth factor (VEGF) Gene polymorphisms and expression with the risk of endometriosis: a case-control study. *Molecular Biology Reports*, **2019**, 46, 3445-3450	2.8	10
120	The Association between Genetic Variation in Wnt Transcription Factor () and Alopecia Areata. *Immunological Investigations*, **2019**, 48, 555-562	2.9	7
119	Comments on: "A Meta-Analysis of the Association between Microrna-196A2 and Risk of Ischemic Stroke and Coronary Artery Disease in Asian Population". *Journal of Stroke and Cerebrovascular Diseases*, **2019**, 28, 1409-1411	2.8	
118	CDH1 and DDR1 common variants confer risk to vitiligo and autoimmune comorbidities. *Gene*, **2019**, 700, 17-22	3.8	8
117	MIF 173 G>C variation was associated with depressive disorder in type 2 diabetes in an Iranian population. *Psychoneuroendocrinology*, **2019**, 104, 243-248	5	7
116	Iranian neonatal diabetes mellitus due to mutation in PDX1 gene: a case report. *Journal of Medical Case Reports*, **2019**, 13, 258	1.2	9

115	A case of H syndrome with a novel mutation in SLC29A3. *Meta Gene*, **2019**, 21, 100599	0.7	1
114	Comments on: "Meta-analysis of association between Arg326Gln (rs1503185) and Gln276Pro (rs1566734) polymorphisms of PTPRJ gene and cancer risk". *Journal of Applied Genetics*, **2019**, 60, 431-433	2.5	1
113	Comments on "Correlation between rs738409 polymorphism and hepatocellular carcinoma: a meta-analysis of 10,330 subjects". *International Journal of Biological Markers*, **2019**, 34, 322-324	2.8	
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	1.4	2
111	Overall corrections and assessments of "Correlations between TLR polymorphisms and inflammatory bowel disease: a meta-analysis of 49 case-control studies". *Immunologic Research*, **2019**, 67, 301-303	4.3	2
110	MicroRNA-binding site polymorphisms and risk of colorectal cancer: A systematic review and meta-analysis. *Cancer Medicine*, **2019**, 8, 7477-7499	4.8	13
109	The Role of ERRFI1+808T/G Polymorphism in Diabetic Nephropathy. *International Journal of Molecular and Cellular Medicine*, **2019**, 8, 49-55	1.2	3
108	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	1.2	2
107	Influence of antioxidants Q gene variants on risk of diabetes mellitus and its complications: a systematic review. *Minerva Endocrinologica*, **2019**, 44, 310-325	1.9	6
106	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	3.9	12
105	Inflammation related miRNAs as an important player between obesity and cancers. *Journal of Diabetes and Metabolic Disorders*, **2019**, 18, 675-692	2.5	4
104	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	2.8	5
103	Over-expression of TGF-1 gene in medication free Schizophrenia. *Psychoneuroendocrinology*, **2019**, 99, 265-270	5	9
102	Haplotypes in vitamin D receptor gene encode risk in diabetic nephropathy. *Gene*, **2019**, 683, 149-152	3.8	7
101	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	7	153
100	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	2.6	8
99	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	2.1	8
98	Association between MTHFR variant and diabetic neuropathy. *Pharmacological Reports*, **2018**, 70, 1-5	3.9	6

97	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	2.1	11
96	Association between epstein barr virus and tongue squamous cell carcinoma in iranian patients. *Pathology Research and Practice*, **2018**, 214, 130-133	3.4	
95	Intronic Deletion in an Iranian Kindred with Hereditary Hypophosphatemic Rickets with Hypercalciuria. *JCRPE Journal of Clinical Research in Pediatric Endocrinology*, **2018**, 10, 343-349	1.9	8
94	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	2.5	1
93	eNOS gene Glu298Asp variant confer risk in sudden sensorineural hearing loss. *Acta Oto-Laryngologica*, **2018**, 138, 904-908	1.6	2
92	The p.Arg435His Variation of IgG3 With High Affinity to FcRn Is Associated With Susceptibility for Pemphigus Vulgaris-Analysis of Four Different Ethnic Cohorts. *Frontiers in Immunology*, **2018**, 9, 1788	8.4	10
91	Association between 318C/T polymorphism of the CTLA-4 gene and systemic lupus erythematosus in Iranian patients. *International Journal of Rheumatic Diseases*, **2017**, 20, 2040-2044	2.3	8
90	Adiponectin gene variants and abdominal obesity in an Iranian population. *Eating and Weight Disorders*, **2017**, 22, 85-90	3.6	9
89	Methylomics of breast cancer: Seeking epimarkers in peripheral blood of young subjects. *Tumor Biology*, **2017**, 39, 1010428317695040	2.9	11
88	Evaluation of the presence of Epstein-Barr virus (EBV) in Iranian patients with thyroid papillary carcinoma. *Pathology Research and Practice*, **2017**, 213, 854-856	3.4	11
87	Vitamin D receptor gene FokI variant in diabetic foot ulcer and its relation with oxidative stress. *Gene*, **2017**, 599, 87-91	3.8	12
86	Liver alpha-amylase gene expression as an early obesity biomarker. *Pharmacological Reports*, **2017**, 69, 229-234	3.9	6
85	Polymorphisms of Antioxidant Genes as a Target for Diabetes Management. *International Journal of Molecular and Cellular Medicine*, **2017**, 6, 135-147	1.2	6
84	Molecular investigation of WFS1 gene exon 8 in Iranian patients with Wolfram syndrome. *International Journal of Diabetes in Developing Countries*, **2016**, 36, 75-80	0.8	4
83	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	0.6	1
82	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,	1.3	2
81	Effect of inulin supplementation in male mice fed with high fat diet on biochemical profile and □amylase gene expression. *Tropical Journal of Pharmaceutical Research*, **2016**, 15, 1197	0.8	3
80	Epistatic interaction between adiponectin and survivin gene polymorphisms in endometrial carcinoma. *Pathology Research and Practice*, **2015**, 211, 293-7	3.4	13

79	Acarbose versus trans-chalcone: comparing the effect of two glycosidase inhibitors on obese mice. *Archives of Endocrinology and Metabolism*, **2015**, 59, 202-9	2.2	14
78	Association between Genetic Variants and Diabetes Mellitus in Iranian Populations: A Systematic Review of Observational Studies. *Journal of Diabetes Research*, **2015**, 2015, 585917	3.9	35
77	Sex-specific association of RANTES gene -403 variant in Meniere disease. *European Archives of Oto-Rhino-Laryngology*, **2015**, 272, 2221-5	3.5	2
76	Association between Macrophage Migration Inhibitory Factor Gene Variation and Response to Glucocorticoid Treatment in Sudden Sensorineural Hearing Loss. *Audiology and Neuro-Otology*, **2015**, 20, 376-82	2.2	11
75	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	0.7	6
74	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	1.1	15
73	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	2.5	1
72	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	2.5	5
71	TGF-1 and IGF-I gene variations in type 1 diabetes microangiopathic complications. *Journal of Diabetes and Metabolic Disorders*, **2014**, 13, 45	2.5	13
70	TNF- and IFN- gene variation and genetic susceptibility to type 1 diabetes and its microangiopathic complications. *Journal of Diabetes and Metabolic Disorders*, **2014**, 13, 46	2.5	13
69	A novel missense mutation in oncostatin M receptor beta causing primary localized cutaneous amyloidosis. *BioMed Research International*, **2014**, 2014, 653724	3	6
68	The antioxidants dilemma: are they potentially immunosuppressants and carcinogens?. *Frontiers in Physiology*, **2014**, 5, 245	4.6	15
67	Association between vitamin D receptor gene polymorphisms (Fok1 and Bsm1) and osteoporosis: a systematic review. *Journal of Diabetes and Metabolic Disorders*, **2014**, 13, 98	2.5	31
66	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, 267-72	3	9
65	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	1.1	1
64	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, 1127-31	4.6	10
63	Survivin gene polymorphism association with tongue squamous cell carcinoma. *Genetic Testing and Molecular Biomarkers*, **2013**, 17, 74-7	1.6	5
62	No evidence of association between CTLA-4 polymorphisms and systemic lupus erythematosus in Iranian patients. *International Journal of Rheumatic Diseases*, **2013**, 16, 681-4	2.3	1

61	eNOS Gene Variant in Patients with Coronary Artery Disease. *Journal of Biomarkers*, **2013**, 2013, 403783	0	7
60	Establishing a cGMP pancreatic islet processing facility: the first experience in Iran. *Cell and Tissue Banking*, **2012**, 13, 569-75	2.2	10
59	Survivin gene polymorphism association with papillary thyroid carcinoma. *Pathology Research and Practice*, **2012**, 208, 100-3	3.4	24
58	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	2.5	8
57	Effect of adiponectin gene polymorphisms on waist circumference in patients with diabetes. *Journal of Diabetes and Metabolic Disorders*, **2012**, 11, 14	2.5	11
56	Detection of KCNJ11 gene mutations in a family with neonatal diabetes mellitus: implications for therapeutic management of family members with long-standing disease. *Molecular Diagnosis and Therapy*, **2012**, 16, 109-14	4.5	8
55	IL-23 Gene Expression in PBMCs of Patients with Coronary Artery Disease. *Disease Markers*, **2012**, 33, 289-293	3.2	13
54	VEGF gene mRNA expression in patients with coronary artery disease. *Molecular Biology Reports*, **2012**, 39, 8595-9	2.8	18
53	TGF- and IL-23 gene expression in unstimulated PBMCs of patients with diabetes. *Endocrine*, **2012**, 41, 430-4	4	18
52	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	4	5
51	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	1.9	11
50	Association of survivin gene polymorphism with endometrial cancer. *International Journal of Gynecological Cancer*, **2012**, 22, 35-7	3.5	17
49	IL-23 gene expression in PBMCs of patients with coronary artery disease. *Disease Markers*, **2012**, 33, 289-93	3.2	10
48	Practical evaluation of late-night salivary cortisol: a real-life approach. *Endocrine*, **2012**, 42, 220-1	4	
47	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, 348-52	7.4	30
46	VEGF gene polymorphism association with diabetic foot ulcer. *Diabetes Research and Clinical Practice*, **2011**, 93, 215-219	7.4	32
45	RANTES gene mRNA expression and its -403 G/A promoter polymorphism in coronary artery disease. *Gene*, **2011**, 487, 103-6	3.8	11
44	Is DNA methylation responsible for immune system dysfunction in schizophrenia?. *Medical Hypotheses*, **2011**, 77, 573-9	3.8	2

43	Novel mutations of wolframin: a report with a look at the protein structure. *Clinical Genetics*, **2011**, 79, 96-9	4	7
42	A new frameshift MEN1 gene mutation associated with familial malignant insulinomas. *Familial Cancer*, **2011**, 10, 343-8	3	13
41	Absence of kl-vs Variant of Klotho Gene in Iranian Cardiac Patients (Comparison to the World Populations). *Disease Markers*, **2011**, 31, 211-214	3.2	3
40	Absence of kl-vs variant of klotho gene in Iranian cardiac patients (comparison to the world populations). *Disease Markers*, **2011**, 31, 211-4	3.2	2
39	HLA-DR Association in Papillary Thyroid Carcinoma. *Disease Markers*, **2010**, 28, 49-53	3.2	12
38	Replication of TCF7L2 rs7903146 association with type 2 diabetes in an Iranian population. *Genetics and Molecular Biology*, **2010**, 33, 449-51	2	22
37	eNOS gene polymorphism association with retinopathy in type 1 diabetes. *Ophthalmic Genetics*, **2010**, 31, 103-7	1.2	21
36	Methylenetetrahydrofolate reductase gene polymorphism in diabetes and obesity. *Molecular Biology Reports*, **2010**, 37, 105-9	2.8	26
35	VEGF gene polymorphism association with diabetic neuropathy. *Molecular Biology Reports*, **2010**, 37, 3625-30	2.8	41
34	HLA-DR association in papillary thyroid carcinoma. *Disease Markers*, **2010**, 28, 49-53	3.2	9
33	Gender-specific differences in the association of adiponectin gene polymorphisms with body mass index. *Review of Diabetic Studies*, **2010**, 7, 241-6	3.6	25
32	Apolipoprotein E gene polymorphism and total serum cholesterol level in Iranian population. *Journal of Postgraduate Medicine*, **2010**, 56, 173-5	0.8	16
31	Associations between HLA-C alleles and papillary thyroid carcinoma. *Cancer Biomarkers*, **2009**, 5, 19-22	3.8	10
30	In vitro modulation of TCF7L2 gene expression in human pancreatic cells. *Molecular Biology Reports*, **2009**, 36, 2329-32	2.8	7
29	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, 465-71	3	16
28	CXCL5 gene polymorphism association with diabetes mellitus. *Molecular Diagnosis and Therapy*, **2008**, 12, 391-4	4.5	8
27	Severe acanthosis nigricans in a 17 year-old female with partial lipodystrophic syndrome. *Journal of Pediatric Endocrinology and Metabolism*, **2008**, 21, 1027-8	1.6	1
26	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	1.6	3

25	Adenosine deaminase gene polymorphism is associated with obesity in Iranian population. *Obesity Research and Clinical Practice*, **2007**, 1, I-II	5.4	5
24	Rapid detection of intercellular adhesion molecule 1 (G241R and K469E) polymorphisms by a novel PCR-SSP assay. *Tissue Antigens*, **2007**, 69, 338-41		4
23	Would blockage of cytokines improve the outcome of pancreatic islet transplantation?. *Medical Hypotheses*, **2006**, 66, 816-9	3.8	14
22	Curcumin inhibits in vitro MCP-1 release from mouse pancreatic islets. *Transplantation Proceedings*, **2006**, 38, 3035-8	1.1	17
21	Optimizing conditions for rat pancreatic islets isolation. *Cytotechnology*, **2005**, 48, 75-8	2.2	11
20	Two polymorphisms in the epithelial cell-derived neutrophil-activating peptide (ENA-78) gene. *Disease Markers*, **2005**, 21, 75-7	3.2	5
19	Lack of association between macrophage migration inhibitory factor gene polymorphism and giant cell arteritis. *Journal of Rheumatology*, **2005**, 32, 74-6	4.1	6
18	MCP-1 gene haplotype association in biopsy proven giant cell arteritis. *Journal of Rheumatology*, **2005**, 32, 507-10	4.1	7
17	Inducible nitric oxide synthase polymorphism is associated with susceptibility to Henoch-Schönlein purpura in northwestern Spain. *Journal of Rheumatology*, **2005**, 32, 1081-5	4.1	24
16	Interleukin 1beta gene polymorphism association with severe renal manifestations and renal sequelae in Henoch-Schönlein purpura. *Journal of Rheumatology*, **2004**, 31, 295-8	4.1	42
15	Lack of association between endothelial nitric oxide synthase polymorphisms and Henoch-Schönlein purpura. *Journal of Rheumatology*, **2004**, 31, 299-301	4.1	7
14	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	4.1	16
13	Genetic markers of disease susceptibility and severity in giant cell arteritis and polymyalgia rheumatica. *Seminars in Arthritis and Rheumatism*, **2003**, 33, 38-48	5.3	120
12	Corticotropin-releasing hormone promoter polymorphisms in patients with rheumatoid arthritis from northwest Spain. *Journal of Rheumatology*, **2003**, 30, 913-7	4.1	14
11	Endothelial nitric oxide synthase haplotype associations in biopsy-proven giant cell arteritis. *Journal of Rheumatology*, **2003**, 30, 2019-22	4.1	14
10	Polymorphism in the STAT6 gene encodes risk for nut allergy. *Genes and Immunity*, **2002**, 3, 220-4	4.4	63
9	Intercellular adhesion molecule-1 gene polymorphisms in isolated polymyalgia rheumatica. *Journal of Rheumatology*, **2002**, 29, 502-4	4.1	25
8	Henoch-Schönlein purpura and cutaneous leukocytoclastic angiitis exhibit different HLA-DRB1 associations. *Journal of Rheumatology*, **2002**, 29, 945-7	4.1	8

7	HLA-B35 association with nephritis in Henoch-Schönlein purpura. *Journal of Rheumatology*, **2002**, 29, 948-9	4.1	31
6	Interleukin 1 receptor antagonist gene polymorphism is associated with severe renal involvement and renal sequelae in Henoch-Schönlein purpura. *Journal of Rheumatology*, **2002**, 29, 1404-7	4.1	47
5	Macrophage migration inhibitory factor gene polymorphism is associated with sarcoidosis in biopsy proven erythema nodosum. *Journal of Rheumatology*, **2002**, 29, 1671-3	4.1	39
4	Interleukin 8 gene polymorphism is associated with increased risk of nephritis in cutaneous vasculitis. *Journal of Rheumatology*, **2002**, 29, 2367-70	4.1	48
3	Adenosine deaminase gene variant in diabetes and obesity. *Journal of Diabetes and Metabolic Disorders*,1	2.5	0
2	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*,1	2.5	
1	Genetic research in Immunogenetics Group of Endocrinology and Metabolism Research Institute. *Journal of Diabetes and Metabolic Disorders*,1	2.5	