Eskandar Taghizadeh

List of Publications by Year
in descending order

27	333	1051969 10	993246 17
papers	citations	h-index	g-index

28	28	28	525
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genetics of Familial Combined Hyperlipidemia (FCHL) Disorder: An Update. Biochemical Genetics, 2022, 60, 453-481.	0.8	4
2	The role of efferocytosis in neuro-degenerative diseases. Neurological Sciences, 2022, 43, 1593-1603.	0.9	4
3	Cellular and Molecular Aspects of Managing Familial Hypercholesterolemia: Recent and Emerging Therapeutic Approaches. Endocrine, Metabolic and Immune Disorders - Drug Targets, 2022, 22, 1018-1028.	0.6	3
4	Epigenetic aspects of multiple sclerosis and future therapeutic options. International Journal of Neuroscience, 2021, 131, 56-64.	0.8	12
5	CD47 Functionalization of Nanoparticles as a Poly(ethylene glycol) Alternative: A Novel Approach to Improve Drug Delivery. Current Drug Targets, 2021, 22, 1750-1759.	1.0	10
6	LncRNAs as putative biomarkers and therapeutic targets for Parkinsonâ€™s disease. Neurological Sciences, 2021, 42, 4007-4015.	0.9	16
7	A Novel Variant in Iranian Patient with Cystinuria: A Case Report. Iranian Journal of Public Health, 2021, 50, 1897-1901.	0.3	1
8	Secondary findings from whole-exome sequencing data in families with familial combined hyperlipidemia (FCHL). Egyptian Journal of Medical Human Genetics, 2021, 22, .	0.5	0
9	A novel variant in LPL gene is associated with familial combined hyperlipidemia. BioFactors, 2020, 46, 94-99.	2.6	10
10	A novel mutation in USF1 gene is associated with familial combined hyperlipidemia. IUBMB Life, 2020, 72, 616-623.	1.5	8
11	Molecular Pathways, Screening and Follow-up of Colorectal Carcinogenesis: An Overview. Current Cancer Therapy Reviews, 2020, 16, 88-96.	0.2	2
12	The atherogenic role of immune cells in familial hypercholesterolemia. IUBMB Life, 2020, 72, 782-789.	1.5	8
13	A novel variant in C5ORF42 gene is associated with Joubert syndrome. Molecular Biology Reports, 2020, 47, 4099-4103.	1.0	2
14	Role of long non-coding RNAs (LncRNAs) in multiple sclerosis: a brief review. Neurological Sciences, 2020, 41, 2443-2451.	0.9	7
15	Limb-girdle Muscular Dystrophy and Therapy: Insights into Cell and Gene-based Approaches. Current Gene Therapy, 2020, 19, 386-394.	0.9	5
16	MiR-492 as an Important Biomarker for Early Diagnosis and Targeted Treatment in Different Cancers. Current Cancer Therapy Reviews, 2020, 16, 269-275.	0.2	2
17	Familial combined hyperlipidemia: An overview of the underlying molecular mechanisms and therapeutic strategies. IUBMB Life, 2019, 71, 1221-1229.	1.5	34
18	CD73; a key ectonucleotidase in the development of breast cancer: Recent advances and perspectives. Journal of Cellular Physiology, 2019, 234, 14622-14632.	2.0	15

#	Article	IF	Citations
19	Involvement of aberrant regulation of epigenetic mechanisms in the pathogenesis of Parkinson's disease and epigeneticâ€based therapies. Journal of Cellular Physiology, 2019, 234, 19307-19319.	2.0	40
20	Molecular mechanisms, prevalence, and molecular methods for familial combined hyperlipidemia disease: A review. Journal of Cellular Biochemistry, 2019, 120, 8891-8898.	1.2	17
21	Cellular and Molecular Aspects of Parkinson Treatment: Future Therapeutic Perspectives. Molecular Neurobiology, 2019, 56, 4799-4811.	1.9	28
22	Prevalence, pathological mechanisms, and genetic basis of limbâ€girdle muscular dystrophies: A review. Journal of Cellular Physiology, 2019, 234, 7874-7884.	2.0	50
23	Macrophage: A Key Therapeutic Target in Atherosclerosis?. Current Pharmaceutical Design, 2019, 25, 3165-3174.	0.9	21
24	Roles of E6 and E7 Human Papillomavirus Proteins in Molecular Pathogenesis of Cervical Cancer. Current Protein and Peptide Science, 2019, 20, 926-934.	0.7	17
25	Limb-girdle Muscular Dystrophy with New Mutation in Sarcoglycan Beta Gene: A Case Report. Iranian Journal of Public Health, 2018, 47, 1953-1957.	0.3	3
26	Distribution of Human Papillomavirus Genotypes among Women in Mashhad, Iran. Intervirology, 2017, 60, 38-42.	1.2	11
27	SULF 1 gene polymorphism, rs6990375 is in significant association with fetus failure in IVF technique. Iranian Journal of Reproductive Medicine, 2015, 13, 215-20.	0.8	2