

Mohammad R Abbaszadegan

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3795585/publications.pdf>

Version: 2024-02-01

116
papers

1,739
citations

257101

24
h-index

360668

35
g-index

123
all docs

123
docs citations

123
times ranked

2391
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Construction and Quantitative Evaluation of a Tissue-Specific Sleeping Beauty by EDL2-Specific Transposase Expression in Esophageal Squamous Carcinoma Cell Line KYSE-30. <i>Molecular Biotechnology</i> , 2023, 65, 350-360. | 1.3 | 1 |
| 2 | Elucidated tumorigenic role of MAML1 and TWIST1 in gastric cancer is associated with Helicobacter pylori infection. <i>Microbial Pathogenesis</i> , 2022, 162, 105304. | 1.3 | 4 |
| 3 | Kindlin1 As a Sex and Location Specific Diagnostic Marker in Gastric Cancer Patients. <i>Iranian Journal of Pathology</i> , 2022, 17, 23-28. | 0.2 | 2 |
| 4 | Cardiac Tamponade: A Rare Manifestation of Familial Mediterranean Fever. <i>Case Reports in Rheumatology</i> , 2022, 2022, 1-5. | 0.2 | 5 |
| 5 | MicroRNA-217: a therapeutic and diagnostic tumor marker. <i>Expert Review of Molecular Diagnostics</i> , 2022, 22, 61-76. | 1.5 | 11 |
| 6 | Long non-coding RNA AC087388.1 as a novel biomarker in colorectal cancer. <i>BMC Cancer</i> , 2022, 22, 196. | 1.1 | 2 |
| 7 | Biallelic Variants in the Ectonucleotidase <i>ENTPD1</i> Cause a Complex Neurodevelopmental Disorder with Intellectual Disability, Distinct White Matter Abnormalities, and Spastic Paraplegia. <i>Annals of Neurology</i> , 2022, 92, 304-321. | 2.8 | 2 |
| 8 | SOX2/SALL4 stemness axis modulates Notch signaling genes to maintain self-renewal capacity of esophageal squamous cell carcinoma. <i>Molecular and Cellular Biochemistry</i> , 2021, 476, 921-929. | 1.4 | 11 |
| 9 | Mechanisms of long non-coding RNA function in colorectal cancer tumorigenesis. <i>Asia-Pacific Journal of Clinical Oncology</i> , 2021, 17, 7-23. | 0.7 | 32 |
| 10 | Genetic and molecular biology of systemic lupus erythematosus among Iranian patients: an overview. <i>Autoimmunity Highlights</i> , 2021, 12, 2. | 3.9 | 5 |
| 11 | A novel mutation in the cathepsin C (CTSC) gene in Iranian family with Papillon-Lefevre syndrome. <i>Clinical and Experimental Dental Research</i> , 2021, 7, 568-573. | 0.8 | 2 |
| 12 | Crosstalk between MMP-13, CD44, and TWIST1 and its role in regulation of EMT in patients with esophageal squamous cell carcinoma. <i>Molecular and Cellular Biochemistry</i> , 2021, 476, 2465-2478. | 1.4 | 12 |
| 13 | MAEL as a diagnostic marker for the early detection of esophageal squamous cell carcinoma. <i>Diagnostic Pathology</i> , 2021, 16, 36. | 0.9 | 1 |
| 14 | Correlation between the immune checkpoints and EMT genes proposes potential prognostic and therapeutic targets in ESCC. <i>Journal of Molecular Histology</i> , 2021, 52, 597-609. | 1.0 | 16 |
| 15 | Role of miRNA gene variants in the susceptibility and pharmacogenetics of colorectal cancer. <i>Pharmacogenomics</i> , 2021, 22, 303-318. | 0.6 | 4 |
| 16 | Methylation as a critical epigenetic process during tumor progressions among Iranian population: an overview. <i>Genes and Environment</i> , 2021, 43, 14. | 0.9 | 1 |
| 17 | Interaction between LINC-ROR and Stemness State in Gastric Cancer Cells with Helicobacter pylori Infection. <i>Iranian Biomedical Journal</i> , 2021, 25, 157-168. | 0.4 | 7 |
| 18 | Genotyping of ABCC8, KCNJ11, and HADH in Iranian Infants with Congenital Hyperinsulinism. <i>Case Reports in Endocrinology</i> , 2021, 2021, 1-6. | 0.2 | 0 |

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 19 | Single nucleotide polymorphisms as the efficient prognostic markers in breast cancer. <i>Current Cancer Drug Targets</i> , 2021, 21, . | 0.8 | 0 |
| 20 | The Level of Mesenchymal-Epithelial Transition Autophosphorylation is Correlated with Esophageal Squamous Cell Carcinoma Migration. <i>Iranian Biomedical Journal</i> , 2021, 25, 243-254. | 0.4 | 0 |
| 21 | Genetically engineered mouse models of esophageal cancer. <i>Experimental Cell Research</i> , 2021, 406, 112757. | 1.2 | 8 |
| 22 | A Systematic Review on the Genotoxic Effects of Selective Serotonin Reuptake Inhibitors. <i>Advances in Experimental Medicine and Biology</i> , 2021, 1286, 115-124. | 0.8 | 3 |
| 23 | Non-collagenous extracellular matrix protein dermatopontin may play a role as another component of transforming growth factor- β^2 signaling pathway in colon carcinogenesis. <i>Iranian Journal of Basic Medical Sciences</i> , 2021, 24, 444-450. | 1.0 | 0 |
| 24 | Allogeneic tumor cell line-based vaccines: A good alternative to autologous and cancer stem cell vaccines in colorectal cancer.. <i>Iranian Journal of Basic Medical Sciences</i> , 2021, 24, 1231-1239. | 1.0 | 1 |
| 25 | MAEL Cancer-Testis Antigen as a Diagnostic Marker in Primary Stages of Gastric Cancer with <i>Helicobacter pylori</i> Infection. <i>Journal of Gastrointestinal Cancer</i> , 2020, 51, 17-22. | 0.6 | 7 |
| 26 | Role of DIDO1 in Progression of Esophageal Squamous Cell Carcinoma. <i>Journal of Gastrointestinal Cancer</i> , 2020, 51, 83-87. | 0.6 | 7 |
| 27 | Whole Exome Sequencing Reveals a Novel Damaging Mutation in Human Fibroblast Activation Protein in a Family with Esophageal Squamous Cell Carcinoma. <i>Journal of Gastrointestinal Cancer</i> , 2020, 51, 179-188. | 0.6 | 5 |
| 28 | In silico evidence of high frequency of miRNA-related SNPs in Esophageal Squamous Cell Carcinoma. <i>Journal of Cellular Physiology</i> , 2020, 235, 966-978. | 2.0 | 3 |
| 29 | TWIST1 upregulates matrix metalloproteinase (MMP) genes family in esophageal squamous carcinoma cells. <i>Gene Expression Patterns</i> , 2020, 37, 119127. | 0.3 | 4 |
| 30 | Integration analysis of long non-coding RNA (lncRNA) role in tumorigenesis of colon adenocarcinoma. <i>BMC Medical Genomics</i> , 2020, 13, 108. | 0.7 | 52 |
| 31 | Novel Deleterious Mutation in Steroid-5 α -Reductase-2 in 46, XY Disorders of Sex Development: Case Report Study. <i>Fetal and Pediatric Pathology</i> , 2020, , 1-8. | 0.4 | 2 |
| 32 | GSTs polymorphisms are associated with epigenetic silencing of CDKN2A gene in esophageal squamous cell carcinoma. <i>Environmental Science and Pollution Research</i> , 2020, 27, 31269-31277. | 2.7 | 4 |
| 33 | Role of extra cellular proteins in gastric cancer progression and metastasis: an update. <i>Genes and Environment</i> , 2020, 42, 18. | 0.9 | 8 |
| 34 | MAML1 promotes ESCC aggressiveness through upregulation of EMT marker TWIST1. <i>Molecular Biology Reports</i> , 2020, 47, 2659-2668. | 1.0 | 14 |
| 35 | Primary Angle Closure Glaucoma-associated Genetic Polymorphisms in Northeast Iran. <i>Journal of Ophthalmic and Vision Research</i> , 2020, 15, 45-52. | 0.7 | 4 |
| 36 | Genetic and molecular bases of esophageal Cancer among Iranians: an update. <i>Diagnostic Pathology</i> , 2019, 14, 97. | 0.9 | 4 |

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 37 | Effects of selective serotonin reuptake inhibitors on DNA damage in patients with depression. <i>Journal of Psychopharmacology</i> , 2019, 33, 1364-1376. | 2.0 | 32 |
| 38 | Novel mutation in AIRE gene with autoimmune polyendocrine syndrome type 1. <i>Immunobiology</i> , 2019, 224, 728-733. | 0.8 | 4 |
| 39 | Mutation analysis of genes related to methylmalonic acidemia: identification of eight novel mutations. <i>Molecular Biology Reports</i> , 2019, 46, 271-285. | 1.0 | 4 |
| 40 | MAML1 regulates EMT markers expression through NOTCH-independent pathway in breast cancer cell line MCF7. <i>Biochemical and Biophysical Research Communications</i> , 2019, 510, 376-382. | 1.0 | 13 |
| 41 | Linkage between EMT and stemness state through molecular association between TWIST1 and NY-ESO1 in esophageal squamous cell carcinoma. <i>Biochimie</i> , 2019, 163, 84-93. | 1.3 | 12 |
| 42 | MEIS1 knockdown may promote differentiation of esophageal squamous carcinoma cell line KYSE30. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e00746. | 0.6 | 12 |
| 43 | TWIST1, MMP21, and HLAG1 co-overexpression is associated with ESCC aggressiveness. <i>Journal of Cellular Biochemistry</i> , 2019, 120, 14838-14846. | 1.2 | 6 |
| 44 | Role of MAML1 in targeted therapy against the esophageal cancer stem cells. <i>Journal of Translational Medicine</i> , 2019, 17, 126. | 1.8 | 32 |
| 45 | Mutation Screening of KCNQ1 and KCNE1 Genes in Iranian Patients With Jervell and Lange-Nielsen Syndrome. <i>Fetal and Pediatric Pathology</i> , 2019, 38, 273-281. | 0.4 | 6 |
| 46 | Homozygous Null TBX4 Mutations Lead to Posterior Amelia with Pelvic and Pulmonary Hypoplasia. <i>American Journal of Human Genetics</i> , 2019, 105, 1294-1301. | 2.6 | 17 |
| 47 | Withdrawal Notice: The Prognostic Value of Prognostic Biomarkers in Esophageal Squamous Cell Carcinoma in Iranian Population. <i>Current Cancer Therapy Reviews</i> , 2019, 15, . | 0.2 | 0 |
| 48 | Ovarian cancer stem cells and targeted therapy. <i>Journal of Ovarian Research</i> , 2019, 12, 120. | 1.3 | 70 |
| 49 | The association between serum irisin levels and cardiovascular disease in diabetic patients. <i>Diabetes and Metabolic Syndrome: Clinical Research and Reviews</i> , 2019, 13, 786-790. | 1.8 | 26 |
| 50 | Induction of T cell-mediated immune response by dendritic cells pulsed with mRNA of sphere-forming cells isolated from patients with gastric cancer. <i>Life Sciences</i> , 2019, 219, 136-143. | 2.0 | 19 |
| 51 | Novel DNA variation of GPR54 gene in familial central precocious puberty. <i>Italian Journal of Pediatrics</i> , 2019, 45, 10. | 1.0 | 13 |
| 52 | ErbB1 and ErbB3 co-over expression as a prognostic factor in gastric cancer. <i>Biological Research</i> , 2019, 52, 2. | 1.5 | 29 |
| 53 | Combination of Genetics and Nanotechnology for Down Syndrome Modification: A Potential Hypothesis and Review of the Literature. <i>Iranian Journal of Public Health</i> , 2019, 48, 371-378. | 0.3 | 2 |
| 54 | The Role of Interleukin-4 and 13 Gene Polymorphisms in Allergic Rhinitis: A Case Control Study. <i>Reports of Biochemistry and Molecular Biology</i> , 2019, 8, 111-118. | 0.5 | 10 |

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 55 | Suppression of dsRNA response genes and innate immunity following Oct4, Stella, and Nanos2 overexpression in mouse embryonic fibroblasts. <i>Cytokine</i> , 2018, 106, 1-11. | 1.4 | 7 |
| 56 | Familial Esophageal Squamous Cell Carcinoma with damaging rare/germline mutations in KCNJ12/KCNJ18 and GPRIN2 genes. <i>Cancer Genetics</i> , 2018, 221, 46-52. | 0.2 | 20 |
| 57 | Four novel mutations of the <i>BCKDHA</i> , <i>BCKDHB</i> and <i>DBT</i> genes in Iranian patients with maple syrup urine disease. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2018, 31, 205-212. | 0.4 | 4 |
| 58 | Role of MAML1 and MEIS1 in Esophageal Squamous Cell Carcinoma Depth of Invasion. <i>Pathology and Oncology Research</i> , 2018, 24, 245-250. | 0.9 | 15 |
| 59 | Cytokine networks and their association with <i>Helicobacter pylori</i> infection in gastric carcinoma. <i>Journal of Cellular Physiology</i> , 2018, 233, 2791-2803. | 2.0 | 32 |
| 60 | Predicting the Correlation of EZH2 and Cancer Stem Cell Markers in Esophageal Squamous Cell Carcinoma. <i>Journal of Gastrointestinal Cancer</i> , 2018, 49, 437-441. | 0.6 | 11 |
| 61 | Mutations in HNF1A Gene are not a Common Cause of Familial Young-Onset Diabetes in Iran. <i>Indian Journal of Clinical Biochemistry</i> , 2018, 33, 91-95. | 0.9 | 6 |
| 62 | Novel mutations and their genotype-phenotype correlations in patients with Noonan syndrome, using next-generation sequencing. <i>Advances in Medical Sciences</i> , 2018, 63, 87-93. | 0.9 | 15 |
| 63 | Genetic and molecular origins of colorectal Cancer among the Iranians: an update. <i>Diagnostic Pathology</i> , 2018, 13, 97. | 0.9 | 6 |
| 64 | WNT and NOTCH signaling pathways as activators for epidermal growth factor receptor in esophageal squamous cell carcinoma. <i>Cellular and Molecular Biology Letters</i> , 2018, 23, 42. | 2.7 | 39 |
| 65 | Contribution of KCTD12 to esophageal squamous cell carcinoma. <i>BMC Cancer</i> , 2018, 18, 853. | 1.1 | 16 |
| 66 | Isolation and identification of chemotherapy-enriched sphere-forming cells from a patient with gastric cancer. <i>Journal of Cellular Physiology</i> , 2018, 233, 7036-7046. | 2.0 | 11 |
| 67 | Childhood Sex-Typed Behavior and Gender Change in Individuals with 46,XY and 46,XX Disorders of Sex Development: An Iranian Multicenter Study. <i>Archives of Sexual Behavior</i> , 2018, 47, 2287-2298. | 1.2 | 11 |
| 68 | Molecular Signaling in Tumorigenesis of Gastric Cancer. <i>Iranian Biomedical Journal</i> , 2018, 22, 217-30. | 0.4 | 27 |
| 69 | Investigation of melanoma-associated antigen A4 cancer/testis antigen clinical relevance in esophageal squamous cell carcinoma. <i>Journal of Cancer Research and Therapeutics</i> , 2018, 14, 1059-1064. | 0.3 | 7 |
| 70 | Ectopic Expression of Human Gene in ESCC Cell Line Using Retroviral System. <i>Avicenna Journal of Medical Biotechnology</i> , 2018, 10, 75-82. | 0.2 | 1 |
| 71 | Promoter Hypermethylation of the Eyes Absent 4 Gene is a Tumor-Specific Epigenetic Biomarker in Iranian Colorectal Cancer Patients. <i>Acta Medica Iranica</i> , 2018, 56, 21-27. | 0.8 | 4 |
| 72 | Contribution of MAML1 in esophageal squamous cell carcinoma tumorigenesis. <i>Annals of Diagnostic Pathology</i> , 2017, 27, 79-82. | 0.6 | 8 |

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 73 | miRNA-Related Polymorphisms in miR-423 (rs6505162) and <i>PEX6</i> (rs1129186) and Risk of Esophageal Squamous Cell Carcinoma in an Iranian Cohort. <i>Genetic Testing and Molecular Biomarkers</i> , 2017, 21, 382-390. | 0.3 | 26 |
| 74 | Novel candidate genes may be possible predisposing factors revealed by whole exome sequencing in familial esophageal squamous cell carcinoma. <i>Tumor Biology</i> , 2017, 39, 101042831769911. | 0.8 | 5 |
| 75 | Crosstalk between SHH and stemness state signaling pathways in esophageal squamous cell carcinoma. <i>Journal of Cell Communication and Signaling</i> , 2017, 11, 147-153. | 1.8 | 8 |
| 76 | Isolation, identification, and characterization of cancer stem cells: A review. <i>Journal of Cellular Physiology</i> , 2017, 232, 2008-2018. | 2.0 | 157 |
| 77 | Sexual orientation and medical history among Iranian people with Complete Androgen Insensitivity Syndrome and Congenital Adrenal Hyperplasia. <i>Journal of Psychosomatic Research</i> , 2017, 92, 55-62. | 1.2 | 16 |
| 78 | Expression analysis of matrix metalloproteinase-13 in human gastric cancer in the presence of <i>Helicobacter Pylori</i> infection. <i>Cancer Biomarkers</i> , 2017, 18, 349-356. | 0.8 | 9 |
| 79 | TWIST1 upregulates the MAGEA4 oncogene. <i>Molecular Carcinogenesis</i> , 2017, 56, 877-885. | 1.3 | 32 |
| 80 | Applying Subtractive Hybridization Technique to Enrich and Amplify Tumor-Specific Transcripts of Esophageal Squamous Cell Carcinoma. <i>Pathology and Oncology Research</i> , 2017, 23, 271-279. | 0.9 | 2 |
| 81 | Negative Regulatory Role of TWIST1 on SNAIL Gene Expression. <i>Pathology and Oncology Research</i> , 2017, 23, 85-90. | 0.9 | 16 |
| 82 | Ectopic expression of TWIST1 upregulates the stemness marker OCT4 in the esophageal squamous cell carcinoma cell line KYSE30. <i>Cellular and Molecular Biology Letters</i> , 2017, 22, 33. | 2.7 | 19 |
| 83 | Biological and Clinicopathological Significance of Cripto-1 Expression in the Progression of Human ESCC. <i>Reports of Biochemistry and Molecular Biology</i> , 2017, 5, 83-90. | 0.5 | 8 |
| 84 | Gene Polymorphisms Associated with Allergic Rhinitis in an Iranian Population. <i>Reports of Biochemistry and Molecular Biology</i> , 2017, 5, 97-102. | 0.5 | 3 |
| 85 | Disease Biomarkers in Gastrointestinal Malignancies. <i>Disease Markers</i> , 2016, 2016, 1-3. | 0.6 | 2 |
| 86 | Presence of the RET Cys634Tyr mutation and Gly691Ser functional polymorphism in Iranian families with multiple endocrine neoplasia type 2A. <i>Hormones</i> , 2016, 15, 65-72. | 0.9 | 2 |
| 87 | Identification of a novel deletion in the MMAA gene in two Iranian siblings with vitamin B12-responsive methylmalonic acidemia. <i>Cellular and Molecular Biology Letters</i> , 2016, 21, 4. | 2.7 | 7 |
| 88 | Correlation of Wnt and NOTCH pathways in esophageal squamous cell carcinoma. <i>Journal of Cell Communication and Signaling</i> , 2016, 10, 129-135. | 1.8 | 47 |
| 89 | Correlation Between Meis1 and Msi1 in Esophageal Squamous Cell Carcinoma. <i>Journal of Gastrointestinal Cancer</i> , 2016, 47, 273-277. | 0.6 | 25 |
| 90 | SOX1 is correlated to stemness state regulator SALL4 through progression and invasiveness of esophageal squamous cell carcinoma. <i>Gene</i> , 2016, 594, 171-175. | 1.0 | 18 |

| # | ARTICLE | IF | CITATIONS |
|-----|--|-----|-----------|
| 91 | Psychosexual Outcome Among Iranian Individuals with 5 α -Reductase Deficiency Type 2 and Its Relationship with Parental Sexism. <i>Journal of Sexual Medicine</i> , 2016, 13, 1629-1641. | 0.3 | 17 |
| 92 | In silico dissection of miRNA targetome polymorphisms and their role in regulating miRNA-mediated gene expression in esophageal cancer. <i>Cell Biochemistry and Biophysics</i> , 2016, 74, 483-497. | 0.9 | 18 |
| 93 | Identification of four novel mutations of the WFS1 gene in Iranian Wolfram syndrome pedigrees. <i>Acta Diabetologica</i> , 2016, 53, 899-904. | 1.2 | 2 |
| 94 | Contribution of EVX1 in Aggressiveness of Esophageal Squamous Cell Carcinoma. <i>Pathology and Oncology Research</i> , 2016, 22, 341-347. | 0.9 | 6 |
| 95 | Role of Msi1 and PYGO2 in esophageal squamous cell carcinoma depth of invasion. <i>Journal of Cell Communication and Signaling</i> , 2016, 10, 49-53. | 1.8 | 29 |
| 96 | Predicting the molecular role of MEIS1 in esophageal squamous cell carcinoma. <i>Tumor Biology</i> , 2016, 37, 1715-1725. | 0.8 | 29 |
| 97 | Loss of heterozygosity and microsatellite instability as predictive markers among Iranian esophageal cancer patients. <i>Iranian Journal of Basic Medical Sciences</i> , 2016, 19, 726-33. | 1.0 | 4 |
| 98 | Linc-ROR and its spliced variants 2 and 4 are significantly up-regulated in esophageal squamous cell carcinoma. <i>Iranian Journal of Basic Medical Sciences</i> , 2016, 19, 1131-1135. | 1.0 | 18 |
| 99 | Variation in the miRNA-433 binding site of FGF20 is a risk factor for Parkinson's disease in Iranian population. <i>Journal of the Neurological Sciences</i> , 2015, 355, 72-74. | 0.3 | 25 |
| 100 | Role of Msi1 and MAML1 in Regulation of Notch Signaling Pathway in Patients with Esophageal Squamous Cell Carcinoma. <i>Journal of Gastrointestinal Cancer</i> , 2015, 46, 365-369. | 0.6 | 29 |
| 101 | Chromosomal analysis of couples with repeated spontaneous abortions in northeastern Iran. <i>International Journal of Fertility & Sterility</i> , 2015, 9, 47-54. | 0.2 | 39 |
| 102 | Presence of the RET Cys634Tyr mutation and Gly691Ser functional polymorphism in Iranian families with multiple endocrine neoplasia type 2A. <i>Hormones</i> , 2015, 15, 65-72. | 0.9 | 1 |
| 103 | Expression analysis of CD44 isoforms S and V3, in patients with esophageal squamous cell carcinoma. <i>Iranian Journal of Basic Medical Sciences</i> , 2015, 18, 380-4. | 1.0 | 5 |
| 104 | Protein modeling of cathepsin C mutations found in Papillon-Lefèvre syndrome. <i>Gene</i> , 2014, 538, 182-187. | 1.0 | 14 |
| 105 | Stemness state regulators SALL4 and SOX2 are involved in progression and invasiveness of esophageal squamous cell carcinoma. <i>Medical Oncology</i> , 2014, 31, 922. | 1.2 | 81 |
| 106 | Association of ADAM33 gene polymorphisms with allergic asthma. <i>Iranian Journal of Basic Medical Sciences</i> , 2014, 17, 716-21. | 1.0 | 5 |
| 107 | Role of Brg1 in progression of esophageal squamous cell carcinoma. <i>Iranian Journal of Basic Medical Sciences</i> , 2014, 17, 912-7. | 1.0 | 9 |
| 108 | Inherited genetic markers for thrombophilia in northeastern Iran (a clinical-based report). <i>Reports of Biochemistry and Molecular Biology</i> , 2014, 2, 76-82. | 0.5 | 1 |

| # | ARTICLE | IF | CITATIONS |
|-----|--|-----|-----------|
| 109 | Two novel mutations in CYP11B1 and modeling the consequent alterations of the translated protein in classic congenital adrenal hyperplasia patients. <i>Endocrine</i> , 2013, 44, 212-219. | 1.1 | 14 |
| 110 | Identification of Xq22.1-23 as a region linked with hereditary recurrent spontaneous abortion in a family. <i>Iranian Journal of Reproductive Medicine</i> , 2013, 11, 659-64. | 0.8 | 0 |
| 111 | DNA damage in oral mucosa cells of patients with fixed orthodontic appliances. <i>Journal of Dentistry of Tehran University of Medical Sciences</i> , 2013, 10, 494-500. | 0.4 | 5 |
| 112 | p16 promoter hypermethylation: A useful serum marker for early detection of gastric cancer. <i>World Journal of Gastroenterology</i> , 2008, 14, 2055. | 1.4 | 79 |
| 113 | Rare gross deletion in T-cell immune regulator-1 gene in Iranian family with infantile malignant osteopetrosis. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2008, 29, 1494-6. | 0.5 | 1 |
| 114 | Stool-based DNA testing, a new noninvasive method for colorectal cancer screening, the first report from Iran. <i>World Journal of Gastroenterology</i> , 2007, 13, 1528. | 1.4 | 40 |
| 115 | Rapid DNA extraction protocol from stool, suitable for molecular genetic diagnosis of colon cancer. <i>Iranian Biomedical Journal</i> , 2007, 11, 203-208. | 0.4 | 9 |
| 116 | Prevalence of Human T-Lymphotropic Virus Type 1 among Blood Donors from Mashhad, Iran. <i>Journal of Clinical Microbiology</i> , 2003, 41, 2593-2595. | 1.8 | 66 |