

# Parvaneh Keshavarz

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2918807/publications.pdf>

Version: 2024-02-01

29  
papers

398  
citations

759233

12  
h-index

752698

20  
g-index

30  
all docs

30  
docs citations

30  
times ranked

673  
citing authors

#	ARTICLE	IF	CITATIONS
1	Haplotype-based association study of Opioid Receptor Kappa-type 1 (OPRK1) gene polymorphisms with nicotine dependence among male smokers. <i>British Journal of Biomedical Science</i> , 2021, 78, 151-153.	1.3	2
2	Association Study of Opioid Receptor Delta-Type 1 (OPRD1) Gene Variants with Nicotine Dependence in an Iranian Population. <i>Journal of Molecular Neuroscience</i> , 2021, 71, 1301-1305.	2.3	0
3	Haplotype-Based Association and In Silico Studies of OPRM1 Gene Variants with Susceptibility to Opioid Dependence Among Addicted Iranians Undergoing Methadone Treatment. <i>Journal of Molecular Neuroscience</i> , 2020, 70, 504-513.	2.3	4
4	Single locus and haplotype association of ENPP1 gene variants with the development of retinopathy among type 2 diabetic patients. <i>International Ophthalmology</i> , 2020, 40, 639-647.	1.4	5
5	Evaluation of beta-thalassemia in the fetus through cffDNA with multiple polymorphisms as a haplotype in the beta-globin gene. <i>Transfusion Clinique Et Biologique</i> , 2020, 27, 243-252.	0.4	1
6	Significant Destructive Interaction of BDNF Val>Met Polymorphism with Stroke Severity and Family History of Dementia for Cognitive Impairments. <i>Acta Neurologica Taiwanica</i> , 2020, 29(3), 67-78.	0.3	1
7	Single and multi-locus association study of TCF7L2 gene variants with susceptibility to type 2 diabetes mellitus in an Iranian population. <i>Gene</i> , 2019, 696, 88-94.	2.2	14
8	A De Novo Deletion of Chromosome 18p with Persistent Limb Tremor and Difficulty Speaking. <i>Caspian Journal of Neurological Sciences</i> , 2019, 5, 41-47.	0.2	0
9	Comparing Oprm1 Gene Promoter Methylation in the Lymphocytes of Male Rats Addicted to Nicotine, Morphine, Methadone, and Buprenorphine. <i>Caspian Journal of Neurological Sciences</i> , 2019, 5, 168-174.	0.2	0
10	Association and in silico studies of ENPP1 gene variants with type 2 diabetes mellitus in a Northern Iranian population. <i>Gene</i> , 2018, 675, 225-232.	2.2	4
11	Assessment of Gene Expression Following Vitrification of 2-cell and Blastocyst Mouse Embryos. <i>Avicenna Journal of Medical Biotechnology</i> , 2018, 10, 120-122.	0.3	0
12	Brain-Derived Neurotrophic Factor (BDNF) Val66met (rs6265) Polymorphism Associated with Global and Multi-Domain Cognitive Impairment in Ischemic Stroke Patients. <i>Activitas Nervosa Superior</i> , 2017, 59, 28-36.	0.4	4
13	Association of OPRD1 Gene Variants with Opioid Dependence in Addicted Male Individuals Undergoing Methadone Treatment in the North of Iran. <i>Journal of Psychoactive Drugs</i> , 2017, 49, 242-251.	1.7	14
14	Association of OPRK1 gene polymorphisms with opioid dependence in addicted men undergoing methadone treatment in an Iranian population. <i>Journal of Addictive Diseases</i> , 2017, 36, 227-235.	1.3	18
15	Brain-derived neurotrophic factor (BDNF) Val66Met polymorphism and post-stroke dementia: a hospital-based study from northern Iran. <i>Neurological Sciences</i> , 2016, 37, 935-942.	1.9	19
16	Association of BDNF G196A Gene Polymorphism with Ischemic Stroke Occurrence and its 6-Month Outcome in an Iranian Population. <i>Topics in Stroke Rehabilitation</i> , 2016, 23, 254-260.	1.9	15
17	Lack of genetic susceptibility of KCNJ11 E23K polymorphism with risk of type 2 diabetes in an Iranian population. <i>Endocrine Research</i> , 2014, 39, 120-125.	1.2	21
18	Lack of association of genetic variation in chromosome region 15q14-22.1 with type 2 diabetes in a Japanese population. <i>BMC Medical Genetics</i> , 2008, 9, 22.	2.1	1

#	ARTICLE	IF	CITATIONS
19	The association of genetic variants in Kr <sup>1/4</sup> ppelâ€like factor <sup>11</sup> and Type <sup>2</sup> diabetes in the Japanese population. <i>Diabetic Medicine</i> , 2008, 25, 19-26.	2.3	14
20	Single nucleotide polymorphisms in genes encoding LKB1 (STK11), TORC2 (CRTC2) and AMPK $\hat{1}\pm$ 2-subunit (PRKAA2) and risk of type 2 diabetes. <i>Molecular Genetics and Metabolism</i> , 2008, 93, 200-209.	1.1	36
21	Transgenic expression of a mutated cyclin-dependent kinase 4 (CDK4/R24C) in pancreatic $\hat{1}$ 2-cells prevents progression of diabetes in db/db mice. <i>Diabetes Research and Clinical Practice</i> , 2008, 82, 33-41.	2.8	17
22	Evaluation of sample size effect on the identification of haplotype blocks. <i>BMC Bioinformatics</i> , 2007, 8, 200.	2.6	5
23	Genetic association of single nucleotide polymorphisms in endonuclease G-like 1 gene with type 2 diabetes in a Japanese population. <i>Diabetologia</i> , 2007, 50, 1218-1227.	6.3	11
24	SNPs in the KCNJ11-ABCC8 gene locus are associated with type 2 diabetes and blood pressure levels in the Japanese population. <i>Journal of Human Genetics</i> , 2007, 52, 781-793.	2.3	84
25	Association of single-nucleotide polymorphisms in the suppressor of cytokine signaling 2 (SOCS2) gene with type 2 diabetes in the Japanese. <i>Genomics</i> , 2006, 87, 446-458.	2.9	35
26	No evidence for association of the ENPP1 (PC-1) K121Q variant with risk of type 2 diabetes in a Japanese population. <i>Journal of Human Genetics</i> , 2006, 51, 559-566.	2.3	53
27	Association study on chromosome 20q11.21-13.13 locus and its contribution to type 2 diabetes susceptibility in Japanese. <i>Human Genetics</i> , 2006, 120, 527-542.	3.8	19
28	The E23K Polymorphism of KCNJ11 and Diabetic Retinopathy in Northern Iran. , 0, 79, .		1
29	A study on methylation of two CpG islands of MAOA gene promoter among opium-addicted males undergoing methadone treatment. <i>Nucleosides, Nucleotides and Nucleic Acids</i> , 0, , 1-10.	1.1	0