

Mahboobeh Nasiri

List of Publications by Year in descending order

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Version: 2024-02-01

24
papers

102
citations

1684188

5
h-index

1372567

10
g-index

24
all docs

24
docs citations

24
times ranked

136
citing authors

#	ARTICLE	IF	CITATIONS
1	CTLA-4 and IL-6 gene polymorphisms: Risk factors for recurrent pregnancy loss. <i>Human Immunology</i> , 2016, 77, 1271-1274.	2.4	25
2	SIRT1 gene polymorphisms associated with decreased risk of atherosclerotic coronary artery disease. <i>Gene</i> , 2018, 672, 16-20.	2.2	14
3	Association of the rs555172 polymorphism in SENCN long non-coding RNA and atherosclerotic coronary artery disease. <i>Journal of Cardiovascular and Thoracic Research</i> , 2017, 9, 170-174.	0.9	12
4	The IL-6 -634C/G polymorphism: a candidate genetic marker for the prediction of idiopathic recurrent pregnancy loss. <i>International Journal of Reproductive BioMedicine</i> , 2016, 14, 103-108.	0.9	9
5	Association of the +49 A/G Polymorphism of CTLA4 Gene with Idiopathic Recurrent Spontaneous Abortion in Women in Southwest of Iran. <i>Journal of Reproduction and Infertility</i> , 2016, 17, 151-6.	1.0	9
6	Association of ERAP 2 gene variants with risk of pre-eclampsia among Iranian women. <i>International Journal of Gynecology and Obstetrics</i> , 2019, 145, 337-342.	2.3	7
7	Association of rs4618210A>G variant in PLCL2 gene with myocardial infarction: A case-control study in Iran. <i>Journal of Cardiovascular and Thoracic Research</i> , 2020, 12, 303-306.	0.9	5
8	Interferon Regulatory Factor 5 Gene Polymorphisms in Iranian Women with Unexplained Recurrent Pregnancy Loss. <i>Immunological Investigations</i> , 2017, 46, 97-107.	2.0	4
9	Association of the ATG9B gene polymorphisms with coronary artery disease susceptibility: A case-control study. <i>Journal of Cardiovascular and Thoracic Research</i> , 2019, 11, 109-115.	0.9	3
10	Association of TIM-3 (rs1036199) and TIM-4 (rs7700944, rs6882076) gene polymorphisms with susceptibility to systemic lupus erythematosus. <i>Meta Gene</i> , 2020, 25, 100749.	0.6	2
11	G-protein coupled-receptor 65 5'UTR gene polymorphism in the pathogenesis of systemic lupus erythematosus. <i>Rheumatology Research</i> , 2017, 2, 139-143.	0.1	2
12	The IL-6 -634C/G polymorphism: a candidate genetic marker for the prediction of idiopathic recurrent pregnancy loss. <i>International Journal of Reproductive BioMedicine</i> , 2016, 14, 103-8.	0.9	2
13	Glutathione Peroxidase 1 Gene Polymorphism in Nephrolithiasis Patients From South of Iran. <i>Iranian Journal of Kidney Diseases</i> , 2017, 11, 29-35.	0.1	2
14	Association of the functional genetic variants of TOX3 gene with breast cancer in Iran: A case-control study. <i>Gene Reports</i> , 2020, 18, 100511.	0.8	1
15	PCSK9 Gene Polymorphisms Associated With the Risk of Myocardial Infarction in Iranian Patients. <i>International Journal of Basic Science in Medicine</i> , 2021, 6, 57-63.	0.3	1
16	Granulocyte colony-stimulating factor gene rs1042658 variant and susceptibility to idiopathic recurrent pregnancy loss: A case-control study. <i>International Journal of Reproductive BioMedicine</i> , 2018, 16, 35-40.	0.9	1
17	Interleukin-17A Genetic Polymorphisms as a Prognostic Markers for Resistance to Visceral Leishmaniasis in the Iranian Population. <i>Archives of Clinical Infectious Diseases</i> , 2019, In Press, .	0.2	1
18	Granulocyte colony-stimulating factor gene rs1042658 variant and susceptibility to idiopathic recurrent pregnancy loss: A case-control study. <i>International Journal of Reproductive BioMedicine</i> , 2018, 16, 35-40.	0.9	1

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19	in Vitro Effect of Methamphetamine on Proliferation, Differentiation and Apoptosis of Adipose Tissue Stem Cells. Journal of Pharmacy and Pharmaceutical Sciences, 2022, 25, 69-76.	2.1	1
20	Association of M55L and Q192R polymorphisms of paraoxonase 1 gene (PON1) with recurrent pregnancy loss risk: A caseâ€“control study. International Journal of Reproductive BioMedicine, 2021, 19, 559-568.	0.9	0
21	FoxO3a gene down-regulation in pathogenesis of pediatric acute lymphoblastic leukemia. Indian Journal of Medical and Paediatric Oncology, 2019, 40, 381.	0.2	0
22	Down-Regulation of PU.1 Gene in Pediatric Acute Lymphoblastic Leukemia Patients from South of Iran. International Journal of Hematology-Oncology and Stem Cell Research, 0, , .	0.3	0
23	Study of rs2858060 polymorphism in Lnc-Ang362 long non-coding sequence as an informative marker in screening of atherosclerotic coronary artery diseases: pilot study. Medical Journal of Tabriz University of Medical Sciences & Health Services, 2019, 41, 38-43.	0.1	0
24	Down-Regulation of Gene in Pediatric Acute Lymphoblastic Leukemia Patients from South of Iran. International Journal of Hematology-Oncology and Stem Cell Research, 2019, 13, 20-24.	0.3	0