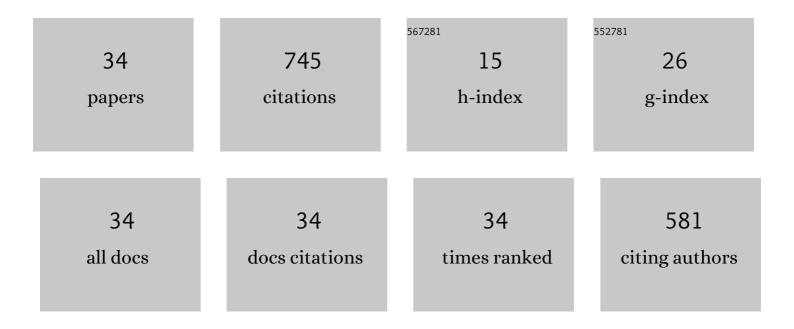
## Jessica Klusek

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Maternal Pragmatic Language Difficulties in the FMR1 Premutation and the Broad Autism Phenotype: Associations with Individual and Family Outcomes. Journal of Autism and Developmental Disorders, 2022, 52, 835-851.	2.7	7
2	Daily Living Skills in Adolescent and Young Adult Males With Fragile X Syndrome. American Journal on Intellectual and Developmental Disabilities, 2022, 127, 64-83.	1.6	1
3	Family history of FXTAS is associated with age-related cognitive-linguistic decline among mothers with the FMR1 premutation. Journal of Neurodevelopmental Disorders, 2022, 14, 7.	3.1	3
4	Cluttering in the Speech of Young Men With Fragile X Syndrome. Journal of Speech, Language, and Hearing Research, 2022, 65, 954-969.	1.6	1
5	Verbal inhibition declines among older women with high FMR1 premutation expansions: A prospective study. Brain and Cognition, 2022, 159, 105851.	1.8	3
6	Response Inhibition Deficits in Women with the FMR1 Premutation are Associated with Age and Fall Risk. Brain and Cognition, 2021, 148, 105675.	1.8	9
7	Family as a Context for Child Development: Mothers with the FMR1 Premutation and Their Children with Fragile X Syndrome. Seminars in Speech and Language, 2021, 42, 277-286.	0.8	0
8	Concurrent Associations between Expressive Language Ability and Independence in Adolescents and Adults with Fragile X Syndrome. Brain Sciences, 2021, 11, 1179.	2.3	3
9	Trajectories of Heart Activity Across Infancy to Early Childhood Differentially Predict Autism and Anxiety Symptoms in Fragile X Syndrome. Frontiers in Psychiatry, 2021, 12, 727559.	2.6	8
10	The <i>FMR1</i> Premutation Phenotype and Mother-Youth Synchrony in Fragile X Syndrome. American Journal on Intellectual and Developmental Disabilities, 2021, 126, 443-459.	1.6	2
11	A novel eyeâ€ŧracking paradigm for indexing social avoidanceâ€related behavior in fragile X syndrome. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2020, 183, 5-16.	1.7	11
12	Inhibition deficits are modulated by age and CGG repeat length in carriers of the FMR1 premutation allele who are mothers of children with fragile X syndrome. Brain and Cognition, 2020, 139, 105511.	1.8	22
13	Vagal Tone as a Putative Mechanism for Pragmatic Competence: An Investigation of Carriers of the FMR1 Premutation. Journal of Autism and Developmental Disorders, 2019, 49, 197-208.	2.7	13
14	Prevalence and Predictors of Anxiety Disorders in Adolescent and Adult Males with Autism Spectrum Disorder and Fragile X Syndrome. Journal of Autism and Developmental Disorders, 2019, 49, 1131-1141.	2.7	36
15	ASD Comorbidity in Fragile X Syndrome: Symptom Profile and Predictors of Symptom Severity in Adolescent and Young Adult Males. Journal of Autism and Developmental Disorders, 2019, 49, 960-977.	2.7	48
16	Gesture Frequency and Function in Infants With Fragile X Syndrome and Infant Siblings of Children With Autism Spectrum Disorder. Journal of Speech, Language, and Hearing Research, 2019, 62, 2386-2399.	1.6	11
17	Reading in Children With Fragile X Syndrome: Phonological Awareness and Feasibility of Intervention. American Journal on Intellectual and Developmental Disabilities, 2018, 123, 193-211.	1.6	4
18	Impaired eye contact in the <i>FMR1</i> premutation is not associated with social anxiety or the broad autism phenotype. Clinical Neuropsychologist, 2018, 32, 1337-1352.	2.3	8

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#	Article	IF	CITATIONS
19	Cortisol profiles differentiated in adolescents and young adult males with fragile X syndrome versus autism spectrum disorder. Developmental Psychobiology, 2018, 60, 78-89.	1.6	11
20	Biobehavioral composite of social aspects of anxiety in young adults with fragile X syndrome contrasted to autism spectrum disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2018, 177, 665-675.	1.7	34
21	Curvilinear Association Between Language Disfluency and FMR1 CGG Repeat Size Across the Normal, Intermediate, and Premutation Range. Frontiers in Genetics, 2018, 9, 344.	2.3	22
22	Developmental Markers of Genetic Liability to Autism in Parents: A Longitudinal, Multigenerational Study. Journal of Autism and Developmental Disorders, 2017, 47, 834-845.	2.7	17
23	Reduced vagal tone in women with the FMR1 premutation is associated with FMR1 mRNA but not depression or anxiety. Journal of Neurodevelopmental Disorders, 2017, 9, 16.	3.1	12
24	Altered sensitivity to social gaze in the FMR1 premutation and pragmatic language competence. Journal of Neurodevelopmental Disorders, 2017, 9, 31.	3.1	20
25	Pragmatic Language Features of Mothers With the <i>FMR1</i> Premutation Are Associated With the Language Outcomes of Adolescents and Young Adults With Fragile X Syndrome. Journal of Speech, Language, and Hearing Research, 2016, 59, 49-61.	1.6	17
26	Phonological awareness and reading in boys with fragile X syndrome. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2015, 56, 30-39.	5.2	19
27	Cardiac autonomic regulation in autism and Fragile X syndrome: A review Psychological Bulletin, 2015, 141, 141-175.	6.1	85
28	Reading and Phonological Skills in Boys with Fragile X Syndrome. Journal of Autism and Developmental Disorders, 2015, 45, 1699-1711.	2.7	15
29	Teaching reading to youth with fragile X syndrome: Should phonemic awareness and phonics instruction be used?. EBP Briefs, 2015, 9, 47-61.	0.5	1
30	A Comparison of Pragmatic Language in Boys With Autism and Fragile X Syndrome. Journal of Speech, Language, and Hearing Research, 2014, 57, 1692-1707.	1.6	84
31	Sex differences and within-family associations in the broad autism phenotype. Autism, 2014, 18, 106-116.	4.1	35
32	Physiological Arousal in Autism and Fragile X Syndrome: Group Comparisons and Links With Pragmatic Language. American Journal on Intellectual and Developmental Disabilities, 2013, 118, 475-495.	1.6	45
33	Social Communication and Theory of Mind in Boys with Autism and Fragile X Syndrome. Frontiers in Psychology, 2012, 3, 266.	2.1	72
34	Defining genetically meaningful language and personality traits in relatives of individuals with fragile X syndrome and relatives of individuals with autism. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2012, 159B, 660-668.	1.7	66