

# Seyed Mohammad Akrami

## List of Publications by Year in descending order

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Version: 2024-02-01

31  
papers

529  
citations

687363

13  
h-index

677142

22  
g-index

33  
all docs

33  
docs citations

33  
times ranked

923  
citing authors

#	ARTICLE	IF	CITATIONS
1	Normal and tumour cervical cells respond differently to vaginal lactobacilli, independent of pH and lactate. <i>Journal of Medical Microbiology</i> , 2013, 62, 1065-1072.	1.8	100
2	IS CONSANGUINEOUS MARRIAGE RELIGIOUSLY ENCOURAGED? ISLAMIC AND IRANIAN CONSIDERATIONS. <i>Journal of Biosocial Science</i> , 2007, 39, 313-316.	1.2	58
3	Inflammation, a significant player of Ataxiaâ€“Telangiectasia pathogenesis?. <i>Inflammation Research</i> , 2018, 67, 559-570.	4.0	37
4	Ataxia telangiectasia syndrome: moonlighting ATM. <i>Expert Review of Clinical Immunology</i> , 2017, 13, 1155-1172.	3.0	36
5	Is There a Significant Trend in Prevalence of Consanguineous Marriage in Tehran? A Review of Three Generations. <i>Journal of Genetic Counseling</i> , 2009, 18, 82-86.	1.6	31
6	Mercury specifically induces LINE-1 activity in a human neuroblastoma cell line. <i>Mutation Research - Genetic Toxicology and Environmental Mutagenesis</i> , 2014, 759, 9-20.	1.7	30
7	In vitro Therapeutic Effects of Low Level Laser at mRNA Level on the Release of Skin Growth Factors from Fibroblasts in Diabetic Mice. <i>Avicenna Journal of Medical Biotechnology</i> , 2014, 6, 113-8.	0.3	29
8	Genetic Counseling in Southern Iran: Consanguinity and Reason for Referral. <i>Journal of Genetic Counseling</i> , 2008, 17, 472-479.	1.6	25
9	The Effect of <i>Lactobacillus crispatus</i> and <i>Lactobacillus rhamnosus</i> Culture Supernatants on Expression of Autophagy Genes and HPV E6 and E7 Oncogenes in The HeLa Cell Line. <i>Cell Journal</i> , 2016, 17, 601-7.	0.2	25
10	Effect of Class Switch Recombination Defect on the Phenotype of Ataxia-Telangiectasia Patients. <i>Immunological Investigations</i> , 2021, 50, 201-215.	2.0	22
11	Exposure of hepatocellular carcinoma cells to low-level As <sub>2</sub> O <sub>3</sub> causes an extra toxicity pathway via L1 retrotransposition induction. <i>Toxicology Letters</i> , 2014, 229, 111-117.	0.8	20
12	The effect of homozygous deletion of the BBOX1 and Fibrin genes on carnitine level and acyl carnitine profile. <i>BMC Medical Genetics</i> , 2014, 15, 75.	2.1	17
13	Effect of Copper Sulfate on Expression of Endogenous L1 Retrotransposons in HepG2 Cells (Hepatocellular Carcinoma). <i>Biological Trace Element Research</i> , 2015, 165, 131-134.	3.5	14
14	Overexpression of <i>HPRT1</i> is associated with poor prognosis in head and neck squamous cell carcinoma. <i>FEBS Open Bio</i> , 2021, 11, 2525-2540.	2.3	14
15	Screening for exonic copy number mutations at MSH2 and MLH1 by MAPH. <i>Familial Cancer</i> , 2005, 4, 145-149.	1.9	10
16	Evaluating the Extent of LINE-1 Mobility Following Exposure to Heavy Metals in HepG2 Cells. <i>Biological Trace Element Research</i> , 2014, 160, 143-151.	3.5	10
17	Effect of heavy metals on silencing of engineered long interspersed element-1 retrotransposon in nondividing neuroblastoma cell line. <i>Iranian Biomedical Journal</i> , 2013, 17, 171-8.	0.7	8
18	2q34-qter duplication and 4q34.2-qter deletion in a patient with developmental delay. <i>European Journal of Medical Genetics</i> , 2012, 55, 203-210.	1.3	7

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19	Deregulation of miR-1, miR486, and let-7a in cytogenetically normal acute myeloid leukemia: association with NPM1 and FLT3 mutation and clinical characteristics. <i>Tumor Biology</i> , 2016, 37, 4841-4847.	1.8	7
20	Genetics of consanguineous marriage: Impact and importance of counseling. <i>Journal of Pediatric Genetics</i> , 2015, 01, 217-220.	0.7	6
21	The spectrum of <i>ATM</i> gene mutations in Iranian patients with ataxia-telangiectasia. <i>Pediatric Allergy and Immunology</i> , 2021, 32, 1316-1326.	2.6	5
22	Genetics of hereditary nonpolyposis colorectal cancer. <i>Archives of Iranian Medicine</i> , 2006, 9, 381-9.	0.6	5
23	Retrotransposons and pediatric genetic disorders: Importance and implications. <i>Journal of Pediatric Genetics</i> , 2015, 03, 009-016.	0.7	3
24	Mitochondrial Variants in Pompe Disease: A Comparison between Classic and Non-Classic Forms. <i>Cell Journal</i> , 2018, 20, 333-339.	0.2	3
25	Mitochondrial Copy Number and D-Loop Variants in Pompe Patients. <i>Cell Journal</i> , 2016, 18, 405-15.	0.2	3
26	Novel trends in genetics: transposable elements and their application in medicine. <i>Archives of Iranian Medicine</i> , 2014, 17, 702-12.	0.6	3
27	Molecular changes in obese and depressive patients are similar to neurodegenerative disorders. <i>Iranian Journal of Neurology</i> , 2017, 16, 192-200.	0.5	1
28	IS CONSANGUINEOUS MARRIAGE RELIGIOUSLY ENCOURAGED? A REPLY TO SAADAT. <i>Journal of Biosocial Science</i> , 2008, 40, 155-155.	1.2	0
29	Genetic Risk Variants for Class Switching Recombination Defects in Ataxia-Telangiectasia Patients. <i>Journal of Clinical Immunology</i> , 2022, 42, 72-84.	3.8	0
30	Review of Prenatal Aneuploidy Screening Uptake Rate and Trends in Iran, and Developed Countries. <i>Journal of Human Genetics and Genomics</i> , 2021, 3, .	0.0	0
31	Plasma 5-miRNA as Biomarkers for Identifying Prostate Cancer Patients. <i>Iranian Journal of Public Health</i> , 2019, 48, 1743-1745.	0.5	0