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	Expression analysis of four long noncoding RNAs in breast cancer. Tumor Biology, 2016, 37, 2933-2940.	0.8	104
2	Glutamate receptor, metabotropic 7 (GRM7) gene variations and susceptibility to autism: A caseâ€“control study. Autism Research, 2016, 9, 1161-1168.	2.1	57
3	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
4	Association of Increased Levels of lncRNA H19 in PBMCs with Risk of Coronary Artery Disease. Cell Journal, 2019, 20, 564-568.	0.2	41
5	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
6	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
7	<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
8	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
9	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
10	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
11	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
12	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
13	Bioinformatics prioritization of SNPs perturbing microRNA regulation of hematological malignancy-implicated genes. Genomics, 2015, 106, 360-366.	1.3	24
14	RIT2 , a susceptibility gene for Parkinson's disease in Iranian population. Neurobiology of Aging, 2014, 35, e27-e28.	1.5	23
15	Investigation of piwi-interacting RNA pathway genes role in idiopathic non-obstructive azoospermia. Scientific Reports, 2018, 8, 142.	1.6	23
16	SOCS gene family expression profile in the blood of multiple sclerosis patients. Journal of the Neurological Sciences, 2017, 375, 481-485.	0.3	22
17	Expression Study and Clinical Correlations of MYC and CCAT2 in Breast Cancer Patients. Iranian Biomedical Journal, 2017, 21, 303-311.	0.4	22
18	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

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19	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
20	The role of parental microRNA alleles in recurrent pregnancy loss: an association study. Reproductive BioMedicine Online, 2017, 34, 325-330.	1.1	20
21	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
22	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
23	The study of MED12 gene mutations in uterine leiomyomas from Iranian patients. Tumor Biology, 2016, 37, 1567-1571.	0.8	19
24	Expression analysis of AFAP1-AS1 and AFAP1 in breast cancer. Cancer Biomarkers, 2018, 22, 49-54.	0.8	19
25	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
26	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
27	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
28	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
29	Mutational screening of theNR5A1in azoospermia. Andrologia, 2015, 47, 395-401.	1.0	15
30	ACE gene rs4343 polymorphism elevates the risk of preeclampsia in pregnant women. Journal of Human Hypertension, 2018, 32, 825-830.	1.0	15
31	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
32	ANRIL and ANRASSF1 long noncoding RNAs are upregulated in gastric cancer. Journal of Cellular Biochemistry, 2019, 120, 12544-12548.	1.2	14
33	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
34	The expression analysis of LATS2 gene in de novo AML patients. Medical Oncology, 2014, 31, 961.	1.2	13
35	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
36	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

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37	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
38	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
39	Thiamine-responsive megaloblastic anemia syndrome with Ebstein anomaly: a case report. European Journal of Pediatrics, 2014, 173, 1663-1665.	1.3	12
40	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
41	Association of SRD5A2 gene mutations with risk of hypospadias in the Iranian population. Journal of Endocrinological Investigation, 2017, 40, 391-396.	1.8	11
42	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
43	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
44	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
45	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
46	Association study of FOXP3 gene and the risk of 0020 pre-eclampsia. Clinical and Experimental Hypertension, 2018, 40, 613-616.	0.5	10
47	Study of Linc00574 Regulatory Effect on the TCTE3 Expression in Sperm Motility. Reproductive Sciences, 2021, 28, 159-165.	1.1	10
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	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
51	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
52	RARâ€related orphan receptor A: One gene with multiple functions related to migraine. CNS Neuroscience and Therapeutics, 2020, 26, 1315-1321.	1.9	9
53	Emerging Roles of Long Non-coding RNAs in Uterine Leiomyoma Pathogenesis: a Review. Reproductive Sciences, 2022, 29, 1086-1101.	1.1	9
54	A bioinformatics approach for identification of miR-100 targets implicated in breast cancer. Cellular and Molecular Biology, 2017, 63, 99-105.	0.3	9

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55	rs508485 Polymorphism Is Associated with Non-obstructive Azoospermia in Iranian Patients. Reports of Biochemistry and Molecular Biology, 2017, 5, 108-111.	0.5	9
56	Expression profiling of breast cancer patients treated with tamoxifen: prognostic or predictive significance. Medical Oncology, 2014, 31, 896.	1.2	8
57	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
58	Angiotensinâ€converting enzyme gene rs4343 polymorphism increases susceptibility to migraine. CNS Neuroscience and Therapeutics, 2017, 23, 698-699.	1.9	8
59	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
60	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
61	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
62	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
63	Treatment Failure in Acute Myeloid Leukemia: Focus on the Role of Extracellular Vesicles. Leukemia Research, 2022, 112, 106751.	0.4	8
64	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
65	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
66	TRAIL gene expression analysis in multiple sclerosis patients. Human Antibodies, 2016, 24, 33-38.	0.6	7
67	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
68	CYP24A1 expression analysis in uterine leiomyoma regarding MED12 mutation profile. Archives of Gynecology and Obstetrics, 2021, 303, 787-792.	0.8	7
69	Aberrant WntÎ²-Catenin Signaling Pathway in Testis of Azoospermic Menâ€. Advanced Pharmaceutical Bulletin, 2015, 5, 373-377.	0.6	7
70	An Iranian family with azoospermia and premature ovarian insufficiency segregating NR5A1 mutation. Climacteric, 2014, 17, 301-303.	1.1	6
71	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
72	A candidate intronic CYP24A1 gene variant affects the risk of colorectal cancer. Biomarkers in Medicine, 2020, 14, 23-29.	0.6	5

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73	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
74	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
75	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
76	Circular RNAs: Novel Biomarkers in Spermatogenesis Defects and Male Infertility. Reproductive Sciences, 2023, 30, 62-71.	1.1	5
77	Autophagy ATG16L1Ârs2241880 impacts the colorectal cancer risk: A caseâ€control study. Journal of Clinical Laboratory Analysis, 2022, 36, e24169.	0.9	5
78	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
79	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
80	Ectopic expression of CYP24A1 circular RNA hsa_circ_0060927 in uterine leiomyomas. Journal of Clinical Laboratory Analysis, 2020, 34, e23114.	0.9	4
81	Could CYP24A1 promoter methylation status affect the gene expression in the colorectal cancer patients?. Meta Gene, 2020, 24, 100656.	0.3	4
82	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
83	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
84	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
85	Karyotypic evolution: cytogenetics follow-up study in childhood acute lymphoblastic leukemia. Asian Pacific Journal of Cancer Prevention, 2003, 4, 358-68.	0.5	4
86	Reduced expression of <scp> CFAP44 </scp> and <scp> CFAP44â€AS1 </scp> may affect sperm motility and morphology. Andrologia, 2022, 54, e14447.	1.0	4
87	The first study of galactose-1-phosphate uridyl transferase mutations in Iranian galactosemia patients. Clinical Biochemistry, 2006, 39, 697-699.	0.8	3
88	Association of Î²-Secretase Functional Polymorphism with Risk of Schizophrenia. Genetic Testing and Molecular Biomarkers, 2017, 21, 248-251.	0.3	3
89	Association analysis of the GABRB3 promoter variant and susceptibility to autism spectrum disorder. Basal Ganglia, 2018, 11, 4-7.	0.3	3
90	Contribution of long noncoding RNA HOTAIR variants to preeclampsia susceptibility in Iranian women. Hypertension in Pregnancy, 2021, 40, 29-35.	0.5	3

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91	CREB-binding protein (CREBBP) and preeclampsia: a new promising target gene. Molecular Biology Reports, 2021, 48, 2117-2122.	1.0	3
92	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
93	TTY2 genes deletions as genetic risk factor of male infertility. Cellular and Molecular Biology, 2017, 63, 57.	0.3	3
94	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
95	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
96	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
97	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
98	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
99	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
100	Cytogenetic abnormalities in the lymphocytes of a female patient with primary breast carcinoma. Cancer Genetics and Cytogenetics, 2002, 132, 169-170.	1.0	2
101	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
102	Expression analysis of CBR3-AS1 and androgen receptor genes in breast cancer. Meta Gene, 2018, 17, 82-87.	0.3	2
103	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
104	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
105	DACT1 variants and colorectal cancer. British Journal of Biomedical Science, 2021, 78, 221-224.	1.2	2
106	MIR17HG Gene Polymorphism and the Risk of Recurrent Spontaneous Abortion. Gene, Cell and Tissue, 2016, 3, .	0.2	2
107	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
108	Overexpression of GABRP Gene in Triple Negative Breast Cancer: Molecular Mechanisms and Interpretation. International Journal of Cancer Management, 2021, 14, .	0.2	2

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109	7.017 Y chromosome microdeletions in Iranian infertile men. Reproductive BioMedicine Online, 2008, 16, s50-s51.	1.1	1
110	Myoclonus dystonia syndrome: a novel Îµ-sarcoglycan gene mutation with variable clinical symptoms. Gene, 2014, 548, 306-307.	1.0	1
111	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
112	A novelSRD5A2mutation in an Iranian family with sex development disorder. Andrologia, 2021, 53, e13847.	1.0	1
113	MiR-206 Target Prediction in Breast Cancer Subtypes by Bioinformatics Tools. International Journal of Cancer Management, 2018, 11, .	0.2	1
114	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
115	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
116	Association between single nucleotide polymorphisms rs12722489 and multiple sclerosis in Iranian patients with multiple sclerosis. Current Journal of Neurology, 0, , .	0.0	1
117	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
118	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
119	Exon 1 Mutational Screening in Iranian Patients with Uterine Leiomyoma. Reports of Biochemistry and Molecular Biology, 2019, 8, 21-24.	0.5	1
120	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
121	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
122	Gnathodiaphyseal dysplasia with a novel genetic variant in a large family from Iran. Molecular Genetics & Genomic Medicine, 0, , .	0.6	1
123	The association between FOXP3 gene variations and autism: True or false positive?. Gene, 2017, 635, 1-2.	1.0	0
124	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
125	Mutational screening of RTK-BRAF genes in de novo adult acute myeloid leukemia. Gene Reports, 2020, 21, 100904.	0.4	0
126	Authors response. Indian Journal of Medical Research, 2012, 135, 138.	0.4	0

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127	An Association Study of rs2829803 Polymorphism in miRNA-155 with Multiple Sclerosis in Patients Referred to Farshchian Hospital of Hamadan City, (Iran). Majallah-i DÄnishgÀh-i ì'UlÅ«m-i PizishkÄ«-i Qum, 2018, 12, 28-34.	0.2	0
128	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
129	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
130	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
131	Coding and Non-Coding RNAs, as Male Fertility and Infertility Biomarkers. International Journal of Fertility & Sterility, 2021, 15, 158-166.	0.2	0
132	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
133	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