

# Mutations in the gene encoding the synaptic scaffolding with autism spectrum disorders

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Citation Report

#	ARTICLE	IF	CITATIONS
2	Autism: the quest for the genes. <i>Expert Reviews in Molecular Medicine</i> , 2007, 9, 1-15.	1.6	44
3	A Neuroligin-3 Mutation Implicated in Autism Increases Inhibitory Synaptic Transmission in Mice. <i>Science</i> , 2007, 318, 71-76.	6.0	932
4	Molecular Windows into Speech and Language Disorders. <i>Folia Phoniatica Et Logopaedica</i> , 2007, 59, 130-140.	0.5	16
5	Ethical implications of array comparative genomic hybridization in complex phenotypes: points to consider in research. <i>Genetics in Medicine</i> , 2007, 9, 626-631.	1.1	43
6	The Possible Interplay of Synaptic and Clock Genes in Autism Spectrum Disorders. <i>Cold Spring Harbor Symposia on Quantitative Biology</i> , 2007, 72, 645-654.	2.0	161
7	Co-inheritance of a novel deletion of the entire SPINK1 gene with a CFTR missense mutation (L997F) in a family with chronic pancreatitis. <i>Molecular Genetics and Metabolism</i> , 2007, 92, 168-175.	0.5	25
8	MeCP2 Controls Excitatory Synaptic Strength by Regulating Glutamatergic Synapse Number. <i>Neuron</i> , 2007, 56, 58-65.	3.8	439
9	Structures of Neuroligin-1 and the Neuroligin-1/Neurexin-1 <sup>β</sup> Complex Reveal Specific Protein-Protein and Protein-Ca <sup>2+</sup> Interactions. <i>Neuron</i> , 2007, 56, 992-1003.	3.8	178
10	Use of array CGH in the evaluation of dysmorphology, malformations, developmental delay, and idiopathic mental retardation. <i>Current Opinion in Genetics and Development</i> , 2007, 17, 182-192.	1.5	293
11	Contribution of SHANK3 Mutations to Autism Spectrum Disorder. <i>American Journal of Human Genetics</i> , 2007, 81, 1289-1297.	2.6	604
12	Strong Association of De Novo Copy Number Mutations with Autism. <i>Science</i> , 2007, 316, 445-449.	6.0	2,497
13	Atypical Antipsychotics in Children with Pervasive Developmental Disorders. <i>Paediatric Drugs</i> , 2007, 9, 249-266.	1.3	26
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15	22q13 microduplication in two patients with common clinical manifestations: A recognizable syndrome?. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 2804-2809.	0.7	40
16	Association of autism with polymorphisms in the paired-like homeodomain transcription factor 1 (PITX1) on chromosome 5q31: a candidate gene analysis. <i>BMC Medical Genetics</i> , 2007, 8, 74.	2.1	35
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19	Autism: highly heritable but not inherited. <i>Nature Medicine</i> , 2007, 13, 534-536.	15.2	83
20	The bridge between dendritic cells and asthma. <i>Nature Medicine</i> , 2007, 13, 536-538.	15.2	12

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22	The Neurobiology of Autism. <i>Brain Pathology</i> , 2007, 17, 434-447.	2.1	373
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32	Autism genetics: strategies, challenges, and opportunities. <i>Autism Research</i> , 2008, 1, 4-17.	2.1	123
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43	Deletion 22q13.3 syndrome. <i>Orphanet Journal of Rare Diseases</i> , 2008, 3, 14.	1.2	161
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58	A complete theory of psychosis and autism as diametric disorders of social brain must consider full range of clinical syndromes. <i>Behavioral and Brain Sciences</i> , 2008, 31, 277-278.	0.4	2
59	Is this conjectural phenotypic dichotomy a plausible outcome of genomic imprinting?. <i>Behavioral and Brain Sciences</i> , 2008, 31, 267-268.	0.4	2

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71	Cortical plasticity: A proposed mechanism by which genomic factors lead to the behavioral and neurological phenotype of autism spectrum and psychotic-spectrum disorders. Behavioral and Brain Sciences, 2008, 31, 276-277.	0.4	9
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