Reza Mirfakhraie

List of Publications by Year
in descending order

133 papers	1,401 citations	393982 19 h-index	476904 29 g-index
138 all docs	138 docs citations	138 times ranked	2291 citing authors

#	Article	IF	Citations
1	Circular RNAs: Novel Biomarkers in Spermatogenesis Defects and Male Infertility. Reproductive Sciences, 2023, 30, 62-71.	1.1	5
2	Emerging Roles of Long Non-coding RNAs in Uterine Leiomyoma Pathogenesis: a Review. Reproductive Sciences, 2022, 29, 1086-1101.	1.1	9
3	Treatment Failure in Acute Myeloid Leukemia: Focus on the Role of Extracellular Vesicles. Leukemia Research, 2022, 112, 106751.	0.4	8
4	Identification of a novel de novo mutation in the CTNNB1 gene in an Iranian patient with intellectual disability. Neurological Sciences, 2022, 43, 2859.	0.9	4
5	The role of FOXC1/FOXCUT/DANCR axis in triple negative breast cancer: a bioinformatics and experimental approach. Molecular Biology Reports, 2022, 49, 2821-2829.	1.0	3
6	Highlighting the interaction between immunomodulatory properties of mesenchymal stem cells and signaling pathways contribute to Graft Versus Host Disease management. Transplant Immunology, 2022, 71, 101524.	0.6	1
7	Autophagy ATG16L1Ârs2241880 impacts the colorectal cancer risk: A caseâ€control study. Journal of Clinical Laboratory Analysis, 2022, 36, e24169.	0.9	5
8	Reduced expression of <scp> CFAP44 </scp> and <scp> CFAP44â€AS1 </scp> may affect sperm motility and morphology. Andrologia, 2022, 54, e14447.	1.0	4
9	Analysis of Cytogenetic Abnormalities in Iranian Patients with Syndromic Autism Spectrum Disorder: A Case Series.. Iranian Journal of Child Neurology, 2022, 16, 117-128.	0.2	0
10	Long non-coding RNA MIR4435-2HG: a key molecule in progression of cancer and non-cancerous disorders. Cancer Cell International, 2022, 22, .	1.8	11
11	A novelSRD5A2mutation in an Iranian family with sex development disorder. Andrologia, 2021, 53, e13847.	1.0	1
12	Novel long noncoding RNAs upregulation may have synergistic effects on theÂCYP24A1 and PFDN4 biomarker role in human colorectal cancer. Journal of Cellular Physiology, 2021, 236, 2051-2057.	2.0	13
13	Study of Linc00574 Regulatory Effect on the TCTE3 Expression in Sperm Motility. Reproductive Sciences, 2021, 28, 159-165.	1.1	10
14	CYP24A1 expression analysis in uterine leiomyoma regarding MED12 mutation profile. Archives of Gynecology and Obstetrics, 2021, 303, 787-792.	0.8	7
15	Dysregulation of vitamin D synthesis pathway genes in colorectal cancer: A caseâ€control study. Journal of Clinical Laboratory Analysis, 2021, 35, e23617.	0.9	5
16	Contribution of long noncoding RNA HOTAIR variants to preeclampsia susceptibility in Iranian women. Hypertension in Pregnancy, 2021, 40, 29-35.	0.5	3
17	CREB-binding protein (CREBBP) and preeclampsia: a new promising target gene. Molecular Biology Reports, 2021, 48, 2117-2122.	1.0	3
18	DACT1 variants and colorectal cancer. British Journal of Biomedical Science, 2021, 78, 221-224.	1.2	2

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19	Long noncoding RNA LINC00978 acts as a potential diagnostic biomarker in patients with colorectal cancer. Experimental and Molecular Pathology, 2021, 122, 104666.	0.9	12
20	The effect of granulocyte colonyâ€stimulating factor dose and administration interval after allogeneic hematopoietic cell transplantation on early engraftment of neutrophil and platelet. Journal of Clinical Laboratory Analysis, 2021, 35, e24060.	0.9	1
21	Coding and Non-Coding RNAs, as Male Fertility and Infertility Biomarkers. International Journal of Fertility & Sterility, 2021, 15, 158-166.	0.2	0
22	Overexpression of GABRP Gene in Triple Negative Breast Cancer: Molecular Mechanisms and Interpretation. International Journal of Cancer Management, 2021, 14, .	0.2	2
23	Differential expression of <i>Hsa-miR-517a/b</i> in placental tissue may contribute to the pathogenesis of preeclampsia. Journal of the Turkish German Gynecology Association, 2021, 22, 273-278.	0.2	0
24	Ectopic expression of CYP24A1 circular RNA hsa_circ_0060927 in uterine leiomyomas. Journal of Clinical Laboratory Analysis, 2020, 34, e23114.	0.9	4
25	A candidate intronic CYP24A1 gene variant affects the risk of colorectal cancer. Biomarkers in Medicine, 2020, 14, 23-29.	0.6	5
26	Mutational screening of RTK-BRAF genes in de novo adult acute myeloid leukemia. Gene Reports, 2020, 21, 100904.	0.4	0
27	RARâ€related orphan receptor A: One gene with multiple functions related to migraine. CNS Neuroscience and Therapeutics, 2020, 26, 1315-1321.	1.9	9
28	Investigating the regulatory function of the ANO1-AS2 on the ANO1 gene in infertile men with asthenozoospermia and terato-asthenozoospermia. Experimental and Molecular Pathology, 2020, 117, 104528.	0.9	8
29	A homozygous missense mutation ofWFS1gene causes Wolfram's syndrome without hearing loss in an Iranian family (a report of clinical heterogeneity). Journal of Clinical Laboratory Analysis, 2020, 34, e23358.	0.9	5
30	Could CYP24A1 promoter methylation status affect the gene expression in the colorectal cancer patients?. Meta Gene, 2020, 24, 100656.	0.3	4
31	The novel homozygous p.Asn197_Ser201del mutation in BTD gene is associated with profound biotinidase deficiency in an Iranian consanguineous family. Molecular Biology Reports, 2020, 47, 4021-4027.	1.0	1
32	Association of lncRNAâ€'p53 regulatory network (lincRNAâ€'p21, lincRNAâ€'ROR and MALAT1) and p53 with the clinicopathological features of colorectal primary lesions and tumors. Oncology Letters, 2020, 19, 3937-3949.	0.8	14
33	Importance of CNOT8 Deadenylase Subunit in DNA Damage Responses Following Ionizing Radiation (IR). Reports of Biochemistry and Molecular Biology, 2020, 9, 163-170.	0.5	0
34	Investigating the relationship between ccfDNA concentration, its integrity, and some individual factors in an Iranian population. Human Antibodies, 2020, 28, 319-326.	0.6	0
35	Effects of Quinacrine on Expression of Hippo signaling Pathway Components (LATS1, LATS2, and YAP) in Human Breast Cancer Stem Cells. Asian Pacific Journal of Cancer Prevention, 2020, 21, 3171-3176.	0.5	0
36	Effects of Quinacrine on Expression of Hippo signaling Pathway Components (LATS1, LATS2, and YAP) in Human Breast Cancer Stem Cells. Asian Pacific Journal of Cancer Prevention, 2020, 21, 3171-3176.	0.5	3

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37	LncRNA SRA1 may play a role in the uterine leiomyoma tumor growth regarding the MED12 mutation pattern . International Journal of Women's Health, 2019, Volume 11, 495-500.	1.1	8
38	The Differential DNA Hypermethylation Patterns of microRNA-137 and microRNA-342 Locus in Early Colorectal Lesions and Tumours. Biomolecules, 2019, 9, 519.	1.8	9
39	The value of the plasma circulating cell-free DNA concentration and integrity index as a clinical tool for prostate cancer diagnosis: a prospective caseâ€“control cohort study in an Iranian population . Cancer Management and Research, 2019, Volume 11, 4549-4556.	0.9	11
40	Association of telomere length with chronic exposure to ionizing radiation among inhabitants of natural high background radiation areas of Ramsar, Iran. International Journal of Radiation Biology, 2019, 95, 1113-1121.	1.0	11
41	ANRIL and ANRASSF1 long noncoding RNAs are upregulated in gastric cancer. Journal of Cellular Biochemistry, 2019, 120, 12544-12548.	1.2	14
42	Expression analysis of selected miRâ€206 targets from the transforming growth factorâ€Î² signaling pathway in breast cancer. Journal of Cellular Biochemistry, 2019, 120, 13545-13553.	1.2	2
43	Altered methylation and expression patterns of genes regulating placental nitric oxide pathway in patients with severe preeclampsia. Human Antibodies, 2019, 27, 117-124.	0.6	2
44	An intron variant in the FLT1 gene increases the risk of preeclampsia in Iranian women. Clinical and Experimental Hypertension, 2019, 41, 697-701.	0.5	8
45	miRâ€30a promoter variation contributes to the increased risk of colorectal cancer in an Iranian population. Journal of Cellular Biochemistry, 2019, 120, 7734-7740.	1.2	9
46	Association of Increased Levels of lncRNA H19 in PBMCs with Risk of Coronary Artery Disease. Cell Journal, 2019, 20, 564-568.	0.2	41
47	A Survey of the Common Mutations and IVS8-Tn Polymorphism of Cystic Fibrosis Transmembrane Conductance Regulator Gene in Infertile Men with Nonobstructive Azoospermia and CBAVD in Iranian Population. Iranian Biomedical Journal, 2019, 23, 92-98.	0.4	4
48	Potential using of microRNA-34A in combination with paclitaxel in colorectal cancer cells. Journal of Cancer Research and Therapeutics, 2019, 15, 32.	0.3	10
49	Investigation of Genes Expression in Acute Myeloid Leukemia. Reports of Biochemistry and Molecular Biology, 2019, 7, 136-141.	0.5	10
50	Exon 1 Mutational Screening in Iranian Patients with Uterine Leiomyoma. Reports of Biochemistry and Molecular Biology, 2019, 8, 21-24.	0.5	1
51	Application of Multiplex Ligation-Dependent Probe Amplification in Determining the Copy Number Alterations of Gene Family Members in Invasive Ductal Breast Carcinoma. Reports of Biochemistry and Molecular Biology, 2019, 8, 91-101.	0.5	3
52	A Survey of the Common Mutations and IVS8-Tn Polymorphism of Cystic Fibrosis Transmembrane Conductance Regulator Gene in Infertile Men with Nonobstructive Azoospermia and CBAVD in Iranian Population. Iranian Biomedical Journal, 2019, 23, 92-8.	0.4	4
53	Evaluation of MALAT1 promoter DNA methylation patterns in early colorectal lesions and tumors. Gastroenterology and Hepatology From Bed To Bench, 2019, 12, S58-S65.	0.6	1
54	Expression analysis of AFAP1-AS1 and AFAP1 in breast cancer. Cancer Biomarkers, 2018, 22, 49-54.	0.8	19

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55	Association analysis of the GABRB3 promoter variant and susceptibility to autism spectrum disorder. Basal Ganglia, 2018, 11, 4-7.	0.3	3
56	Investigation of piwi-interacting RNA pathway genes role in idiopathic non-obstructive azoospermia. Scientific Reports, 2018, 8, 142.	1.6	23
57	Association study of FOXP3 gene and the risk of 0020 pre-eclampsia. Clinical and Experimental Hypertension, 2018, 40, 613-616.	0.5	10
58	Expression analysis of CBR3-AS1 and androgen receptor genes in breast cancer. Meta Gene, 2018, 17, 82-87.	0.3	2
59	Premutations of FMR1 CGG repeats are not related to idiopathic premature ovarian failure in Iranian patients: A case control study. Gene, 2018, 676, 189-194.	1.0	7
60	Neuropilin-1 expression is associated with lymph node metastasis in breast cancer tissues. Cancer Management and Research, 2018, Volume 10, 1969-1974.	0.9	19
61	Idiopathic Premature Ovarian Failure and its association to the abnormal longitudinal changes of telomere length in a population of Iranian Infertile Women: A pilot study. Meta Gene, 2018, 18, 58-61.	0.3	4
62	The rs4846049 polymorphism in the 3’UTR region of the MTHFR gene increases the migraine susceptibility in an Iranian population. Journal of Pain Research, 2018, Volume 11, 145-149.	0.8	11
63	ACE gene rs4343 polymorphism elevates the risk of preeclampsia in pregnant women. Journal of Human Hypertension, 2018, 32, 825-830.	1.0	15
64	Association between Long Noncoding RNA ANRIL Expression Variants and Susceptibility to Coronary Artery Disease. International Journal of Molecular and Cellular Medicine, 2018, 7, 1-7.	1.1	25
65	An Association Study between Longitudinal Changes of Leukocyte Telomere and the Risk of Azoospermia in a Population of Iranian Infertile Men. Iranian Biomedical Journal, 2018, 22, 231-236.	0.4	6
66	MiR-206 Target Prediction in Breast Cancer Subtypes by Bioinformatics Tools. International Journal of Cancer Management, 2018, 11, .	0.2	1
67	An Association Study of rs2829803 Polymorphism in miRNA-155 with Multiple Sclerosis in Patients Referred to Farshchian Hospital of Hamadan City, (Iran). Majallah-i DÄnishgÄh-i Ì'UlÅ«m-i PizishkÄ«-i Qum, 2018, 12, 28-34.	0.2	0
68	An Association Study between Longitudinal Changes of Leukocyte Telomere and the Risk of Azoospermia in a Population of Iranian Infertile Men. Iranian Biomedical Journal, 2018, 22, 231-6.	0.4	1
69	Single nucleotide polymorphisms in the FOXP3 gene are associated with increased risk of relapsing-remitting multiple sclerosis. Human Antibodies, 2017, 24, 85-90.	0.6	17
70	Association of Î²-Secretase Functional Polymorphism with Risk of Schizophrenia. Genetic Testing and Molecular Biomarkers, 2017, 21, 248-251.	0.3	3
71	SOCS gene family expression profile in the blood of multiple sclerosis patients. Journal of the Neurological Sciences, 2017, 375, 481-485.	0.3	22
72	Angiotensinâ€converting enzyme gene rs4343 polymorphism increases susceptibility to migraine. CNS Neuroscience and Therapeutics, 2017, 23, 698-699.	1.9	8

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73	The role of parental microRNA alleles in recurrent pregnancy loss: an association study. Reproductive BioMedicine Online, 2017, 34, 325-330.	1.1	20
74	SN38-PEG-PLGA-verapamil nanoparticles inhibit proliferation and downregulate drug transporter ABCG2 gene expression in colorectal cancer cells. Progress in Biomaterials, 2017, 6, 137-145.	1.8	16
75	Expression analysis of long non-coding A T B and its putative target in breast cancer. Breast Disease, 2017, 37, 11-20.	0.4	20
76	The association between FOXP3 gene variations and autism: True or false positive?. Gene, 2017, 635, 1-2.	1.0	0
77	Association of SRD5A2 gene mutations with risk of hypospadias in the Iranian population. Journal of Endocrinological Investigation, 2017, 40, 391-396.	1.8	11
78	Correlation between MTHFR Genotype Polymorphisms and Different Types of Migraine Headache in Iranian Patients: A Case Control Study. Journal of Neurology and Neuroscience, 2017, 8, .	0.4	0
79	Genetic variations in UGT2B28, UGT2B17, UGT2B15 genes and the risk of prostate cancer: A case-control study. Gene, 2017, 634, 47-52.	1.0	8
80	A bioinformatics approach for identification of miR-100 targets implicated in breast cancer. Cellular and Molecular Biology, 2017, 63, 99-105.	0.3	9
81	Expression Study and Clinical Correlations of MYC and CCAT2 in Breast Cancer Patients. Iranian Biomedical Journal, 2017, 21, 303-311.	0.4	22
82	Mutational Analysis of FLT3 Internal Tandem Duplication and D835 in De novo Adult Acute Myeloid Leukemia. Journal of Advances in Medicine and Medical Research, 2017, 24, 1-9.	0.1	1
83	TTY2 genes deletions as genetic risk factor of male infertility. Cellular and Molecular Biology, 2017, 63, 57.	0.3	3
84	rs508485 Polymorphism Is Associated with Non-obstructive Azoospermia in Iranian Patients. Reports of Biochemistry and Molecular Biology, 2017, 5, 108-111.	0.5	9
85	Evaluation of Placental mir-155-5p and Long Non-coding RNA sONE Expression in Patients with Severe Pre-eclampsia. International Journal of Molecular and Cellular Medicine, 2017, 6, 22-30.	1.1	3
86	Evaluation of promoter methylation status of MLH1 gene in Iranian patients with colorectal tumors and adenoma polyps. Gastroenterology and Hepatology From Bed To Bench, 2017, 10, S117-S121.	0.6	1
87	Association of CpG-SNP and 3'UTR-SNP of WFS1 with the Risk of Type 2 Diabetes Mellitus in an Iranian Population. International Journal of Molecular and Cellular Medicine, 2017, 6, 197-203.	1.1	3
88	Combination of cold atmospheric plasma and iron nanoparticles in breast cancer: gene expression and apoptosis study. OncoTargets and Therapy, 2016, Volume 9, 5911-5917.	1.0	44
89	Glutamate receptor, metabotropic 7 (GRM7) gene variations and susceptibility to autism: A caseâ€“control study. Autism Research, 2016, 9, 1161-1168.	2.1	57
90	Genetic variants and expression study of FOXP3 gene in acute coronary syndrome in Iranian patients. Cell Biochemistry and Function, 2016, 34, 158-162.	1.4	8

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91	TRAIL gene expression analysis in multiple sclerosis patients. Human Antibodies, 2016, 24, 33-38.	0.6	7
92	RAR-related orphan receptor A (RORA): A new susceptibility gene for multiple sclerosis. Journal of the Neurological Sciences, 2016, 369, 259-262.	0.3	38
93	Comparative expression analysis of hypoxiaâ€inducible factorâ€alpha and its natural occurring antisense in breast cancer tissues and adjacent noncancerous tissues. Cell Biochemistry and Function, 2016, 34, 572-578.	1.4	29
94	<scp>Docetaxel</scp>â€“Chitosan nanoparticles for breast cancer treatment: cell viability and gene expression study. Chemical Biology and Drug Design, 2016, 88, 850-858.	1.5	32
95	The pre-mir-27a variant rs895819 may contribute to type 2 diabetes mellitus susceptibility in an Iranian cohort. Journal of Endocrinological Investigation, 2016, 39, 1187-1193.	1.8	25
96	A Bioinformatics Approach to the Identification of Variants Associated with Type 1 and Type 2 Diabetes Mellitus that Reside in Functionally Validated miRNAs Binding Sites. Biochemical Genetics, 2016, 54, 211-221.	0.8	13
97	Analysis of the androgen receptor CAG repeats length in Iranian patients with idiopathic non-obstructive azoospermia. Asian Pacific Journal of Reproduction, 2016, 5, 71-74.	0.2	2
98	MiRNA-Related Polymorphisms in miR-146a and TCF21 Are Associated with Increased Susceptibility to Coronary Artery Disease in an Iranian Population. Genetic Testing and Molecular Biomarkers, 2016, 20, 241-248.	0.3	38
99	Expression analysis of four long noncoding RNAs in breast cancer. Tumor Biology, 2016, 37, 2933-2940.	0.8	104
100	The study of MED12 gene mutations in uterine leiomyomas from Iranian patients. Tumor Biology, 2016, 37, 1567-1571.	0.8	19
101	MIR17HG Gene Polymorphism and the Risk of Recurrent Spontaneous Abortion. Gene, Cell and Tissue, 2016, 3, .	0.2	2
102	Lactobacilli Differentially Modulate mTOR and Wnt/ Î²-Catenin Pathways in Different Cancer Cell Lines. Iranian Journal of Cancer Prevention, 2016, In Press, e5369.	0.7	28
103	Lactobacilli Modulate Hypoxia-Inducible Factor (HIF)-1 Regulatory Pathway in Triple Negative Breast Cancer Cell Line. Cell Journal, 2016, 18, 237-244.	0.2	13
104	MAP3K1 May be a Promising Susceptibility Gene for Type 2 Diabetes Mellitus in an Iranian Population. International Journal of Molecular and Cellular Medicine, 2016, 5, 134-140.	1.1	3
105	Regulation of BAX/BCL2 gene expression in breast cancer cells by docetaxel-loaded human serum albumin nanoparticles. Medical Oncology, 2015, 32, 208.	1.2	21
106	Bioinformatics prioritization of SNPs perturbing microRNA regulation of hematological malignancy-implicated genes. Genomics, 2015, 106, 360-366.	1.3	24
107	Novel missense mutation in the GALNS gene in an affected patient with severe form of mucopolysaccharidosis type IVA. Clinica Chimica Acta, 2015, 450, 121-124.	0.5	4
108	Mutational screening of theNR5A1in azoospermia. Andrologia, 2015, 47, 395-401.	1.0	15

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109	Aberrant Wnt/Î²-Catenin Signaling Pathway in Testis of Azoospermic Menâ€‹. Advanced Pharmaceutical Bulletin, 2015, 5, 373-377.	0.6	7
110	Promyelocytic Leukemia (PML) Gene Mutations may not Contribute to Gastric Adenocarcinoma Development. Asian Pacific Journal of Cancer Prevention, 2015, 16, 3523-3525.	0.5	3
111	RIT2 , a susceptibility gene for Parkinson's disease in Iranian population. Neurobiology of Aging, 2014, 35, e27-e28.	1.5	23
112	An Iranian family with azoospermia and premature ovarian insufficiency segregating NR5A1 mutation. Climacteric, 2014, 17, 301-303.	1.1	6
113	Thiamine-responsive megaloblastic anemia syndrome with Ebstein anomaly: a case report. European Journal of Pediatrics, 2014, 173, 1663-1665.	1.3	12
114	Prognostic and predictive value of copy number alterations in invasive breast cancer as determined by multiplex ligation-dependent probe amplification. Cellular Oncology (Dordrecht), 2014, 37, 107-118.	2.1	14
115	Expression profiling of breast cancer patients treated with tamoxifen: prognostic or predictive significance. Medical Oncology, 2014, 31, 896.	1.2	8
116	Myoclonus dystonia syndrome: a novel Îµ-sarcoglycan gene mutation with variable clinical symptoms. Gene, 2014, 548, 306-307.	1.0	1
117	The expression analysis of LATS2 gene in de novo AML patients. Medical Oncology, 2014, 31, 961.	1.2	13
118	Expression of Glycogen synthase kinase 3-Î² (GSK3-Î²) gene in azoospermic men. Iranian Journal of Reproductive Medicine, 2014, 12, 313-20.	0.8	5
119	Association Study of rs3184504 C>T Polymorphism in Patients With Coronary Artery Disease. International Journal of Molecular and Cellular Medicine, 2014, 3, 157-65.	1.1	2
120	Lack of association between LXRÎ± and LXRÎ² gene polymorphisms and prevalence of metabolic syndrome: A caseâ€“control study of an Iranian population. Gene, 2013, 532, 288-293.	1.0	7
121	Expression Analysis of Two Cancer-testis Genes, FBXO39 and TDRD4, in Breast Cancer Tissues and Cell Lines. Asian Pacific Journal of Cancer Prevention, 2013, 14, 6625-6629.	0.5	31
122	Authors response. Indian Journal of Medical Research, 2012, 135, 138.	0.4	0
123	A Novel Mutation in the Transactivation-Regulating Domain of the Androgen Receptor in a Patient With Azoospermia. Journal of Andrology, 2011, 32, 367-370.	2.0	20
124	The prevalence of common CFTR mutations in Iranian infertile men with non-CAVD obstructive azoospermia by using ARMS PCR techniques. Journal of Assisted Reproduction and Genetics, 2011, 28, 1087-1090.	1.2	12
125	BAK, BAX, and NBK/BIK Proapoptotic Gene Alterations in Iranian Patients with Ataxia Telangiectasia. Journal of Clinical Immunology, 2010, 30, 132-137.	2.0	7
126	High prevalence of AZFb microdeletion in Iranian patients with idiopathic non-obstructive azoospermia. Indian Journal of Medical Research, 2010, 132, 265-70.	0.4	18

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127	7.017 Y chromosome microdeletions in Iranian infertile men. Reproductive BioMedicine Online, 2008, 16, s50-s51.	1.1	1
128	The first study of galactose-1-phosphate uridyl transferase mutations in Iranian galactosemia patients. Clinical Biochemistry, 2006, 39, 697-699.	0.8	3
129	First Study of CF Mutations in the CFTR Gene of Iranian Patients: Detection of F508, G542X, W1282X, A120T, R117H, and R347H Mutations. Journal of Tropical Pediatrics, 2004, 50, 359-361.	0.7	19
130	Karyotypic evolution: cytogenetics follow-up study in childhood acute lymphoblastic leukemia. Asian Pacific Journal of Cancer Prevention, 2003, 4, 358-68.	0.5	4
131	Cytogenetic abnormalities in the lymphocytes of a female patient with primary breast carcinoma. Cancer Genetics and Cytogenetics, 2002, 132, 169-170.	1.0	2
132	Association between single nucleotide polymorphisms rs12722489 and multiple sclerosis in Iranian patients with multiple sclerosis. Current Journal of Neurology, 0, , .	0.0	1
133	Gnathodiaphyseal dysplasia with a novel genetic variant in a large family from Iran. Molecular Genetics & Genomic Medicine, 0, , .	0.6	1