Ehsan Razmara

List of Publications by Year
in descending order

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39 papers	458 citations	949033 11 h-index	939365 18 g-index
40 all docs	40 docs citations	40 times ranked	550 citing authors

#	Article	IF	Citations
1	A novel missense variant in the LMNB2 gene causes progressive myoclonus epilepsy. Acta Neurologica Belgica, 2022, 122, 659-667.	0.5	6
2	Novel phenotype and genotype spectrum of NARS2 and literature review of previous mutations. Irish Journal of Medical Science, 2022, 191, 1877-1890.	0.8	13
3	SMAD4 contributes to chondrocyte and osteocyte development. Journal of Cellular and Molecular Medicine, 2022, 26, 1-15.	1.6	8
4	Association between TBXT rs2305089 polymorphism and chordoma in Iranian patients identified by a developed Tâ€ARMSâ€PCR assay. Journal of Clinical Laboratory Analysis, 2022, 36, e24150.	0.9	2
5	Novel phenotype and genotype spectrum of WDR62 in two patients with associated primary autosomal recessive microcephaly. Irish Journal of Medical Science, 2022, 191, 2733-2741.	0.8	2
6	Strategies to overcome the side effects of chimeric antigen receptor T cell therapy. Annals of the New York Academy of Sciences, 2022, 1510, 18-35.	1.8	3
7	KrÃ¼ppel-like factors in bone biology. Cellular Signalling, 2022, 93, 110308.	1.7	7
8	Docosahexaenoic acid reverses the promoting effects of breast tumor cell-derived exosomes on endothelial cell migration and angiogenesis. Life Sciences, 2021, 264, 118719.	2.0	25
9	The oncogenic and tumor suppressive roles of RNAâ€binding proteins in human cancers. Journal of Cellular Physiology, 2021, 236, 6200-6224.	2.0	17
10	Gravesâ€™ disease: introducing new genetic and epigenetic contributors. Journal of Molecular Endocrinology, 2021, 66, R33-R55.	1.1	21
11	A novel deletion variant in CLN3 with highly variable expressivity is responsible for juvenile neuronal ceroid lipofuscinoses. Acta Neurologica Belgica, 2021, 121, 737-748.	0.5	4
12	Novel manifestations of Warburg micro syndrome type 1 caused by a new splicing variant of RAB3GAP1: a case report. BMC Neurology, 2021, 21, 180.	0.8	3
13	How Transmembrane Inner Ear (TMIE) plays role in the auditory system: A mystery to us. Journal of Cellular and Molecular Medicine, 2021, 25, 5869-5883.	1.6	8
14	Functions of the SNAI family in chondrocyteâ€toâ€osteocyte development. Annals of the New York Academy of Sciences, 2021, 1503, 5-22.	1.8	12
15	Identification of a sixâ€microRNA signature as a potential diagnostic biomarker in breast cancer tissues. Journal of Clinical Laboratory Analysis, 2021, 35, e24010.	0.9	11
16	Prevalence and Genotype Distribution of Human Papillomavirus Infection among 12 076 Iranian Women. International Journal of Infectious Diseases, 2021, 111, 295-302.	1.5	7
17	Colorectal cancer cell-derived extracellular vesicles transfer miR-221-3p to promote endothelial cell angiogenesis via targeting suppressor of cytokine signaling 3. Life Sciences, 2021, 285, 119937.	2.0	25
18	Novel Homozygous Pathogenic Mutations of LAMA 2 Gene in Patients with Congen ital Muscular Dystrophy. Iranian Journal of Child Neurology, 2021, 15, 101-106.	0.2	0

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19	Upregulation of the long noncoding RNAs DSCAMâ€AS1 and MANCR is a potential diagnostic marker for breast carcinoma. Biotechnology and Applied Biochemistry, 2020, , .	1.4	11
20	A novel variant of ST3GAL3 causes nonâ€syndromic autosomal recessive intellectual disability in Iranian patients. Journal of Gene Medicine, 2020, 22, e3253.	1.4	9
21	Novel homozygous variants in the TMC1 and CDH23 genes cause autosomal recessive nonsyndromic hearing loss. Molecular Genetics & Genomic Medicine, 2020, 8, e1550.	0.6	5
22	Novel neuroclinical findings of autosomal recessive primary microcephaly 15 in a consanguineous Iranian family. European Journal of Medical Genetics, 2020, 63, 104096.	0.7	11
23	A panel of six-circulating miRNA signature in serum and its potential diagnostic value in colorectal cancer. Life Sciences, 2020, 258, 118226.	2.0	34
24	Advances of exosome isolation techniques in lung cancer. Molecular Biology Reports, 2020, 47, 7229-7251.	1.0	17
25	Novel imaging and clinical phenotypes of CONDSIAS disorder caused by a homozygous frameshift variant of ADPRHL2: a case report. BMC Neurology, 2020, 20, 291.	0.8	13
26	A homozygote variant in the tRNA splicing endonuclease subunit 54 causes pontocerebellar hypoplasia in a consanguineous Iranian family. Molecular Genetics & Genomic Medicine, 2020, 8, e1413.	0.6	7
27	Three Novel Variants identified in FBN1 and TGFBR2 in seven Iranian families with suspected Marfan syndrome. Molecular Genetics & Genomic Medicine, 2020, 8, e1274.	0.6	4
28	Non-coding RNAs underlying chemoresistance in gastric cancer. Cellular Oncology (Dordrecht), 2020, 43, 961-988.	2.1	29
29	Homozygous inâ€frame variant of SCL6A3 causes dopamine transporter deficiency syndrome in a consanguineous family. Annals of Human Genetics, 2020, 84, 315-323.	0.3	12
30	Wholeâ€exome sequencing identified a novel variant in an Iranian patient affected by pycnodysostosis. Molecular Genetics & Genomic Medicine, 2020, 8, e1118.	0.6	5
31	A novel missense variant in GPT2 causes non-syndromic autosomal recessive intellectual disability in a consanguineous Iranian family. European Journal of Medical Genetics, 2020, 63, 103853.	0.7	5
32	Identification of RELN variant p.(Ser2486Gly) in an Iranian family with ankylosing spondylitis; the first association of RELN and AS. European Journal of Human Genetics, 2020, 28, 754-762.	1.4	14
33	Functional Analysis of RELN S2486G Mutation and its Contribution to Pathogenesis of Ankylosing Spondylitis. Archives of Iranian Medicine, 2020, 23, 688-696.	0.2	3
34	Non-Coding RNAs in Cartilage Development: An Updated Review. International Journal of Molecular Sciences, 2019, 20, 4475.	1.8	53
35	S3440P Substitution in C-Terminal Region of Human Reelin Dramatically Impairs Secretion of Reelin from HEK 293T cells. Cellular and Molecular Biology, 2019, 65, 12-16.	0.3	7
36	S3440P Substitution in C-Terminal Region of Human Reelin Dramatically Impairs Secretion of Reelin from HEK 293T cells. Cellular and Molecular Biology, 2019, 65, 12-16.	0.3	5

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37	Whole-exome sequencing identifies R1279X of MYH6 gene to be associated with congenital heart disease. BMC Cardiovascular Disorders, 2018, 18, 137.	0.7	18
38	The expression analysis of Fra-1 gene and IL-11 protein in Iranian patients with ulcerative colitis. BMC Immunology, 2018, 19, 17.	0.9	15
39	The first case of NSHL by direct impression on gene and identification of one novel mutation in in the Iranian families. Iranian Journal of Basic Medical Sciences, 2018, 21, 333-341.	1.0	7