

Molly Losh

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7096552/publications.pdf>

Version: 2024-02-01

58
papers

3,978
citations

218381

26
h-index

168136

53
g-index

62
all docs

62
docs citations

62
times ranked

3360
citing authors

#	ARTICLE	IF	CITATIONS
1	The Broad Autism Phenotype Questionnaire. <i>Journal of Autism and Developmental Disorders</i> , 2007, 37, 1679-1690.	1.7	451
2	Narrative ability in high-functioning children with autism or Asperger's syndrome. <i>Journal of Autism and Developmental Disorders</i> , 2003, 33, 239-251.	1.7	350
3	"Frog, where are you?" Narratives in children with specific language impairment, early focal brain injury, and Williams syndrome. <i>Brain and Language</i> , 2004, 88, 229-247.	0.8	337
4	Defining key features of the broad autism phenotype: A comparison across parents of multiple and single incidence autism families. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2008, 147B, 424-433.	1.1	245
5	Anorexia nervosa and autism spectrum disorders: Guided investigation of social cognitive endophenotypes. <i>Psychological Bulletin</i> , 2007, 133, 976-1006.	5.5	244
6	Neuropsychological Profile of Autism and the Broad Autism Phenotype. <i>Archives of General Psychiatry</i> , 2009, 66, 518.	13.8	238
7	"The frog ate the bug and made his mouth sad": narrative competence in children with autism. <i>Journal of Abnormal Child Psychology</i> , 2000, 28, 193-204.	3.5	232
8	Understanding of emotional experience in autism: Insights from the personal accounts of high-functioning children with autism. <i>Developmental Psychology</i> , 2006, 42, 809-818.	1.2	178
9	Social-cognition and the broad autism phenotype: identifying genetically meaningful phenotypes. <i>Journal of Child Psychology and Psychiatry and Allied Disciplines</i> , 2007, 48, 105-112.	3.1	170
10	Associated features in females with an FMR1 premutation. <i>Journal of Neurodevelopmental Disorders</i> , 2014, 6, 30.	1.5	116
11	Brief Report: Vocational Outcomes for Young Adults with Autism Spectrum Disorders at Six Months After Virtual Reality Job Interview Training. <i>Journal of Autism and Developmental Disorders</i> , 2015, 45, 3364-3369.	1.7	109
12	Cardiac autonomic regulation in autism and Fragile X syndrome: A review. <i>Psychological Bulletin</i> , 2015, 141, 141-175.	5.5	85
13	Current Developments in the Genetics of Autism: From Phenome to Genome. <i>Journal of Neuropathology and Experimental Neurology</i> , 2008, 67, 829-837.	0.9	84
14	A Comparison of Pragmatic Language in Boys With Autism and Fragile X Syndrome. <i>Journal of Speech, Language, and Hearing Research</i> , 2014, 57, 1692-1707.	0.7	84
15	Consistency between research and clinical diagnoses of autism among boys and girls with fragile X syndrome. <i>Journal of Intellectual Disability Research</i> , 2014, 58, 940-952.	1.2	84
16	Longitudinal profiles of expressive vocabulary, syntax and pragmatic language in boys with fragile X syndrome or Down syndrome. <i>International Journal of Language and Communication Disorders</i> , 2013, 48, 432-443.	0.7	83
17	Quantifying Narrative Ability in Autism Spectrum Disorder: A Computational Linguistic Analysis of Narrative Coherence. <i>Journal of Autism and Developmental Disorders</i> , 2014, 44, 3016-3025.	1.7	75
18	Social Communication and Theory of Mind in Boys with Autism and Fragile X Syndrome. <i>Frontiers in Psychology</i> , 2012, 3, 266.	1.1	72

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19	Lower birth weight indicates higher risk of autistic traits in discordant twin pairs. <i>Psychological Medicine</i> , 2012, 42, 1091-1102.	2.7	66
20	Defining genetically meaningful language and personality traits in relatives of individuals with fragile X syndrome and relatives of individuals with autism. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2012, 159B, 660-668.	1.1	66
21	A developmental, longitudinal investigation of autism phenotypic profiles in fragile X syndrome. <i>Journal of Neurodevelopmental Disorders</i> , 2016, 8, 47.	1.5	52
22	Physiological Arousal in Autism and Fragile X Syndrome: Group Comparisons and Links With Pragmatic Language. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2013, 118, 475-495.	0.8	45
23	What's the story? A computational analysis of narrative competence in autism. <i>Autism</i> , 2018, 22, 335-344.	2.4	36
24	Sex differences and within-family associations in the broad autism phenotype. <i>Autism</i> , 2014, 18, 106-116.	2.4	35
25	Elevated Polygenic Burden for Autism Spectrum Disorder Is Associated With the Broad Autism Phenotype in Mothers of Individuals With Autism Spectrum Disorder. <i>Biological Psychiatry</i> , 2021, 89, 476-485.	0.7	32
26	Signaling of noncomprehension in communication breakdowns in fragile X syndrome, Down syndrome, and autism spectrum disorder. <i>Journal of Communication Disorders</i> , 2017, 65, 22-34.	0.8	31
27	A Multimethod Analysis of Pragmatic Skills in Children and Adolescents With Fragile X Syndrome, Autism Spectrum Disorder, and Down Syndrome. <i>Journal of Speech, Language, and Hearing Research</i> , 2018, 61, 3023-3037.	0.7	31
28	An Acoustic Characterization of Prosodic Differences in Autism Spectrum Disorder and First-Degree Relatives. <i>Journal of Autism and Developmental Disorders</i> , 2020, 50, 3032-3045.	1.7	29
29	Links between looking and speaking in autism and first-degree relatives: insights into the expression of genetic liability to autism. <i>Molecular Autism</i> , 2018, 9, 51.	2.6	27
30	Mechanisms of voice control related to prosody in autism spectrum disorder and first-degree relatives. <i>Autism Research</i> , 2019, 12, 1192-1210.	2.1	25
31	Rapid automatized naming as an index of genetic liability to autism. <i>Journal of Neurodevelopmental Disorders</i> , 2010, 2, 109-116.	1.5	24
32	Eye-voice span during rapid automatized naming: evidence of reduced automaticity in individuals with autism spectrum disorder and their siblings. <i>Journal of Neurodevelopmental Disorders</i> , 2014, 6, 33.	1.5	21
33	The Broad Autism Phenotype. , 2011, , 457-476.		20
34	Perception of affect in biological motion cues in anorexia nervosa. <i>International Journal of Eating Disorders</i> , 2013, 46, 12-22.	2.1	19
35	Developmental Markers of Genetic Liability to Autism in Parents: A Longitudinal, Multigenerational Study. <i>Journal of Autism and Developmental Disorders</i> , 2017, 47, 834-845.	1.7	17
36	Understanding Social Communication Differences in Autism Spectrum Disorder and First-Degree Relatives: A Study of Looking and Speaking. <i>Journal of Autism and Developmental Disorders</i> , 2020, 50, 2128-2141.	1.7	17

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37	A case of autism and uniparental disomy of chromosome 1. <i>Human Genetics</i> , 2005, 117, 200-206.	1.8	14
38	A Duck Wearing Boots?! Pragmatic Language Strategies for Repairing Communication Breakdowns Across Genetically Based Neurodevelopmental Disabilities. <i>Journal of Speech, Language, and Hearing Research</i> , 2018, 61, 1440-1454.	0.7	14
39	A constellation of eye-tracking measures reveals social attention differences in ASD and the broad autism phenotype. <i>Molecular Autism</i> , 2022, 13, 18.	2.6	14
40	Pragmatic Language in Autism and Fragile X Syndrome: Genetic and Clinical Applications. <i>Perspectives on Language Learning and Education</i> , 2012, 19, 48-55.	0.2	13
41	Cross-linguistic patterns of speech prosodic differences in autism: A machine learning study. <i>PLoS ONE</i> , 2022, 17, e0269637.	1.1	13
42	Language processing skills linked to FMR1 variation: A study of gaze-language coordination during rapid automatized naming among women with the FMR1 premutation. <i>PLoS ONE</i> , 2019, 14, e0219924.	1.1	11
43	Lifelong Tone Language Experience does not Eliminate Deficits in Neural Encoding of Pitch in Autism Spectrum Disorder. <i>Journal of Autism and Developmental Disorders</i> , 2021, 51, 3291-3310.	1.7	11
44	Systematic Screening for Subtelomeric Anomalies in a Clinical Sample of Autism. <i>Journal of Autism and Developmental Disorders</i> , 2007, 37, 703-708.	1.7	10
45	Common-variant associations with fragile X syndrome. <i>Molecular Psychiatry</i> , 2019, 24, 338-344.	4.1	8
46	Longitudinal analysis of communication repair skills across three neurodevelopmental disabilities. <i>International Journal of Language and Communication Disorders</i> , 2020, 55, 26-42.	0.7	8
47	The Phenotypic Profile Associated With the FMR1 Premutation in Women: An Investigation of Clinical-Behavioral, Social-Cognitive, and Executive Abilities. <i>Frontiers in Psychiatry</i> , 2021, 12, 718485.	1.3	8
48	Physiological regulation and social-emotional processing in female carriers of the FMR1 premutation. <i>Physiology and Behavior</i> , 2020, 214, 112746.	1.0	7
49	Expression and Characterization of Human Fragile X Mental Retardation Protein Isoforms and Interacting Proteins in Human Cells. <i>Proteomics Insights</i> , 2019, 10, 117864181882526.	2.0	5
50	Verbal entrainment in autism spectrum disorder and first-degree relatives. <i>Scientific Reports</i> , 2022, 12, .	1.6	4
51	A cross-cultural study showing deficits in gaze-language coordination during rapid automatized naming among individuals with ASD. <i>Scientific Reports</i> , 2021, 11, 13401.	1.6	3
52	Response to: Genichi Sugihara, Kenji J. Tsuchiya, Nori Takei, Letter to the Editor: Broad Autism Phenotype from Schizophrenia-Spectrum Disorders. <i>Journal of Autism and Developmental Disorders</i> , 2008, 38, 2000-2001.	1.7	1
53	A Unique Visual Attention Profile Associated With the FMR1 Premutation. <i>Frontiers in Genetics</i> , 2021, 12, 591211.	1.1	1
54	A Longitudinal Study of Parent-Child Interactions and Language Outcomes in Fragile X Syndrome and Other Neurodevelopmental Disorders. <i>Frontiers in Psychiatry</i> , 2021, 12, 718572.	1.3	1

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55	Neural Processing of Speech Sounds in ASD and First-Degree Relatives. Journal of Autism and Developmental Disorders, 0, , .	1.7	1
56	Understanding Social Communication Differences in ASD and First-Degree Relatives. , 2021, , 4956-4963.		0
57	Understanding Social-Communication Differences in Autism Spectrum Disorder and First-Degree Relatives. , 2020, , 1-8.		0
58	Childhood Academic Performance: A Potential Marker of Genetic Liability to Autism. Journal of Autism and Developmental Disorders, 2022, , 1.	1.7	0