Bridget Fernandez

List of Publications by Year
in descending order

12 papers	3,870 citations	858243 12 h-index	1336881 12 g-index
12 all docs	12 docs citations	12 times ranked	6671 citing authors

#	Article	IF	Citations
1	A molecular genetic study of autism and related phenotypes in extended pedigrees. Journal of Neurodevelopmental Disorders, 2013, 5, 30.	1.5	23
2	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.	0.7	65
3	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.	3.3	118
4	Disruption at the PTCHD1 Locus on Xp22.11 in Autism Spectrum Disorder and Intellectual Disability. Science Translational Medicine, 2010, 2, 49ra68.	5.8	178
5	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.	1.4	33
6	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.	0.6	19
7	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
8	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
9	Structural Variation of Chromosomes in Autism Spectrum Disorder. American Journal of Human Genetics, 2008, 82, 477-488.	2.6	1,641
10	Reduction in Neural-Tube Defects after Folic Acid Fortification in Canada. New England Journal of Medicine, 2007, 357, 135-142.	13.9	693
11	Founder and Recurrent CDH1 Mutations in Families With Hereditary Diffuse Gastric Cancer. JAMA - Journal of the American Medical Association, 2007, 297, 2360.	3.8	394
12	Contribution of SHANK3 Mutations to Autism Spectrum Disorder. American Journal of Human Genetics, 2007, 81, 1289-1297.	2.6	604