

M Anwar Iqbal

List of Publications by Year in descending order

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Version: 2024-02-01

9
papers

144
citations

1684188
5
h-index

1720034
7
g-index

9
all docs

9
docs citations

9
times ranked

396
citing authors

#	ARTICLE	IF	CITATIONS
1	Interstitial 20p13 microdeletion including <i>PRNP</i> and adjacent genes in a fetus with congenital abnormalities—First case report. <i>Clinical Case Reports (discontinued)</i> , 2021, 9, e04082.	0.5	0
2	<i>CHL1</i> deletion is associated with cognitive and language disabilities — Case report and review of literature. <i>Molecular Genetics & Genomic Medicine</i> , 2021, 9, e1725.	1.2	8
3	Allelic and dosage effects of NHS in X-linked cataract and Nance—Horan syndrome: a family study and literature review. <i>Molecular Cytogenetics</i> , 2021, 14, 48.	0.9	3
4	MEIS2 (15q14) gene deletions in siblings with mild developmental phenotypes and bifid uvula: documentation of mosaicism in an unaffected parent. <i>Molecular Cytogenetics</i> , 2021, 14, 58.	0.9	2
5	Novel 1.3 Mb germline duplication in chromosome 8q21.11 by microarray comparative genomic hybridization plus single nucleotide polymorphism analysis in an adult patient with pancytopenia and urinary bladder complications. <i>Clinical Case Reports (discontinued)</i> , 2018, 6, 1947-1952.	0.5	0
6	Assessing copy number aberrations and copy neutral loss of heterozygosity across the genome as best practice: An evidence based review of clinical utility from the cancer genomics consortium (CGC) working group for myelodysplastic syndrome, myelodysplastic/myeloproliferative and myeloproliferative neoplasms. <i>Cancer Genetics</i> , 2018, 228-229, 197-217.	0.4	25
7	Pathogenetics of alveolar capillary dysplasia with misalignment of pulmonary veins. <i>Human Genetics</i> , 2016, 135, 569-586.	3.8	85
8	A 0.7-Mb de novo duplication at 7q21.3 including the genes <i>DLX5</i> and <i>DLX6</i> in a patient with split-hand/split-foot malformation. <i>American Journal of Medical Genetics, Part A</i> , 2012, 158A, 3201-3206.	1.2	11
9	Interstitial del(20)(q11.2q12)—Clinical and molecular cytogenetic characterization. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 1880-1884.	1.2	10