Saeedeh Salimi

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9089915/publications.pdf

Version: 2024-02-01

304743 434195 1,647 115 22 31 citations h-index g-index papers 117 117 117 2309 docs citations times ranked citing authors all docs

#	Article	lF	CITATIONS
1	Activity of antioxidant enzymes in seminal plasma and their relationship with lipid peroxidation of spermatozoa. International Braz J Urol: Official Journal of the Brazilian Society of Urology, 2008, 34, 485-491.	1.5	73
2	Lipid composition of spermatozoa in normozoospermic and asthenozoospermic males. Prostaglandins Leukotrienes and Essential Fatty Acids, 2007, 77, 45-50.	2.2	65
3	Different Profile of Serum Leptin between Early Onset and Late Onset Preeclampsia. Disease Markers, 2014, 2014, 1-7.	1.3	49
4	Docosahexaenoic acid sensitizes Ramos cells to Gamma-irradiation-induced apoptosis through involvement of PPAR-Î ³ activation and NF-Î ⁸ B suppression. Molecular and Cellular Biochemistry, 2008, 317, 113-120.	3.1	46
5	Involvement of PPAR- \hat{i}^3 and p53 in DHA-induced apoptosis in Reh cells. Molecular and Cellular Biochemistry, 2007, 304, 71-77.	3.1	43
6	Relationship between seminal antioxidant enzymes and the phospholipid and fatty acid composition of spermatozoa. Reproductive BioMedicine Online, 2008, 16, 649-656.	2.4	40
7	Association between vitamin D receptor polymorphisms and haplotypes with pulmonary tuberculosis. Biomedical Reports, 2015, 3, 189-194.	2.0	36
8	Vitamin D Receptor Gene Polymorphism and the Risk of Multiple Sclerosis in South Eastern of Iran. Journal of Molecular Neuroscience, 2015, 56, 572-576.	2.3	34
9	The neuroprotective effects of hydro-alcoholic extract of olive (Olea europaea L.) leaf on rotenone-induced Parkinson's disease in rat. Metabolic Brain Disease, 2018, 33, 79-88.	2.9	32
10	Association of Angiotensin-Converting Enzyme Intron 16 Insertion/Deletion and Angiotensin II Type 1 Receptor A1166C Gene Polymorphisms with Preeclampsia in South East of Iran. Journal of Biomedicine and Biotechnology, $2011, 2011, 1$ -6.	3.0	31
11	Interleukin- 1^2 (IL- 1^2) & amp; IL-4 gene polymorphisms in patients with systemic lupus erythematosus (SLE) & amp; their association with susceptibility to SLE. Indian Journal of Medical Research, 2016, 143, 591.	1.0	30
12	Polymorphisms of the folate metabolizing enzymes: Association with SLE susceptibility and in silico analysis. Gene, 2017, 637, 161-172.	2.2	29
13	Effects of deficit irrigation on some physiological traits, production and fruit quality of  Mazafati' date palm and the fruit wilting and dropping disorder. Agricultural Water Management, 2018, 209, 219-227.	5.6	29
14	Possible Association of IL-4 VNTR Polymorphism with Susceptibility to Preeclampsia. BioMed Research International, 2014, 2014, 1-5.	1.9	28
15	The early-onset preeclampsia is associated with MTHFR and FVL polymorphisms. Archives of Gynecology and Obstetrics, 2015, 291, 1303-1312.	1.7	28
16	Association between the CD14 gene C-159T polymorphism and serum soluble CD14 with pulmonary tuberculosis. International Journal of Tuberculosis and Lung Disease, 2012, 16, 1383-1387.	1.2	25
17	The association of the placental Hypoxia-inducible factor $1-\hat{l}\pm$ polymorphisms and HIF1- $\hat{l}\pm$ mRNA expression with preeclampsia. Placenta, 2018, 67, 31-37.	1.5	25
18	Genetic and epigenetic analysis of the BAX and BCL2 in the placenta of pregnant women complicated by preeclampsia. Apoptosis: an International Journal on Programmed Cell Death, 2019, 24, 301-311.	4.9	25

#	Article	IF	CITATIONS
19	Association between <i>TLR4 </i> and <i> TLR9 </i> Gene Polymorphisms with Development of Pulmonary Tuberculosis in Zahedan, Southeastern Iran. Scientific World Journal, The, 2013, 2013, 1-7.	2.1	24
20	Association of FAS and FAS Ligand Genes Polymorphism and Risk of Systemic Lupus Erythematosus. Scientific World Journal, The, 2013, 2013, 1-6.	2.1	24
21	Association of plasma nitric oxide concentration and endothelial nitric oxide synthase T-786C gene polymorphism in coronary artery disease. Pathophysiology, 2012, 19, 157-162.	2.2	23
22	Comparison of Salivary Cortisol and αâ€amylase Levels and Psychological Profiles in Patients with Burning Mouth Syndrome. Special Care in Dentistry, 2017, 37, 120-125.	0.8	23
23	Angiotensin converting enzyme DD genotype not associated with increased risk of coronary artery disease in the Iranian population. Pathophysiology, 2010, 17, 163-167.	2.2	22
24	Biomarkers identified from serum proteomic analysis for the differential diagnosis of systemic lupus erythematosus. Lupus, 2015, 24, 582-587.	1.6	22
25	Association of the osteopontin rs1126616 polymorphism and a higher serum osteopontin level with lupus nephritis. Biomedical Reports, 2016, 4, 355-360.	2.0	22
26	Vascular endothelial growth factor (<i>VEGF</i>)â€634G/C polymorphism was associated with severe preâ€eclampsia and lower serum VEGF level. Journal of Obstetrics and Gynaecology Research, 2015, 41, 1877-1883.	1.3	21
27	The association of the placental MTHFR 3′â€UTR polymorphisms, promoter methylation, and MTHFR expression with preeclampsia. Journal of Cellular Biochemistry, 2018, 119, 1346-1354.	2.6	20
28	Analysis of polymorphisms, promoter methylation, and mRNA expression profile of maternal and placental P53 and P21 genes in preeclamptic and normotensive pregnant women. Journal of Biomedical Science, 2019, 26, 92.	7.0	20
29	Impact of HOTAIR variants on preeclampsia susceptibility based on blood and placenta and in silico analysis. IUBMB Life, 2019, 71, 1367-1381.	3.4	20
30	<i>XRCC1</i> Arg399Gln and Arg194Trp Polymorphisms and Risk of Systemic Lupus Erythematosus in an Iranian Population: A Pilot Study. BioMed Research International, 2014, 2014, 1-5.	1.9	19
31	Effect of Achillea wilhelmsii extract on expression of the human telomerase reverse transcriptase mRNA in the PC3 prostate cancer cell line. Biomedical Reports, 2017, 7, 251-256.	2.0	19
32	The long nonâ€coding RNA H19 rs217727 polymorphism is associated with PE susceptibility. Journal of Cellular Biochemistry, 2018, 119, 5473-5480.	2.6	19
33	Association Between Functional Polymorphisms of DNA Double-Strand Breaks in Repair Genes <i>XRCC5</i> , <i>XRCC6</i> and <i>XRCC7</i> with the Risk of Systemic Lupus Erythematosus in South East Iran. DNA and Cell Biology, 2015, 34, 360-366.	1.9	17
34	Antiproliferative and Antioxidant Effects of Withania coagulans Extract on Benign Prostatic Hyperplasia in Rats. Nephro-Urology Monthly, 2016, 8, e33180.	0.1	17
35	Genetic polymorphisms and haplotypes of the DJ-1 gene promoter associated with the susceptibility to male infertility. Journal of Assisted Reproduction and Genetics, 2017, 34, 1673-1682.	2.5	16
36	The role of TNF- $\hat{l}\pm$ and TLR4 polymorphisms in the placenta of pregnant women complicated by preeclampsia and in silico analysis. International Journal of Biological Macromolecules, 2019, 134, 1205-1215.	7. 5	16

#	Article	IF	Citations
37	Ageâ€dependent association of <i><scp>MDM2</scp></i> promoter polymorphisms and uterine leiomyoma in <scp>S</scp> outhâ€ <scp>E</scp> ast <scp>I</scp> ran: A preliminary report. Journal of Obstetrics and Gynaecology Research, 2015, 41, 729-734.	1.3	15
38	KE and EE Genotypes of ICAM-1 Gene K469E Polymorphism Is Associated with Severe Preeclampsia. Disease Markers, 2014, 2014, 1-5.	1.3	14
39	Association of functional polymorphisms in <i>FAS</i> and <i>FAS</i> \cdot\i> Ligand genes promoter with preâ€eclampsia. Journal of Obstetrics and Gynaecology Research, 2014, 40, 1167-1173.	1.3	14
40	The placental vascular endothelial growth factor polymorphisms and preeclampsia/preeclampsia severity. Clinical and Experimental Hypertension, 2017, 39, 606-611.	1.3	14
41	Genetic polymorphisms of miRNA <i>let7aâ€2</i> and <i>priâ€mirâ€34b/c</i> are associated with an increased risk of papillary thyroid carcinoma and clinical/pathological features. Journal of Cellular Biochemistry, 2019, 120, 8640-8647.	2.6	14
42	The possible role of maternal and placental vitamin D receptor polymorphisms and haplotypes in pathogenesis of preeclampsia. Clinical and Experimental Hypertension, 2020, 42, 171-176.	1.3	14
43	The effects of placental long noncoding RNA H19 polymorphisms and promoter methylation on H19 expression in association with preeclampsia susceptibility. IUBMB Life, 2020, 72, 413-425.	3.4	14
44	Relationships between Dicer 1 polymorphism and expression levels in the etiopathogenesis of preeclampsia. Journal of Cellular Biochemistry, 2018, 119, 5563-5570.	2.6	13
45	The Drosha rs10719 T>C polymorphism is associated with preeclampsia susceptibility. Clinical and Experimental Hypertension, 2018, 40, 440-445.	1.3	13
46	Endothelial nitric oxide synthase gene Glu298Asp polymorphism in patients with coronary artery disease. Annals of Saudi Medicine, 2010, 30, 33-37.	1.1	13
47	Combination Effect of GSTM1, GSTT1 and GSTP1 Polymorphisms and Risk of Systemic Lupus Erythematosus. Iranian Journal of Public Health, 2015, 44, 814-21.	0.5	13
48	Endothelial nitric oxide synthase gene intron4 VNTR polymorphism in patients with coronary artery disease in Iran. Indian Journal of Medical Research, 2006, 124, 683-8.	1.0	13
49	Lack of evidence for constribution of intron4a/b polymorphism of endothelial nitric oxide synthase (NOS3) gene to plasma nitric oxide levels. Acta Cardiologica, 2008, 63, 229-234.	0.9	12
50	Lack of relationship between endothelial nitric oxide synthase gene 4b/a and T-786C polymorphisms with preeclampsia in southeast of Iran. Archives of Gynecology and Obstetrics, 2012, 285, 405-409.	1.7	12
51	The Effect of Experimental Thyroid Dysfunction on Markers of Oxidative Stress in Rat Pancreas. Drug Development Research, 2016, 77, 199-205.	2.9	12
52	Association of the placental VEGF promoter polymorphisms and VEGF mRNA expression with preeclampsia. Clinical and Experimental Hypertension, 2019, 41, 274-279.	1.3	12
53	Association between miRNA-152 polymorphism and risk of preeclampsia susceptibility. Archives of Gynecology and Obstetrics, 2019, 299, 475-480.	1.7	12
54	Renalase rs10887800 polymorphism is associated with severe preâ€eclampsia in southeast Iranian women. Journal of Cellular Biochemistry, 2019, 120, 3277-3285.	2.6	12

#	Article	IF	CITATIONS
55	The effect of GPx-1 rs1050450 and MnSOD rs4880 polymorphisms on PE susceptibility: a case- control study. Molecular Biology Reports, 2019, 46, 6099-6104.	2.3	11
56	The effect of TP53 and P21 gene polymorphisms on papillary thyroid carcinoma susceptibility and clinical/pathological features. IUBMB Life, 2020, 72, 922-930.	3.4	11
57	The effects of the genetic polymorphisms of antioxidant enzymes on susceptibility to papillary thyroid carcinoma. IUBMB Life, 2020, 72, 1045-1053.	3.4	11
58	Long non-coding RNA <i>ANRIL</i> polymorphisms in papillary thyroid cancer and its severity. British Journal of Biomedical Science, 2021, 78, 58-62.	1.3	11
59	Association of XRCC1 Arg399GIn and Tp53 Arg72Pro polymorphisms and increased risk of uterine leiomyoma - A case-control study. Genetics and Molecular Biology, 2015, 38, 444-449.	1.3	10
60	Genetic variants in 3′â€UTRs of MTHFR in the pregnancies complicated with preeclampsia and bioinformatics analysis. Journal of Cellular Biochemistry, 2018, 119, 773-781.	2.6	10
61	Association between selenium, cadmium, and arsenic levels and genetic polymorphisms in DNA repair genes (XRCC5, XRCC6) in gastric cancerous and non-cancerous tissue. Journal of Trace Elements in Medicine and Biology, 2019, 55, 89-95.	3.0	10
62	Lower Total Serum Protein, Albumin and Zinc in Depression in an Iranian Population. Journal of Medical Sciences (Faisalabad, Pakistan), 2008, 8, 587-590.	0.0	10
63	Vitamin D Receptor rs2228570 and rs731236 Polymorphisms are Susceptible Factors for Systemic Lupus Erythematosus. Advanced Biomedical Research, 2019, 8, 48.	0.5	10
64	Endothelial nitric oxide synthase gene Glu298Asp polymorphism in patients with coronary artery disease. Annals of Saudi Medicine, 2010, 30, 33-7.	1.1	10
65	Relationship between Estradiol and Antioxidant Enzymes Activity of Ischemic Stroke. Journal of Biomedicine and Biotechnology, 2009, 2009, 1-5.	3.0	9
66	The effect of miR-146a rs2910164 and miR-149 rs2292832 polymorphisms on preeclampsia susceptibility. Molecular Biology Reports, 2019, 46, 4529-4536.	2.3	9
67	Hypomethylation of the miRNA-34a gene promoter is associated with Severe Preeclampsia. Clinical and Experimental Hypertension, 2019, 41, 118-122.	1.3	9
68	Association of CTLA4 (rs4553808) and PTPN22 (rs2476601) gene polymorphisms with Hashimoto's thyroiditis disease: A case-control study and an In-silico analysis. Meta Gene, 2020, 24, 100693.	0.6	9
69	Effect of mobile phone usage time on total antioxidant capacity of saliva and salivary immunoglobulin a. Iranian Journal of Public Health, 2014, 43, 480-4.	0.5	9
70	Endothelial nitric oxide synthase gene Glu298Asp polymorphism and risk of preeclampsia in South East of Iran. African Journal of Biotechnology, 2011, 10, 10712-10717.	0.6	8
71	Association of L55M and Q192R Polymorphisms of Paraoxonase-1 Gene withÂPreeclampsia. Archives of Medical Research, 2011, 42, 324-328.	3.3	8
72	Adenosine deaminase activity in fertile and infertile men. Andrologia, 2012, 44, 586-589.	2.1	8

#	Article	IF	CITATIONS
73	The effects of p21 gene C98A polymorphism on development of uterine leiomyoma in southeast Iranian women. Tumor Biology, 2016, 37, 12497-12502.	1.8	8
74	The association of the placental CASPASEâ€3 gene polymorphisms and preeclampsia susceptibility and inâ€silico analysis. Journal of Cellular Biochemistry, 2018, 119, 6756-6764.	2.6	8
75	The association of pri-miRNA- 26a1 rs7372209 polymorphism and Preeclampsia susceptibility. Clinical and Experimental Hypertension, 2019, 41, 268-273.	1.3	8
76	Role of MDM2 309T>G (rs2279744) and I/D (rs3730485) polymorphisms and haplotypes in risk of papillary thyroid carcinoma, tumor stage, tumor size, and early onset of tumor: A case control study. Journal of Cellular Physiology, 2019, 234, 12934-12940.	4.1	8
77	Association between Genetic Polymorphisms in microRNA Machinery Genes and Risk of Papillary Thyroid Carcinoma. Pathology and Oncology Research, 2020, 26, 1235-1241.	1.9	8
78	Genetic variants of <scp><i>HOTAIR</i></scp> are associated with susceptibility to recurrent spontaneous abortion: A preliminary case–control study. Journal of Obstetrics and Gynaecology Research, 2021, 47, 3767-3778.	1.3	8
79	Association of polymorphisms and haplotypes in the cytochrome P450 1B1 gene with uterine leiomyoma: A case control study. Biomedical Reports, 2015, 3, 201-206.	2.0	7
80	Association of interleukinâ€1 receptor antagonist VNTR polymorphism and risk of preâ€eclampsia in southeast Iranian population. Journal of Obstetrics and Gynaecology Research, 2016, 42, 142-147.	1.3	7
81	The MDM2 promoter T309G polymorphism was associated with preeclampsia susceptibility. Journal of Assisted Reproduction and Genetics, 2017, 34, 951-956.	2.5	7
82	Association of HOTAIR gene polymorphisms and haplotypes with uterine leiomyoma susceptibility in southeast of Iran. Molecular Biology Reports, 2019, 46, 4271-4277.	2.3	7
83	The effects of DICER1 and DROSHA polymorphisms on susceptibility to recurrent spontaneous abortion. Journal of Clinical Laboratory Analysis, 2020, 34, e23079.	2.1	7
84	The relationships between maternal and placental polymorphisms of miR-196a2 and miRNA-499 genes and preeclampsia. British Journal of Biomedical Science, 2020, 77, 191-195.	1.3	7
85	Association of IL- $1^{\hat{1}^2}$, NLRP3, and COX-2 Gene Polymorphisms with Autoimmune Thyroid Disease Risk and Clinical Features in the Iranian Population. BioMed Research International, 2021, 2021, 1-10.	1.9	7
86	Cyclin D1 G870A polymorphism: Association with uterine leiomyoma risk and in silico analysis. Biomedical Reports, 2017, 6, 237-241.	2.0	6
87	Estrogen receptor alpha Xbal GG genotype was associated with severe preeclampsia. Clinical and Experimental Hypertension, 2017, 39, 220-224.	1.3	6
88	Association between $\langle scp \rangle ER \langle scp \rangle \hat{l} \pm polymorphisms$ and systemic lupus erythematosus: susceptibility and $\langle i \rangle$ in silico $\langle i \rangle$ analysis. International Journal of Rheumatic Diseases, 2018, 21, 214-222.	1.9	6
89	Common Variations in Prothrombotic Genes and Susceptibility to Ischemic Stroke in Young Patients: A Case-Control Study in Southeast Iran. Medicina (Lithuania), 2019, 55, 47.	2.0	6
90	Prooxidant-Antioxidant Balance in Patients with Systemic Lupus Erythematosus and Its Relationship with Clinical and Laboratory Findings. Autoimmune Diseases, 2016, 2016, 1-5.	0.6	5

#	Article	IF	Citations
91	Association of <i><scp>eNOS</scp></i> gene polymorphisms and systemic lupus erythematosus in southeast Iran. International Journal of Rheumatic Diseases, 2016, 19, 606-612.	1.9	5
92	5-Aminolevulinic acid moderates environmental stress-induced bunch wilting and stress markers in date palm. Acta Physiologiae Plantarum, 2018, 40, 1.	2.1	5
93	Effects of the <scp>MTOR</scp> and <scp>AKT1</scp> genes polymorphisms on papillary thyroid cancer development. IUBMB Life, 2020, 72, 2601-2610.	3.4	5
94	Association of ACE I/D and AGTR1 A1166C Gene Polymorphisms and Risk of Uterine Leiomyoma: A Case-Control Study. Asian Pacific Journal of Cancer Prevention, 2019, 20, 2595-2599.	1.2	5
95	The â^'2549 insertion/deletion polymorphism of VEGF gene associated with uterine leiomyoma susceptibility in women from Southeastern Iran. Ginekologia Polska, 2017, 88, 115-119.	0.7	5
96	The effect of the placental DROSHA rs10719 and rs6877842 polymorphisms on PE susceptibility and mRNA expression. Journal of Human Hypertension, 2019, 33, 552-558.	2.2	4
97	The effect of CASP3 rs4647610 and rs4647602 polymorphisms on tumour size and cancer stage in papillary thyroid carcinoma. British Journal of Biomedical Science, 2020, 77, 129-134.	1.3	4
98	Functional miR29a polymorphism is associated with protection against recurrent spontaneous abortion: A case-control study and bioinformatics analysis. Gene Reports, 2021, 23, 101108.	0.8	4
99	Association Between Interleukin 4 Gene Seventy-Base-Pair Variable Number of Tandem Repeats Polymorphism and Uterine Leiomyoma. Gene, Cell and Tissue, 2014, 1, .	0.2	4
100	Association of Polymorphisms in miR146a, an Inflammation-Associated MicroRNA, with the Risk of Idiopathic Recurrent Spontaneous Miscarriage: A Case-Control Study. Disease Markers, 2022, 2022, 1-10.	1.3	4
101	The ID genotype of MDM2 40 bp insertion/deletion polymorphism was associated with lower risk of SLE. Postgraduate Medical Journal, 2017, 93, 758-761.	1.8	3
102	The Impact of TRAIL (C1595T and G1525A) and DR4 (rs20576) Gene Polymorphisms on Systemic Lupus Erythematosus. Biochemical Genetics, 2020, 58, 649-659.	1.7	3
103	Comparison of CAT-21A/T Gene Polymorphism in Women with Preeclampsia and Control Group. Advanced Biomedical Research, 2018, 7, 133.	0.5	3
104	Carriage of 2R allele at VNTR polymorphous site of XRCC5 gene increases risk of multiple sclerosis in an Iranian population. Russian Journal of Genetics, 2017, 53, 147-152.	0.6	2
105	Association of H19 rs3741219 polymorphism with the susceptibility to uterine leiomyomas. Gene Reports, 2020, 19, 100623.	0.8	2
106	Combination effect of cytochrome P450 1A1 gene polymorphisms on uterine leiomyoma: A case-control study. Bosnian Journal of Basic Medical Sciences, 2016, 16, 209-214.	1.0	2
107	Association of FAS A-670G Polymorphism and Risk of Uterine Leiomyoma in a Southeast Iranian Population. Reports of Biochemistry and Molecular Biology, 2016, 5, 51-55.	1.4	2
108	Lack of Association Between IL-1 Receptor Antagonist Gene 86bp VNTR Polymorphism and Leiomyoma. Gene, Cell and Tissue, 2014, 1 , .	0.2	1

#	Article	IF	CITATIONS
109	The Association of Endothelial Nitric Oxide Synthase Gene Polymorphisms and Preeclampsia Susceptibility. Gene, Cell and Tissue, 2015, 2, .	0.2	1
110	TLR8 and TLR9 Polymorphisms and Pulmonary Tuberculosis. Gene, Cell and Tissue, 2015, 2, .	0.2	1
111	Association of Pvull T> C and Xbal A> G Polymorphisms of Estrogen Receptor α Gene with Uterine Leiomyoma: A Case-Control Study. Gene, Cell and Tissue, 2018, In Press, .	0.2	1
112	Evaluation of relationship between methylene tetrahydrofolate reductase gene C677T polymorphism and preeclampsia. Clinical Biochemistry, 2011, 44, S292-S293.	1.9	0
113	The Effect of Renalase rs2576178 and rs10887800 Polymorphisms on Ischemic Stroke Susceptibility in Young Patients (<50 Years): A Case-Control Study and In Silico Analysis. Disease Markers, 2021, 2021, 1-6.	1.3	0
114	Salivary Atopy Biomarkers in Patients with Geographic Tongue. European Journal of General Dentistry, 0, , .	0.4	0
115	Association study of TPH1 (rs1800532) and TPH2 (rs4570625) Polymorphisms in Type 1 Bipolar Disorder in Iran. Gene, Cell and Tissue, 2019, In Press, .	0.2	0