Raphael Bernier

List of Publications by Year
in descending order

151	16,497	31976 53	17592 121
papers	citations	h-index	g-index

161	161	161	18991
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Sex Differences in Autism: Examining Intrinsic and Extrinsic Factors in Children and Adolescents Enrolled in a National ASD Cohort. Journal of Autism and Developmental Disorders, 2023, 53, 1305-1318.	2.7	15
2	The Early Start Denver Model Intervention and Mu Rhythm Attenuation in Autism Spectrum Disorders. Journal of Autism and Developmental Disorders, 2022, 52, 3304-3313.	2.7	7
3	Use of a Best Practice Alert (BPA) to Increase Diversity Within a US-Based Autism Research Cohort. Journal of Autism and Developmental Disorders, 2022, , 1.	2.7	3
4	Utilization of a Best Practice Alert (BPA) at Point-of-Care for Recruitment into a US-Based Autism Research Study. Journal of Autism and Developmental Disorders, 2022, , 1.	2.7	4
5	The Autism Biomarkers Consortium for Clinical Trials: evaluation of a battery of candidate eye-tracking biomarkers for use in autism clinical trials. Molecular Autism, 2022, 13, 15.	4.9	28
6	Patterns of intervention utilization among school-aged children on the autism spectrum: Findings from a multi-site research consortium. Research in Autism Spectrum Disorders, 2022, 94, 101950.	1.5	5
7	Do Biological Sex and Early Developmental Milestones Predict the Age of First Concerns and Eventual Diagnosis in Autism Spectrum Disorder?. Autism Research, 2021, 14, 156-168.	3.8	21
8	Neurophysiology. , 2021, , 3144-3152.		0
9	A Genetics-First Approach to Dissecting the Heterogeneity of Autism: Phenotypic Comparison of Autism Risk Copy Number Variants. American Journal of Psychiatry, 2021, 178, 77-86.	7.2	62
10	Understanding Vaccine Hesitancy Among Parents of Children With Autism Spectrum Disorder and Parents of Children With Non-Autism Developmental Delays. Journal of Child Neurology, 2021, 36, 911-918.	1.4	20
11	Modeling temporal dynamics of face processing in youth and adults. Social Neuroscience, 2021, 16, 345-361.	1.3	3
12	Differences in the number of de novo mutations between individuals are due to small family-specific effects and stochasticity. Genome Research, 2021, 31, 1513-1518.	5.5	6
13	Resting state EEG in youth with ASD: age, sex, and relation to phenotype. Journal of Neurodevelopmental Disorders, 2021, 13, 33.	3.1	22
14	Parental Beliefs About Causes of Autism Spectrum Disorder: An Investigation of a Research Measure Using Principal Component Analysis. Research in Autism Spectrum Disorders, 2021, 87, 101825.	1.5	3
15	Face Perception. , 2021, , 1937-1944.		1
16	Saliva RNA Biomarkers of Gastrointestinal Dysfunction in Children With Autism and Neurodevelopmental Disorders: Potential Implications for Precision Medicine. Frontiers in Psychiatry, 2021, 12, 824933.	2.6	4
17	Neurocognitive and Neurobehavioral Phenotype of Youth with Schaaf-Yang Syndrome. Journal of Autism and Developmental Disorders, 2020, 50, 2491-2500.	2.7	10
18	Language characterization in 16p11.2 deletion and duplication syndromes. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2020, 183, 380-391.	1.7	16

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19	Beliefs about causes of autism and vaccine hesitancy among parents of children with autism spectrum disorder. Vaccine, 2020, 38, 6327-6333.	3.8	19
20	Transcriptional subtyping explains phenotypic variability in genetic subtypes of autism spectrum disorder. Development and Psychopathology, 2020, 32, 1353-1361.	2.3	3
21	Vaccine Hesitancy and Attributions for Autism among Racially and Ethnically Diverse Groups of Parents of Children with Autism Spectrum Disorder: A Pilot Study. Autism Research, 2020, 13, 1790-1796.	3.8	9
22	Health-related issues in youth with autism spectrum disorder. Children's Health Care, 2020, 49, 355-360.	0.9	0
23	Insufficient Evidence for â€œAutism-Specificâ€•Genes. American Journal of Human Genetics, 2020, 106, 587-595.	6.2	110
24	Vaccine hesitancy and illness perceptions: comparing parents of children with autism spectrum disorder to other parent groups. Children's Health Care, 2020, 49, 385-402.	0.9	10
25	Early infantile epileptic encephalopathy due to biallelic pathogenic variants in <scp>PIGQ</scp>: Report of seven new subjects and review of the literature. Journal of Inherited Metabolic Disease, 2020, 43, 1321-1332.	3.6	15
26	Evaluating heterogeneity in <scp>ASD</scp> symptomatology, cognitive ability, and adaptive functioning among 16p11.2 <scp>CNV</scp> carriers. Autism Research, 2020, 13, 1300-1310.	3.8	23
27	BAZ2B haploinsufficiency as a cause of developmental delay, intellectual disability, and autism spectrum disorder. Human Mutation, 2020, 41, 921-925.	2.5	11
28	Beliefs in vaccine as causes of autism among SPARK cohort caregivers. Vaccine, 2020, 38, 1794-1803.	3.8	12
29	Person Ability Scores as an Alternative to Norm-Referenced Scores as Outcome Measures in Studies of Neurodevelopmental Disorders. American Journal on Intellectual and Developmental Disabilities, 2020, 125, 475-480.	1.6	30
30	Evaluating Two Common Strategies for Research Participant Recruitment Into Autism Studies: Observational Study. Journal of Medical Internet Research, 2020, 22, e16752.	4.3	10
31	Parent-led, stepped-care cognitive-behavioral therapy for youth with autism and co-occurring anxiety: study rationale and method. Revista Brasileira De Psiquiatria, 2020, 42, 638-645.	1.7	0
32	Screening and Referral Practices for Autism Spectrum Disorder in Primary Pediatric Care. , 2020, , 92-100.		0
33	Towards a consensus on developmental regression. Neuroscience and Biobehavioral Reviews, 2019, 107, 3-5.	6.1	14
34	The fetal origins of mental illness. American Journal of Obstetrics and Gynecology, 2019, 221, 549-562.	1.3	190
35	On measuring regression in autism spectrum disorder: A commentary. Neuroscience and Biobehavioral Reviews, 2019, 104, 116-117.	6.1	0
36	Screening and Referral Practices for Autism Spectrum Disorder in Primary Pediatric Care. Pediatrics, 2019, 144, e20183326.	2.1	58

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37	Psychiatric disorders in children with 16p11.2 deletion and duplication. Translational Psychiatry, 2019, 9, 8.	4.8	93
38	Family-based exome sequencing and case-control analysis implicate CEP41 as an ASD gene. Translational Psychiatry, 2019, 9, 4.	4.8	13
39	Phenotypic and biochemical analysis of an international cohort of individuals with variants in NAA10 and NAA15. Human Molecular Genetics, 2019, 28, 2900-2919.	2.9	46
40	Trauma and Autism Spectrum Disorder: Review, Proposed Treatment Adaptations and Future Directions. Journal of Child and Adolescent Trauma, 2019, 12, 529-547.	1.9	48
41	It takes two! Exploring sex differences in parenting neurobiology and behaviour. Journal of Neuroendocrinology, 2019, 31, e12721.	2.6	27
42	Using latent class analysis to identify treatmentâ€use subgroups among parents of children with autism spectrum disorder. Autism Research, 2019, 12, 843-854.	3.8	2
43	Long-term Risk of Neuropsychiatric Disease After Exposure to Infection In Utero. JAMA Psychiatry, 2019, 76, 594.	11.0	180
44	Side Effects and Behavioral Outcomes Following High-Dose Carnitine Supplementation Among Young Males With Autism Spectrum Disorder: A Pilot Study. Global Pediatric Health, 2019, 6, 2333794X1983069.	0.7	12
45	Clinical Presentation of a Complex Neurodevelopmental Disorder Caused by Mutations in ADNP. Biological Psychiatry, 2019, 85, 287-297.	1.3	108
46	SPARK: A US Cohort of 50,000 Families to Accelerate Autism Research. Neuron, 2018, 97, 488-493.	8.1	265
47	Using Teacher Ratings to Investigate Developmental Regression as a Potential Indicator of School-Age Symptoms in Students with Autism Spectrum Disorder. School Mental Health, 2018, 10, 77-90.	2.1	4
48	Visual systemizing preference in children with autism: A randomized controlled trial of intranasal oxytocin. Development and Psychopathology, 2018, 30, 511-521.	2.3	14
49	The diagnosis conundrum: Comparison of crowdsourced and expert assessments of toddlers with high and low risk of autism spectrum disorder. Autism Research, 2018, 11, 1629-1634.	3.8	4
50	Autism Treatment: Familiesâ€™ Use Varies Across U.S. Regions. Journal of Disability Policy Studies, 2018, 29, 97-107.	1.5	6
51	Weaker neural suppression in autism spectrum disorder. Journal of Vision, 2018, 18, 548.	0.3	1
52	Hyper-upregulation of abnormally low neural response along the visual pathway in autism. Journal of Vision, 2018, 18, 344.	0.3	0
53	Parent Perceptions About Autism Spectrum Disorder Influence Treatment Choices. Focus on Autism and Other Developmental Disabilities, 2017, 32, 305-318.	1.3	33
54	Next Generation Sequencing Mitochondrial DNA Analysis in Autism Spectrum Disorder. Autism Research, 2017, 10, 1338-1343.	3.8	31

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55	Developmental trajectories for young children with 16p11.2 copy number variation. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2017, 174, 367-380.	1.7	42
56	Intellectual Functioning and Autism Spectrum Disorder: Can Profiles Inform Identification of Subpopulations?. Review Journal of Autism and Developmental Disorders, 2017, 4, 339-349.	3.4	3
57	Exonic Mosaic Mutations Contribute Risk for Autism Spectrum Disorder. American Journal of Human Genetics, 2017, 101, 369-390.	6.2	151
58	Gene Disrupting Mutations Associated with Regression in Autism Spectrum Disorder. Journal of Autism and Developmental Disorders, 2017, 47, 3600-3607.	2.7	26
59	Interactive Effects of Prenatal Antidepressant Exposure and Likely Gene Disrupting Mutations on the Severity of Autism Spectrum Disorder. Journal of Autism and Developmental Disorders, 2017, 47, 3489-3496.	2.7	9
60	The Cognitive and Behavioral Phenotypes of Individuals with CHRNA7 Duplications. Journal of Autism and Developmental Disorders, 2017, 47, 549-562.	2.7	68
61	Severity of ASD symptoms and their correlation with the presence of copy number variations and exposure to first trimester ultrasound. Autism Research, 2017, 10, 472-484.	3.8	22
62	Early Intervention Before Autism Diagnosis in Children Referred to a Regional Autism Clinic. Journal of Developmental and Behavioral Pediatrics, 2016, 37, 15-19.	1.1	26
63	Autism Spectrum Disorder, Developmental and Psychiatric Features in 16p11.2 Duplication. Journal of Autism and Developmental Disorders, 2016, 46, 2734-2748.	2.7	47
64	Executive function predicts the development of play skills for verbal preschoolers with autism spectrum disorders. Autism Research, 2016, 9, 1274-1284.	3.8	27
65	The Number of Genomic Copies at the 16p11.2 Locus Modulates Language, Verbal Memory, and Inhibition. Biological Psychiatry, 2016, 80, 129-139.	1.3	78
66	De novo loss-of-function mutations in WAC cause a recognizable intellectual disability syndrome and learning deficits in Drosophila. European Journal of Human Genetics, 2016, 24, 1145-1153.	2.8	34
67	Musings on the puzzle piece. Autism, 2016, 20, 250-250.	4.1	1
68	Parental report of vaccine receipt in children with autism spectrum disorder: Do rates differ by pattern of ASD onset?. Vaccine, 2016, 34, 1335-1342.	3.8	10
69	Brief Report: Further Examination of Self-Injurious Behaviors in Children and Adolescents with Autism Spectrum Disorders. Journal of Autism and Developmental Disorders, 2016, 46, 1872-1879.	2.7	21
70	The complex behavioral phenotype of 15q13.3 microdeletion syndrome. Genetics in Medicine, 2016, 18, 1111-1118.	2.4	45
71	Maternal Modifiers and Parent-of-Origin Bias of the Autism-Associated 16p11.2 CNV. American Journal of Human Genetics, 2016, 98, 45-57.	6.2	55
72	Defining the Effect of the 16p11.2 Duplication on Cognition, Behavior, and Medical Comorbidities. JAMA Psychiatry, 2016, 73, 20.	11.0	195

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73	Clinical phenotype of the recurrent 1q21.1 copy-number variant. Genetics in Medicine, 2016, 18, 341-349.	2.4	134
74	Age-related trends in treatment use for children with autism spectrum disorder. Research in Autism Spectrum Disorders, 2015, 15-16, 29-41.	1.5	33
75	A computer-generated animated face stimulus set for psychophysiological research. Behavior Research Methods, 2015, 47, 562-570.	4.0	17
76	Whole exome sequencing in extended families with autism spectrum disorder implicates four candidate genes. Human Genetics, 2015, 134, 1055-1068.	3.8	49
77	Cognitive Profiles in Youth with Autism Spectrum Disorder: An Investigation of Base Rate Discrepancies using the Differential Ability Scalesâ€"Second Edition. Journal of Autism and Developmental Disorders, 2015, 45, 1978-1988.	2.7	26
78	The Role of Parental Cognitive, Behavioral, and Motor Profiles in Clinical Variability in Individuals With Chromosome 16p11.2 Deletions. JAMA Psychiatry, 2015, 72, 119.	11.0	112
79	Excess of rare, inherited truncating mutations in autism. Nature Genetics, 2015, 47, 582-588.	21.4	531
80	Self-motion perception in autism is compromised by visual noise but integrated optimally across multiple senses. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 6461-6466.	7.1	69
81	Epigenetics of Autism-related Impairment. Journal of Developmental and Behavioral Pediatrics, 2015, 36, 61-67.	1.1	83
82	Emergence of Autism Spectrum Disorder in Children from Simplex Families: Relations to Parental Perceptions of Etiology. Journal of Autism and Developmental Disorders, 2015, 45, 1451-1463.	2.7	28
83	The Cognitive and Behavioral Phenotype of the 16p11.2 Deletion in a Clinically Ascertained Population. Biological Psychiatry, 2015, 77, 785-793.	1.3	198
84	Guidelines and Best Practices for Electrophysiological Data Collection, Analysis and Reporting in Autism. Journal of Autism and Developmental Disorders, 2015, 45, 425-443.	2.7	75
85	A Multi-Rater Study on Being Teased Among Children/Adolescents With Autism Spectrum Disorder (ASD) and Their Typically Developing Siblings. Focus on Autism and Other Developmental Disabilities, 2014, 29, 195-205.	1.3	30
86	Nonverbal and Verbal Cognitive Discrepancy Profiles in Autism Spectrum Disorders: Influence of Age and Gender. American Journal on Intellectual and Developmental Disabilities, 2014, 119, 84-99.	1.6	64
87	The transcriptional regulator ADNP links the BAF (SWI/SNF) complexes with autism. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2014, 166, 315-326.	1.6	68
88	Psychotropic medication use among children with autism spectrum disorders within the Simons Simplex Collection: Are core features of autism spectrum disorder related?. Autism, 2014, 18, 933-942.	4.1	24
89	Update on diagnostic classification in autism. Current Opinion in Psychiatry, 2014, 27, 105-109.	6.3	46
90	The Autism Simplex Collection: an international, expertly phenotyped autism sample for genetic and phenotypic analyses. Molecular Autism, 2014, 5, 34.	4.9	31

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91	Expression of the Broad Autism Phenotype in Simplex Autism Families from the Simons Simplex Collection. Journal of Autism and Developmental Disorders, 2014, 44, 2392-2399.	2.7	31
92	A Genotype-First Approach to Defining the Subtypes of a Complex Disease. Cell, 2014, 156, 872-877.	28.9	195
93	Convergence of Genes and Cellular Pathways Dysregulated in Autism Spectrum Disorders. American Journal of Human Genetics, 2014, 94, 677-694.	6.2	819
94	Disruptive CHD8 Mutations Define a Subtype of Autism Early in Development. Cell, 2014, 158, 263-276.	28.9	637
95	Developmental regression among children with autism spectrum disorder: Onset, duration, and effects on functional outcomes. Research in Autism Spectrum Disorders, 2014, 8, 890-898.	1.5	24
96	No increase in autism-associated genetic events in children conceived by assisted reproduction. Fertility and Sterility, 2014, 102, 388-393.	1.0	10
97	EEG Mu Rhythm in Typical and Atypical Development. Journal of Visualized Experiments, 2014, , .	0.3	2
98	Assessment of the Core Features of ASD. Autism and Child Psychopathology Series, 2014, , 65-86.	0.2	3
99	Negative Priming Effect. , 2013, , 1987-1987.		0
100	The role of imitation in the observed heterogeneity in EEG mu rhythm in autism and typical development. Brain and Cognition, 2013, 82, 69-75.	1.8	82
101	Behavioral Approaches. , 2013, , 430-430.		0
102	Mental Retardation (Former Term). , 2013, , 1841-1841.		0
103	Natural Language Paradigm. , 2013, , 1976-1978.		0
104	Developmental changes in mu suppression to observed and executed actions in autism spectrum disorders. Social Cognitive and Affective Neuroscience, 2013, 8, 300-304.	3.0	84
105	TM4SF20 Ancestral Deletion and Susceptibility to a Pediatric Disorder of Early Language Delay and Cerebral White Matter Hyperintensities. American Journal of Human Genetics, 2013, 93, 197-210.	6.2	43
106	The Broader Autism Phenotype in Simplex and Multiplex Families. Journal of Autism and Developmental Disorders, 2013, 43, 1597-1605.	2.7	52
107	Refinement and Discovery of New Hotspots of Copy-Number Variation Associated with Autism Spectrum Disorder. American Journal of Human Genetics, 2013, 92, 221-237.	6.2	279
108	A quarter century of progress on the early detection and treatment of autism spectrum disorder. Development and Psychopathology, 2013, 25, 1455-1472.	2.3	117

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109	Behavior Summarized Evaluation-Revised (BSE-R). , 2013, , 423-427.		0
110	Behavior Analysis. , 2013, , 405-409.		0
111	Mutual Gaze. , 2013, , 1966-1967.		0
112	â€œWhat Do You Like/Dislike About the Treatments Youâ€™re Currently Using?â€• Focus on Autism and Other Developmental Disabilities, 2012, 27, 51-60.	1.3	92
113	A 600â€…kb deletion syndrome at 16p11.2 leads to energy imbalance and neuropsychiatric disorders. Journal of Medical Genetics, 2012, 49, 660-668.	3.2	251
114	A Multisite Study of the Clinical Diagnosis of Different Autism Spectrum Disorders. Archives of General Psychiatry, 2012, 69, 306.	12.3	385
115	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
116	Multiplex Targeted Sequencing Identifies Recurrently Mutated Genes in Autism Spectrum Disorders. Science, 2012, 338, 1619-1622.	12.6	1,133
117	Sporadic autism exomes reveal a highly interconnected protein network of de novo mutations. Nature, 2012, 485, 246-250.	27.8	1,960
118	Relationship between the social functioning of children with autism spectrum disorders and their siblingsâ€™ competencies/problem behaviors. Research in Autism Spectrum Disorders, 2012, 6, 646-653.	1.5	18
119	Social attention: a possible early indicator of efficacy in autism clinical trials. Journal of Neurodevelopmental Disorders, 2012, 4, 11.	3.1	115
120	Emotional and Behavioral Adjustment in Typically Developing Siblings of Children with Autism Spectrum Disorders. Journal of Autism and Developmental Disorders, 2012, 42, 1393-1402.	2.7	36
121	Utility of the Social Communication Questionnaire-Current and Social Responsiveness Scale as Teacher-Report Screening Tools for Autism Spectrum Disorders. Journal of Autism and Developmental Disorders, 2012, 42, 1705-1716.	2.7	73
122	Evidence for involvement of GNB1L in autism. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 61-71.	1.7	28
123	Evidence for broader autism phenotype characteristics in parents from multipleâ€incidence autism families. Autism Research, 2012, 5, 13-20.	3.8	76
124	Exome sequencing in sporadic autism spectrum disorders identifies severe de novo mutations. Nature Genetics, 2011, 43, 585-589.	21.4	1,080
125	Expanding the clinical spectrum of the 16p11.2 chromosomal rearrangements: three patients with syringomyelia. European Journal of Human Genetics, 2011, 19, 152-156.	2.8	47
126	Genome-scan for IQ discrepancy in autism: evidence for loci on chromosomes 10 and 16. Human Genetics, 2011, 129, 59-70.	3.8	44

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127	The Broader Autism Phenotype and Its Implications on the Etiology and Treatment of Autism Spectrum Disorders. Autism Research & Treatment, 2011, 2011, 1-19.	0.5	103
128	Relative Burden of Large CNVs on a Range of Neurodevelopmental Phenotypes. PLoS Genetics, 2011, 7, e1002334.	3.5	293
129	Some, but not complete, reassurance on the safety of folic acid fortification. American Journal of Clinical Nutrition, 2010, 92, 1287-1288.	4.7	3
130	Psychopathology, Families, and Culture: Autism. Child and Adolescent Psychiatric Clinics of North America, 2010, 19, 855-867.	1.9	116
131	Neurobiological correlates of social functioning in autism. Clinical Psychology Review, 2010, 30, 733-748.	11.4	103
132	Autism genome-wide copy number variation reveals ubiquitin and neuronal genes. Nature, 2009, 459, 569-573.	27.8	1,270
133	Common genetic variants on 5p14.1 associate with autism spectrum disorders. Nature, 2009, 459, 528-533.	27.8	912
134	The MTHFR 677Câ†'T polymorphism and behaviors in children with autism: exploratory genotypeâ€"phenotype correlations. Autism Research, 2009, 2, 98-108.	3.8	57
135	Parental reports on the efficacy of treatments and therapies for their children with autism spectrum disorders. Research in Autism Spectrum Disorders, 2009, 3, 528-537.	1.5	97
136	â€œMy greatest joy and my greatest heart ache:â€•Parentsâ€™ own words on how having a child in the autism spectrum has affected their lives and their familiesâ€™ lives. Research in Autism Spectrum Disorders, 2009, 3, 670-684.	1.5	285
137	Level of Functioning in Autism Spectrum Disorders: Phenotypic Congruence Among Affected Siblings. Journal of Autism and Developmental Disorders, 2008, 38, 1019-1027.	2.7	17
138	Genetic and environmental influences on symptom domains in twins and siblings with autism. Research in Autism Spectrum Disorders, 2008, 2, 320-331.	1.5	29
139	Parental reports on the prevalence of co-occurring intellectual disability among children with autism spectrum disorders. Research in Autism Spectrum Disorders, 2008, 2, 546-556.	1.5	12
140	Screening Cases Within a Statewide Autism Registry. Focus on Autism and Other Developmental Disabilities, 2008, 23, 148-154.	1.3	3
141	Lack of evidence for increased genetic loading for autism among families of affected females. Autism, 2007, 11, 279-286.	4.1	43
142	Parental reports on the use of treatments and therapies for children with autism spectrum disorders. Research in Autism Spectrum Disorders, 2007, 1, 195-209.	1.5	192
143	Early Responsiveness to Intensive Behavioural Intervention Predicts Outcomes Among Preschool Children with Autism. International Journal of Disability Development and Education, 2007, 54, 151-175.	1.1	20
144	Quantitative Assessment of Autism Symptom-related Traits in Probands and Parents: Broader Phenotype Autism Symptom Scale. Journal of Autism and Developmental Disorders, 2007, 37, 523-536.	2.7	118

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145	Is There a â€˜Regressive Phenotypeâ€™ of Autism Spectrum Disorder Associated with the Measles-Mumps-Rubella Vaccine? A CPEA Study. Journal of Autism and Developmental Disorders, 2006, 36, 299-316.	2.7	117
146	Familial Autoimmune Thyroid Disease as a Risk Factor for Regression in Children with Autism Spectrum Disorder: A CPEA Study. Journal of Autism and Developmental Disorders, 2006, 36, 317-324.	2.7	99
147	ERP Evidence of Atypical Face Processing in Young Children with Autism. Journal of Autism and Developmental Disorders, 2006, 36, 881-890.	2.7	166
148	How many doctors does it take to make an autism spectrum diagnosis?. Autism, 2006, 10, 439-451.	4.1	216
149	Individuals with Autism Spectrum Disorder Show Normal Responses to a Fear Potential Startle Paradigm. Journal of Autism and Developmental Disorders, 2005, 35, 575-583.	2.7	67
150	Congenital Versus Regressive Onset of Autism Spectrum Disorders. Focus on Autism and Other Developmental Disabilities, 2005, 20, 169-179.	1.3	33
151	Early Regression in Social Communication in Autism Spectrum Disorders: A CPEA Study. Developmental Neuropsychology, 2005, 27, 311-336.	1.4	147