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List of Publications by Year
in descending order

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9 papers	200 citations	1478505 6 h-index	1474206 9 g-index
9 all docs	9 docs citations	9 times ranked	322 citing authors

#	Article	IF	Citations
1	Yield of arrayâ€<scp>CGH</scp> analysis in Tunisian children with autism spectrum disorder. Molecular Genetics & Genomic Medicine, 2022, 10, .	1.2	8
2	FARPâ€1 deletion is associated with lack of response to autism treatment by early start denver model in a multiplex family. Molecular Genetics & Genomic Medicine, 2020, 8, e1373.	1.2	10
3	Appropriateness of arrayâ€CGH in the ADHD clinics: A comparative study. Genes, Brain and Behavior, 2020, 19, e12651.	2.2	4
4	An Interstitial 17q11.2 de novo Deletion Involving the CDK5R1 Gene in a High-Functioning Autistic Patient. Molecular Syndromology, 2018, 9, 247-252.	0.8	2
5	Functional characterization of CDK5 and CDK5R1 mutations identified in patients with non-syndromic intellectual disability. Journal of Human Genetics, 2016, 61, 283-293.	2.3	12
6	High frequency of copy number imbalances in Rubinsteinâ€“Taybi patients negative to CREBBP mutational analysis. European Journal of Human Genetics, 2010, 18, 768-775.	2.8	13
7	Premature chromatid separation is not a useful diagnostic marker for Cornelia de Lange syndrome. Chromosome Research, 2009, 17, 763-771.	2.2	49
8	High frequency of mosaic CREBBP deletions in Rubinsteinâ€“Taybi syndrome patients and mapping of somatic and germ-line breakpoints. Genomics, 2007, 90, 567-573.	2.9	42
9	Rubinstein-Taybi Syndrome: spectrum of CREBBP mutations in Italian patients. BMC Medical Genetics, 2006, 7, 77.	2.1	60