Marsha D Speevak

List of Publications by Year
in descending order

16	338	1039880 9	940416 16
papers	citations	h-index	g-index

16	16	16	529
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	OTUD7A Regulates Neurodevelopmental Phenotypes in the 15q13.3 Microdeletion Syndrome. American Journal of Human Genetics, 2018, 102, 278-295.	2.6	81
2	Detection of submicroscopic aberrations in patients with unexplained mental retardation by fluorescence in situ hybridization using multiple subtelomeric probes. Genetics in Medicine, 2001, 3, 416-421.	1.1	49
3	Nonâ€syndromic language delay in a child with disruption in the Protocadherin11X/Y gene pair. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2011, 156, 484-489.	1.1	40
4	Stable Transfer of Zebrafish Chromosome Segments into Mouse Cells. Genomics, 1996, 33, 57-64.	1.3	36
5	Mosaic r(13) in an infant with aprosencephaly. American Journal of Medical Genetics Part A, 1993, 47, 531-533.	2.4	24
6	Midtrimester genetic amniocentesis in eastern Ontario: a review from 1970 to 1985.. Journal of Medical Genetics, 1987, 24, 335-343.	1.5	19
7	Identification of the origin of double minutes in normal human cells by laser-based chromosome microdissection approach. Human Genetics, 1995, 96, 39-43.	1.8	18
8	The detection of chromosome anomalies by QFâ€PCR and residual risks as compared to Gâ€banded analysis. Prenatal Diagnosis, 2011, 31, 454-458.	1.1	14
9	Construction and analysis of microcell hybrids containing dual selectable tagged human chromosomes. Cytogenetic and Genome Research, 1995, 69, 63-65.	0.6	11
10	Charcotâ€“Marieâ€“Tooth 1B caused by expansion of a familial myelin protein zero (MPZ) gene duplication. European Journal of Medical Genetics, 2013, 56, 566-569.	0.7	10
11	Alternatively spliced, truncated human BRCA2 isoforms contain a novel coding exon. European Journal of Human Genetics, 2003, 11, 951-954.	1.4	9
12	Human Chromosome 3 Mediates Growth Arrest and Suppression of Apoptosis in Microcell Hybrids. Molecular and Cellular Biology, 1996, 16, 2214-2225.	1.1	6
13	A novel complex mutation in MSH2 contributes to both Muir-Torre and Lynch Syndrome. Journal of Human Genetics, 2010, 55, 37-41.	1.1	6
14	Prenatal genomic microarray and sequencing in Canadian medical practice: towards consensus. Journal of Medical Genetics, 2015, 52, 585-586.	1.5	6
15	Alpha-1-antitrypsin molecular testing in Canada: A seven year, multi-centre comparison. Clinical Biochemistry, 2020, 81, 27-33.	0.8	5
16	Identification of chromosomes implicated in suppression of apoptosis in somatic cell hybrids. Biochemistry and Cell Biology, 1994, 72, 655-662.	0.9	4