

Sevim KarakaÅ ÆelÄ°k

List of Publications by Year in descending order

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41
papers

431
citations

840728

11
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839512

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44
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44
docs citations

44
times ranked

648
citing authors

#	ARTICLE	IF	CITATIONS
1	Polymorphisms of ACE (I/D) and ACE2 receptor gene (Rs2106809, Rs2285666) are not related to the clinical course of COVID-19: A case study. <i>Journal of Medical Virology</i> , 2021, 93, 5947-5952.	5.0	64
2	The association of FOXO3A gene polymorphisms with serum FOXO3A levels and oxidative stress markers in vitiligo patients. <i>Gene</i> , 2014, 536, 129-134.	2.2	38
3	Glutathione S-Transferase Gene Polymorphisms in Presbycusis. <i>Otology and Neurotology</i> , 2005, 26, 392-397.	1.3	36
4	The Association Between Polymorphic Genotypes of Glutathione S-Transferases and COPD in the Turkish Population. <i>Biochemical Genetics</i> , 2006, 44, 307-319.	1.7	34
5	Interleukin 18 gene polymorphism is a risk factor for multiple sclerosis. <i>Molecular Biology Reports</i> , 2014, 41, 1653-1658.	2.3	20
6	Increased Serum Levels of IL-28 and IL-29 and the Protective Effect of IL28B rs8099917 Polymorphism in Patients with Hashimoto's Thyroiditis. <i>Immunological Investigations</i> , 2016, 45, 668-678.	2.0	19
7	The effect of FOXO gene family variants and global DNA methylation on RRMS disease. <i>Gene</i> , 2020, 726, 144172.	2.2	19
8	A bioinformatic approach to investigating cytokine genes and their receptor variants in relation to COVID-19 progression. <i>International Journal of Immunogenetics</i> , 2021, 48, 211-218.	1.8	17
9	Genetic variants of estrogen beta and leptin receptors may cause gynecomastia in adolescent. <i>Gene</i> , 2014, 541, 101-106.	2.2	15
10	N-Acetyltransferase 2 Phenotype May Be Associated with Susceptibility to Age-Related Cataract. <i>Current Eye Research</i> , 2005, 30, 835-839.	1.5	14
11	IL28B, IL29 and micro-RNA 548 in subacute sclerosing panencephalitis as a rare disease. <i>Gene</i> , 2018, 678, 73-78.	2.2	14
12	The role of GNLY gene polymorphisms in psoriasis pathogenesis. <i>Anais Brasileiros De Dermatologia</i> , 2019, 94, 198-203.	1.1	13
13	PD-1 Gene Polymorphism in Children with Subacute Sclerosing Panencephalitis. <i>Neuropediatrics</i> , 2013, 44, 187-190.	0.6	12
14	Insulin receptor substrate-2 gene polymorphism: is it associated with endometrial cancer?. <i>Gynecological Endocrinology</i> , 2010, 26, 378-382.	1.7	9
15	Higher Levels of Serum TLR2 and TLR4 in Patients with Hashimoto's Thyroiditis. <i>Endocrine, Metabolic and Immune Disorders - Drug Targets</i> , 2020, 20, 118-126.	1.2	9
16	The G1057D polymorphism of insulin receptor substrate-2 associated with gestational diabetes mellitus. <i>Gynecological Endocrinology</i> , 2014, 30, 165-168.	1.7	8
17	Novel SNARE Complex Polymorphisms Associated with Multiple Sclerosis: Signs of Synaptopathy in Multiple Sclerosis. <i>Balkan Medical Journal</i> , 2019, 36, 174-178.	0.8	8
18	GPR30 Gene Polymorphisms Are Associated with Gynecomastia Risk in Adolescents. <i>Hormone Research in Paediatrics</i> , 2015, 83, 177-182.	1.8	7

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19	Functional association of interleukin-18 gene -607 C/A promoter polymorphisms with endometriosis. <i>Fertility and Sterility</i> , 2011, 95, 298-300.	1.0	6
20	Genetic variants of synaptic vesicle and presynaptic plasma membrane proteins in idiopathic generalized epilepsy. <i>Journal of Receptor and Signal Transduction Research</i> , 2014, 34, 38-43.	2.5	6
21	May TLR4 Asp299Gly and IL17 His161Arg polymorphism be associated with progression of primary measles infection to subacute sclerosing panencephalitis?. <i>Gene</i> , 2014, 547, 186-190.	2.2	6
22	Rare Types of Turner Syndrome: Clinical Presentation and Cytogenetics in Five Cases. <i>Laboratory Medicine</i> , 2012, 43, 197-204.	1.2	5
23	IRS-2 G1057D polymorphism in Turkish patients with colorectal cancer. <i>Przegląd Gastroenterologiczny</i> , 2014, 2, 88-92.	0.7	5
24	SMAD2, SMAD3 and TGF- β 2 GENE expressions in women suffering from urge urinary incontinence and pelvic organ prolapse. <i>Molecular Biology Reports</i> , 2021, 48, 1401-1407.	2.3	5
25	Homozygous Ala65Pro Mutation with V89L Polymorphism in SRD5A2 Deficiency. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2016, 8, 218-223.	0.9	5
26	Association between Psoriasis Disease and IFN- γ Gene Polymorphisms. <i>Immunological Investigations</i> , 2022, 51, 1772-1784.	2.0	5
27	Association of G1057D variant of insulin receptor substrate-2 with endometriosis. <i>Fertility and Sterility</i> , 2010, 94, 1622-1626.	1.0	4
28	Association Between IL28B, IL29 Gene Polymorphisms and Clinical Manifestations of Behçet's Disease. <i>Immunological Investigations</i> , 2021, 50, 906-913.	2.0	4
29	Glutathione S-Transferase Z1 (GSTZ1) Gene Polymorphism in Gastric Cancer: A Preliminary Study in a Turkish Population. <i>Laboratory Medicine</i> , 2014, 45, 37-42.	1.2	3
30	Evaluation of serum trail level and DR4 gene variants as biomarkers for vitiligo patients. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2016, 30, e97-e98.	2.4	3
31	In Vitro Effects of Propofol on Cytotoxic, Apoptotic and PI3K-Akt Signaling Pathway Genes on Brain Cancer Cells. <i>Anti-Cancer Agents in Medicinal Chemistry</i> , 2022, 22, 356-361.	1.7	3
32	Association of NOD1, NOD2, PYDC1 and PYDC2 genes with Behçet's disease susceptibility and clinical manifestations. <i>Ophthalmic Genetics</i> , 2021, 42, 691-697.	1.2	3
33	Partial trisomy 4q and partial monosomy 9p in a girl with choanal atresia and various dysmorphic findings. <i>Gene</i> , 2015, 568, 211-214.	2.2	1
34	Lack of association between sirtuin gene variants and endometrial cancer. <i>Meta Gene</i> , 2019, 19, 56-59.	0.6	1
35	Prodynorphin (PDYN) gene polymorphisms in Turkish patients with methamphetamine use disorder, changes in PDYN serum levels in withdrawal and the relationship between PDYN, temperament and depression. <i>Journal of Ethnicity in Substance Abuse</i> , 2020, , 1-16.	0.9	1
36	Association between IL-18 gene polymorphisms and Hashimoto thyroiditis. <i>Molecular Biology Reports</i> , 2021, 48, 6703-6708.	2.3	1

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37	An in-vitro investigation of the effect of perfluorooctane sulphonate on cell lines of embryonic origin. <i>Molecular Biology Reports</i> , 2014, 41, 3755-3759.	2.3	0
38	Serum paraoxonase level and paraoxonase polymorphism in patients with acromegaly. <i>Redox Report</i> , 2016, 21, 281-286.	4.5	0
39	The investigation of BTLA single-nucleotide polymorphisms in patients with Behcet disease in Elazığ province. <i>Turkish Journal of Biochemistry</i> , 2020, 45, 323-327.	0.5	0
40	Glutathione S-Transferase Z1 Gene Polymorphism in Turkish Population. <i>Erciyes Tıp Dergisi</i> , 2012, 34, 160-164.	0.1	0
41	Effects of Interactions among Gene Polymorphisms of the Renin-Angiotensin-Aldosterone System on Hypertension in Turkish People from Southeast Anatolia. <i>Romanian Journal of Laboratory Medicine</i> , 2019, 27, 159-168.	0.2	0