Bridget Fernandez

List of Publications by Year
in descending order

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12 papers	3,870 citations	759233 12 h-index	1199594 12 g-index
12 all docs	12 docs citations	12 times ranked	6060 citing authors

#	Article	IF	Citations
1	Structural Variation of Chromosomes in Autism Spectrum Disorder. American Journal of Human Genetics, 2008, 82, 477-488.	6.2	1,641
2	Reduction in Neural-Tube Defects after Folic Acid Fortification in Canada. New England Journal of Medicine, 2007, 357, 135-142.	27.0	693
3	Contribution of SHANK3 Mutations to Autism Spectrum Disorder. American Journal of Human Genetics, 2007, 81, 1289-1297.	6.2	604
4	Founder and Recurrent CDH1 Mutations in Families With Hereditary Diffuse Gastric Cancer. JAMA - Journal of the American Medical Association, 2007, 297, 2360.	7.4	394
5	Disruption at the PTCHD1 Locus on Xp22.11 in Autism Spectrum Disorder and Intellectual Disability. Science Translational Medicine, 2010, 2, 49ra68.	12.4	178
6	A common X-linked inborn error of carnitine biosynthesis may be a risk factor for nondysmorphic autism. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 7974-7981.	7.1	118
7	Spina bifida before and after folic acid fortification in Canada. Birth Defects Research Part A: Clinical and Molecular Teratology, 2008, 82, 622-626.	1.6	80
8	Cerebellar and posterior fossa malformations in patients with autismâ€associated chromosome 22q13 terminal deletion. American Journal of Medical Genetics, Part A, 2013, 161, 131-136.	1.2	65
9	Profound, prelingual nonsyndromic deafness maps to chromosome 10q21 and is caused by a novel missense mutation in the Usher syndrome type IF gene PCDH15. European Journal of Human Genetics, 2009, 17, 554-564.	2.8	33
10	A molecular genetic study of autism and related phenotypes in extended pedigrees. Journal of Neurodevelopmental Disorders, 2013, 5, 30.	3.1	23
11	Impact of folic acid food fortification on the birth prevalence of lipomyelomeningocele in Canada. Birth Defects Research Part A: Clinical and Molecular Teratology, 2008, 82, 106-109.	1.6	22
12	Copy number variation analysis and sequencing of the X-linked mental retardation gene TSPAN7/TM4SF2 in patients with autism spectrum disorder. Psychiatric Genetics, 2009, 19, 154-155.	1.1	19