# Mohsen Ghadami

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5939017/publications.pdf

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| 21 | 321 | 933264 | 839398 |
|---|---|---|---|
| papers | citations | 10 h-index | 18 g-index |

| 21 | 21 | 21 | 856 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2018, 141, 1450-1458. | 1.5 | 90 |
| 2 | Bone Marrow Transplantation Restores Follicular Maturation and Steroid Hormones Production in a Mouse Model for Primary Ovarian Failure. PLoS ONE, 2012, 7, e32462. | 1.1 | 33 |
| 3 | Mutations in the histamineN-methyltransferase gene,HNMT, are associated with nonsyndromic autosomal recessive intellectual disability. Human Molecular Genetics, 2015, 24, 5697-5710. | 1.4 | 27 |
| 4 | Circulating free DNA concentration as a marker of disease recurrence and metastatic potential in lung cancer. Clinical and Translational Medicine, 2019, 8, 14. | 1.7 | 22 |
| 5 | Confirmation of genetic homogeneity of syndactyly type 1 in an Iranian family. American Journal of Medical Genetics Part A, 2001, 104, 147-151. | 2.4 | 21 |
| 6 | Demethylation and alterations in the expression level of the cell cycleâ€“related genes as possible mechanisms in arsenic trioxideâ€“induced cell cycle arrest in human breast cancer cells. Tumor Biology, 2017, 39, 101042831769225. | 0.8 | 17 |
| 7 | Association of SP-C gene codon 186 polymorphism (rs1124) and risk of RDS. Journal of Maternal-Fetal and Neonatal Medicine, 2017, 30, 2585-2589. | 0.7 | 16 |
| 8 | Novel EXOSC9 variants cause pontocerebellar hypoplasia type 1D with spinal motor neuronopathy and cerebellar atrophy. Journal of Human Genetics, 2021, 66, 401-407. | 1.1 | 15 |
| 9 | Association of SP-B gene 9306 A/G polymorphism (rs7316) and risk of RDS. Journal of Maternal-Fetal and Neonatal Medicine, 2018, 31, 2965-2970. | 0.7 | 14 |
| 10 | The first cohort of Iranian patients with hyper immunoglobulin E syndrome: A longâ€term followâ€up and genetic analysis. Pediatric Allergy and Immunology, 2019, 30, 469-478. | 1.1 | 14 |
| 11 | MicroRNAs networks in thyroid cancers: focus on miRNAs related to the fascin. Journal of Diabetes and Metabolic Disorders, 2013, 12, 31. | 0.8 | 11 |
| 12 | Investigation of ITGB2 gene in 12 new cases of leukocyte adhesion deficiency-type I revealed four novel mutations from Iran. Archives of Iranian Medicine, 2015, 18, 760-4. | 0.2 | 9 |
| 13 | RPE65 and retinal dystrophy: Report of new and recurrent mutations. Journal of Gene Medicine, 2020, 22, e3154. | 1.4 | 7 |
| 14 | Essential genes in thyroid cancers: focus on fascin. Journal of Diabetes and Metabolic Disorders, 2013, 12, 32. | 0.8 | 6 |
| 15 | Investigation of promoter methylation of FSCN1 gene and FSCN1 protein expression in differentiated thyroid carcinomas. Molecular Biology Reports, 2020, 47, 2161-2169. | 1.0 | 5 |
| 16 | Plasma Levels of MicroRNA-146a-5p, MicroRNA-24-3p, and MicroRNA125a-5p as Potential Diagnostic Biomarkers for Rheumatoid Arthritis. Iranian Journal of Allergy, Asthma and Immunology, 2021, 20, 326-337. | 0.3 | 5 |
| 17 | Evaluation of the Prevalence of Exons 1 And 10 Polymorphisms of Gene and Its Relationship with IVF Success. Journal of Reproduction and Infertility, 2019, 20, 218-224. | 1.0 | 4 |
| 18 | Study of cytomegalovirus infection in idiopathic infertility men referred to Shariati hospital, Tehran, Iran. Iranian Journal of Reproductive Medicine, 2014, 12, 151-4. | 0.8 | 2 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Variable Clinical Manifestations of COVID-19; Viral and Human Genomes Talk. Iranian Journal of Allergy, Asthma and Immunology, 2020, 19, 456-470. | 0.3 | 2 |
| 20 | Cell Free Tumoral DNA Versus Paraffin Block Epidermal Growth Factor Receptor Mutation Detection in Patients with Non-Small Cell Lung Cancer. Asian Pacific Journal of Cancer Prevention, 2019, 20, 3591-3596. | 0.5 | 1 |
| 21 | RET Proto-Oncogene Mutational Analysis in 45 Iranian Patients Affected with Medullary Thyroid Carcinoma: Report of a New Variant. Journal of Thyroid Research, 2021, 2021, 1-9. | 0.5 | 0 |