Christine Tyson

List of Publications by Year
in descending order

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8 papers	175 citations	1307594 7 h-index	1588992 8 g-index
8 all docs	8 docs citations	8 times ranked	500 citing authors

#	Article	IF	Citations
1	A distinct neurodevelopmental syndrome with intellectual disability, autism spectrum disorder, characteristic facies, and macrocephaly is caused by defects in CHD8. Journal of Human Genetics, 2019, 64, 271-280.	2.3	35
2	Copy number variation of the REXO1L1 gene cluster; euchromatic deletion variant or susceptibility factor?. European Journal of Human Genetics, 2017, 25, 8-9.	2.8	1
3	Expansion of a 12-kb VNTR containing the REXO1L1 gene cluster underlies the microscopically visible euchromatic variant of 8q21.2. European Journal of Human Genetics, 2014, 22, 458-463.	2.8	10
4	Population-based characterization of the genetic landscape of chronic lymphocytic leukemia patients referred for cytogenetic testing in British Columbia, Canada: the role of provincial laboratory standardization. Cancer Genetics, 2014, 207, 316-325.	0.4	13
5	Genotypeâ€“phenotype analysis of 18q12.1-q12.2 copy number variation in autism. European Journal of Medical Genetics, 2013, 56, 420-425.	1.3	10
6	Understanding the impact of 1q21.1 copy number variant. Orphanet Journal of Rare Diseases, 2011, 6, 54.	2.7	48
7	Submicroscopic deletions of 11q24-25 in individuals without Jacobsen syndrome: re-examination of the critical region by high-resolution array-CGH. Molecular Cytogenetics, 2008, 1, 23.	0.9	25
8	Elucidation of a cryptic interstitial 7q31.3 deletion in a patient with a language disorder and mild mental retardation by array-CGH. American Journal of Medical Genetics Part A, 2004, 129A, 254-260.	2.4	33