Chris Oliver

List of Publications by Year in descending order

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230 papers

8,318 citations

50276 46 h-index 71685 **76** g-index

241 all docs

241 docs citations

times ranked

241

4487 citing authors

#	Article	IF	Citations
1	Risk markers associated with challenging behaviours in people with intellectual disabilities: a meta-analytic study. Journal of Intellectual Disability Research, 2003, 47, 405-416.	2.0	548
2	Down's Syndrome and Alzheimer's disease: a review. Psychological Medicine, 1986, 16, 307-322.	4.5	322
3	Prevalence of autism spectrum disorder phenomenology in genetic disorders: a systematic review and meta-analysis. Lancet Psychiatry,the, 2015, 2, 909-916.	7.4	280
4	Diagnosis and management of Cornelia de Lange syndrome: first international consensus statement. Nature Reviews Genetics, 2018, 19, 649-666.	16.3	223
5	The prevalence and phenomenology of selfâ€injurious and aggressive behaviour in genetic syndromes. Journal of Intellectual Disability Research, 2011, 55, 109-120.	2.0	212
6	The Prevalence and Phenomenology of Repetitive Behavior in Genetic Syndromes. Journal of Autism and Developmental Disorders, 2009, 39, 572-588.	2.7	202
7	A four year prospective study of age-related cognitive change in adults with Down's syndrome. Psychological Medicine, 1998, 28, 1365-1377.	4.5	148
8	Selfâ€injurious behaviour in individuals with autism spectrum disorder and intellectual disability. Journal of Intellectual Disability Research, 2012, 56, 476-489.	2.0	142
9	Predictors of Psychological Morbidity in Parents of Children with Intellectual Disabilities. Journal of Pediatric Psychology, 2008, 33, 1129-1136.	2.1	127
10	Self-Injurious Behavior, Self-Restraint, and Compulsive Behaviors in Cornelia de Lange Syndrome. American Journal on Intellectual and Developmental Disabilites, 2002, 107, 146.	2.4	114
11	Self-Injurious Behaviour in Children with Learning Disabilities: Recent Advances in Assessment and Intervention. Journal of Child Psychology and Psychiatry and Allied Disciplines, 1995, 36, 909-927.	5.2	109
12	Delineation of Behavioral Phenotypes in Genetic Syndromes: Characteristics of Autism Spectrum Disorder, Affect and Hyperactivity. Journal of Autism and Developmental Disorders, 2011, 41, 1019-1032.	2.7	88
13	Early Development of Self-Injurious Behavior: An Empirical Study. American Journal on Intellectual and Developmental Disabilites, 2001, 106, 189.	2.4	84
14	Preliminary analysis of the psychometric properties of the Mood, Interest & Pleasure Questionnaire (MIPQ) for adults with severe and profound learning disabilities. British Journal of Clinical Psychology, 2003, 42, 81-93.	3.5	84
15	Task-switching deficits and repetitive behaviour in genetic neurodevelopmental disorders: Data from children with Prader–Willi syndrome chromosome 15 q11–q13 deletion and boys with Fragile X syndrome. Cognitive Neuropsychology, 2009, 26, 172-194.	1.1	84
16	Prevalence of Autism Spectrum Phenomenology in Cornelia de Lange and Cri du Chat Syndromes. American Journal on Intellectual and Developmental Disabilites, 2008, 113, 278.	2.4	83
17	The Association Between Repetitive, Self-Injurious and Aggressive Behavior in Children With Severe Intellectual Disability. Journal of Autism and Developmental Disorders, 2012, 42, 910-919.	2.7	83
18	The assessment of mood in adults who have severe or profound mental retardation. Clinical Psychology Review, 2003, 23, 225-245.	11.4	81

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19	Prevalence of autism spectrum disorder symptomatology and related behavioural characteristics in individuals with Down syndrome. Autism, 2013, 17, 390-404.	4.1	81
20	The expression and assessment of emotions and internal states in individuals with severe or profound intellectual disabilities. Clinical Psychology Review, 2011, 31, 293-306.	11.4	79
21	Behavioural phenotype of Cornelia de Lange syndrome: case–control study. British Journal of Psychiatry, 2008, 193, 466-470.	2.8	77
22	Identification of early selfâ€injurious behaviour in young children with intellectual disability. Journal of Intellectual Disability Research, 1999, 43, 149-163.	2.0	75
23	The behavioural phenotype of Angelman syndrome. Journal of Intellectual Disability Research, 2006, 50, 33-53.	2.0	73
24	Assessing the Severity of Challenging Behaviour: Psychometric Properties of the Challenging Behaviour Interview. Journal of Applied Research in Intellectual Disabilities, 2003, 16, 53-61.	2.0	71
25	Selfâ€injurious behaviour in people with mental handicap: a total population study. Journal of Intellectual Disability Research, 1987, 31, 147-162.	2.0	70
26	Phenotypes and genotypes in individuals with <i>SMC1A</i> variants. American Journal of Medical Genetics, Part A, 2017, 173, 2108-2125.	1.2	69
27	Measurement tools for mental health problems and mental well-being in people with severe or profound intellectual disabilities: A systematic review. Clinical Psychology Review, 2017, 57, 32-44.	11.4	69
28	Down's syndrome and the links with Alzheimer's disease Journal of Neurology, Neurosurgery and Psychiatry, 1995, 59, 111-114.	1.9	68
29	Associations between repetitive questioning, resistance to change, temper outbursts and anxiety in Prader–Willi and Fragileâ€X syndromes. Journal of Intellectual Disability Research, 2009, 53, 265-278.	2.0	67
30	Psychological well-being in parents of children with Angelman, Cornelia de Lange and Cri du Chat syndromes. Journal of Intellectual Disability Research, 2011, 55, 397-410.	2.0	64
31	Genomic imprinting and the expression of affect in Angelman syndrome: what's in the smile?. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2007, 48, 571-579.	5. 2	62
32	The relationship between acquired impairments of executive function and behaviour change in adults with Down syndrome. Journal of Intellectual Disability Research, 2010, 54, 393-405.	2.0	61
33	The age related prevalence of aggression and self-injury in persons with an intellectual disability: A review. Research in Developmental Disabilities, 2013, 34, 764-775.	2.2	61
34	The Chronicity of Self-Injurious Behaviour: A Long-Term Follow-Up of a Total Population Study. Journal of Applied Research in Intellectual Disabilities, 2011, 24, 105-117.	2.0	60
35	Effects of Environmental Events on Smiling and Laughing Behavior in Angelman Syndrome. American Journal on Intellectual and Developmental Disabilites, 2002, 107, 194.	2.4	59
36	The early development of self-injurious behaviour: evaluating the role of social reinforcement. Journal of Intellectual Disability Research, 2005, 49, 591-599.	2.0	59

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#	Article	IF	Citations
37	A cross-syndrome cohort comparison of sleep disturbance in children with Smith-Magenis syndrome, Angelman syndrome, autism spectrum disorder and tuberous sclerosis complex. Journal of Neurodevelopmental Disorders, 2018, 10, 9.	3.1	59
38	Self-injurious behavior. Neuroscience and Biobehavioral Reviews, 2018, 84, 483-491.	6.1	57
39	The Neuropsychological Assessment of Age Related Cognitive Deficits in Adults with Down's Syndrome. Journal of Applied Research in Intellectual Disabilities, 1998, 11, 255-272.	2.0	56
40	Characteristics of autism spectrum disorder in Cornelia de Lange syndrome. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2012, 53, 883-891.	5.2	56
41	The association between repetitive behaviours, impulsivity and hyperactivity in people with intellectual disability. Journal of Intellectual Disability Research, 2010, 54, 1078-1092.	2.0	55
42	Differential Effects of Severe Self-injurious Behaviour on the Behaviour of Others. Behavioural and Cognitive Psychotherapy, 1992, 20, 355-365.	1.2	53
43	Anxiety Disorders in Williams Syndrome Contrasted with Intellectual Disability and the General Population: A Systematic Review and Meta-Analysis. Journal of Autism and Developmental Disorders, 2017, 47, 3765-3777.	2.7	53
44	Self-injurious behavior in Rett syndrome: Interactions between features of Rett syndrome and operant conditioning. Journal of Autism and Developmental Disorders, 1993, 23, 91-109.	2.7	52
45	Health and sleep problems in Cornelia de Lange Syndrome: a case control study. Journal of Intellectual Disability Research, 2008, 52, 458-468.	2.0	52
46	Persistence of self-injurious behaviour in autism spectrum disorder over 3Âyears: a prospective cohort study of risk markers. Journal of Neurodevelopmental Disorders, 2016, 8, 21.	3.1	52
47	Sleep duration and sleep quality in people with and without intellectual disability: A meta-analysis. Sleep Medicine Reviews, 2018, 40, 135-150.	8.5	51
48	The relationship between levels of mood, interest and pleasure and 'challenging behaviour' in adults with severe and profound intellectual disability. Journal of Intellectual Disability Research, 2002, 46, 191-197.	2.0	50
49	The behavioural phenotype of Smith–Magenis syndrome: evidence for a gene–environment interaction. Journal of Intellectual Disability Research, 2008, 52, 830-841.	2.0	49
50	The assessment of foodâ€related problems in individuals with Praderâ€Willi syndrome. British Journal of Clinical Psychology, 2003, 42, 379-392.	3.5	48
51	Delineating the Profile of Autism Spectrum Disorder Characteristics in Cornelia de Lange and Fragile X Syndromes. American Journal on Intellectual and Developmental Disabilities, 2013, 118, 55-73.	1.6	48
52	Selfâ€injurious Behaviour and People with Intellectual Disabilities: Assessing the Behavioural Knowledge and Causal Explanations of Care Staff. Journal of Applied Research in Intellectual Disabilities, 1996, 9, 229-239.	2.0	46
53	Selfâ€injurious behaviour in Cornelia de Lange syndrome: 1. Prevalence and phenomenology. Journal of Intellectual Disability Research, 2009, 53, 575-589.	2.0	46
54	Neural correlates of task switching in paternal 15q11–q13 deletion Prader–Willi syndrome. Brain Research, 2010, 1363, 128-142.	2.2	46

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55	The relationship between specific cognitive impairment and behaviour in Prader–Willi syndrome. Journal of Intellectual Disability Research, 2011, 55, 152-171.	2.0	46
56	Self-injurious behaviour in young children with Lesch-Nyhan syndrome. Developmental Medicine and Child Neurology, 2001, 43, 745.	2.1	46
57	Environmental Influences on the Behavioral Phenotype of Angelman Syndrome. American Journal on Intellectual and Developmental Disabilites, 2006, 111, 311.	2.4	45
58	Experimental functional analysis of aggression in children with Angelman syndrome. Research in Developmental Disabilities, 2009, 30, 1095-1106.	2.2	45
59	Prevalence, phenomenology, aetiology and predictors of challenging behaviour in Smithâ€Magenis syndrome. Journal of Intellectual Disability Research, 2011, 55, 138-151.	2.0	45
60	The importance of understanding the behavioural phenotypes of genetic syndromes associated with intellectual disability. Paediatrics and Child Health (United Kingdom), 2014, 24, 468-472.	0.4	45
61	Practitioner Review: Selfâ€injurious behaviour in children with developmental delay. Journal of Child Psychology and Psychiatry and Allied Disciplines, 2015, 56, 1042-1054.	5.2	45
62	The prevalence of aggression in genetic syndromes: A review. Research in Developmental Disabilities, 2014, 35, 1051-1071.	2.2	43
63	Is Theory of Mind Understanding Impaired in Males with Fragile X Syndrome?. Journal of Abnormal Child Psychology, 2007, 35, 17-28.	3.5	42
64	Sleep disorders in rare genetic syndromes: a meta-analysis of prevalence and profile. Molecular Autism, 2021, 12, 18.	4.9	42
65	Positive impact and its relationship to well-being in parents of children with intellectual disability: a literature review. International Journal of Developmental Disabilities, 2015, 61, 1-19.	2.0	41
66	Visual preference for social stimuli in individuals with autism or neurodevelopmental disorders: an eye-tracking study. Molecular Autism, 2016, 7, 24.	4.9	41
67	Self-injury, aggression and destruction in children with severe intellectual disability: Incidence, persistence and novel, predictive behavioural risk markers. Research in Developmental Disabilities, 2016, 49-50, 291-301.	2.2	41
68	Temper outbursts in <scp>P</scp> rader– <scp>W</scp> illi syndrome: causes, behavioural and emotional sequence and responses by carers. Journal of Intellectual Disability Research, 2014, 58, 134-150.	2.0	40
69	A Comparative Study of Sociability in Angelman, Cornelia de Lange, Fragile X, Down and Rubinstein Taybi Syndromes and Autism Spectrum Disorder. American Journal on Intellectual and Developmental Disabilities, 2016, 121, 465-486.	1.6	40
70	Diverse Profiles of Anxiety Related Disorders in Fragile X, Cornelia de Lange and Rubinstein–Taybi Syndromes. Journal of Autism and Developmental Disorders, 2017, 47, 3728-3740.	2.7	40
71	Interventions for mental health problems in children and adults with severe intellectual disabilities: a systematic review. BMJ Open, 2018, 8, e021911.	1.9	40
72	Social Anxiety in Cornelia de Lange Syndrome. Journal of Autism and Developmental Disorders, 2009, 39, 1155-1162.	2.7	39

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73	A national survey of Rett syndrome: behavioural characteristics. Journal of Neurodevelopmental Disorders, 2015, 7, 11.	3.1	38
74	The effects of staff training on staff confidence and challenging behavior in services for people with autism spectrum disorders. Research in Autism Spectrum Disorders, 2008, 2, 311-319.	1.5	37
75	Social Behavior and Characteristics of Autism Spectrum Disorder in Angelman, Cornelia de Lange, and Cri du Chat Syndromes. American Journal on Intellectual and Developmental Disabilities, 2013, 118, 262-283.	1.6	37
76	Self-injury and aggression in tuberous sclerosis complex: cross syndrome comparison and associated risk markers. Journal of Neurodevelopmental Disorders, 2014, 6, 10.	3.1	37
77	The association between environmental events and self-injurious behaviour in Cornelia de Lange syndrome. Journal of Intellectual Disability Research, 2005, 49, 269-277.	2.0	36
78	A review of defining and measuring sociability in children with intellectual disabilities. Research in Developmental Disabilities, 2011, 32, 11-24.	2.2	36
79	Phenotype–environment interactions in genetic syndromes associated with severe or profound intellectual disability. Research in Developmental Disabilities, 2011, 32, 404-418.	2.2	34
80	"You Have to Sit and Explain it All, and Explain Yourself.―Mothers' Experiences of Support Services for Their Offspring with a Rare Genetic Intellectual Disability Syndrome. Journal of Genetic Counseling, 2011, 20, 165-177.	1.6	34
81	Causal Models of Clinically Significant Behaviors in Angelman, Cornelia de Lange, Prader–Willi and Smith–Magenis Syndromes. International Review of Research in Developmental Disabilities, 2013, 44, 167-211.	0.8	34
82	Self-Injurious Behaviour in People with Learning Disabilities: Determinants and Interventions. International Review of Psychiatry, 1990, 2, 101-116.	2.8	33
83	Behavioural characteristics associated with dementia assessment referrals in adults with Down syndrome. Journal of Intellectual Disability Research, 2008, 52, 358-368.	2.0	33
84	A specific pathway can be identified between genetic characteristics and behaviour profiles in Prader–Willi syndrome via cognitive, environmental and physiological mechanisms. Journal of Intellectual Disability Research, 2009, 53, 493-500.	2.0	32
85	Repetitive Behavior in Rubinstein–Taybi Syndrome: Parallels with Autism Spectrum Phenomenology. Journal of Autism and Developmental Disorders, 2015, 45, 1238-1253.	2.7	32
86	Effects of Social Context on Social Interaction and Self-Injurious Behavior in Cornelia de Lange Syndrome. American Journal on Intellectual and Developmental Disabilites, 2006, 111, 184.	2.4	31
87	Self-injurious, aggressive and destructive behaviour in children with severe intellectual disability: Prevalence, service need and service receipt in the UK. Research in Developmental Disabilities, 2015, 45-46, 307-315.	2.2	31
88	Predictors of Self-Injurious Behavior and Self-Restraint in Autism Spectrum Disorder: Towards a Hypothesis of Impaired Behavioral Control. Journal of Autism and Developmental Disorders, 2017, 47, 701-713.	2.7	31
89	The Development of a Health Status Measure for Self-report by People with Intellectual Disabilities. Journal of Applied Research in Intellectual Disabilities, 2005, 18, 143-150.	2.0	30
90	Structural and Environmental Characteristics of Stereotyped Behaviors. American Journal on Intellectual and Developmental Disabilites, 2003, 108, 391.	2.4	29

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91	Effects of adult familiarity on social behaviours in Angelman syndrome. Journal of Intellectual Disability Research, 2011, 55, 339-350.	2.0	29
92	A Behavioural Assessment of Social Anxiety and Social Motivation in Fragile X, Cornelia de Lange and Rubinstein-Taybi Syndromes. Journal of Autism and Developmental Disorders, 2020, 50, 127-144.	2.7	28
93	Low mood and challenging behaviour in people with severe and profound intellectual disabilities. Journal of Intellectual Disability Research, 2011, 55, 182-189.	2.0	27
94	Self-injurious behaviour in people with intellectual disability and autism spectrum disorder. Current Opinion in Psychiatry, 2017, 30, 97-101.	6.3	27
95	Age related change in social behavior in children with Angelman syndrome. American Journal of Medical Genetics, Part A, 2011, 155, 1290-1297.	1.2	26
96	Persistence and predictors of self-injurious behaviour in autism: a ten-year prospective cohort study. Molecular Autism, 2020, 11, 8.	4.9	26
97	The Treatment of Severe Self-Injurious Behavior by the Systematic Fading of Restraints: Effects on Self-Injury, Self-Restraint, Adaptive Behavior, and Behavioral Correlates of Affect 11This research was supported by a grant from The Mental Health Foundation. The authors wish to acknowledge the help given by the following people: the staff of Lambeth Walk day centre, Vicky Turk, Alex Proto, Roy Smith,	2.2	25
98	Diane tangidge, and Angela Daniels. Research in Developmental Disabilities, 1990, 19, 143-165. Effects of Training on Controllability Attributions of Behavioural Excesses and Deficits Shown by adults with Down Syndrome and Dementia. Journal of Applied Research in Intellectual Disabilities, 2007, 20, 64-68.	2.0	25
99	The behavioral characteristics of Sotos syndrome. American Journal of Medical Genetics, Part A, 2015, 167, 2945-2956.	1.2	25
100	A national survey of Rett syndrome: Age, clinical characteristics, current abilities, and health. American Journal of Medical Genetics, Part A, 2015, 167, 1493-1500.	1.2	25
101	Prospective study of autism phenomenology and the behavioural phenotype of Phelan–McDermid syndrome: comparison to fragile X syndrome, Down syndrome and idiopathic autism spectrum disorder. Journal of Neurodevelopmental Disorders, 2017, 9, 37.	3.1	25
102	A molecular to molar analysis of communicative and problem behaviors. Research in Developmental Disabilities, 1999, 20, 197-213.	2.2	24
103	Self-injurious behaviour in people with intellectual disability. Current Opinion in Psychiatry, 2010, 23, 412-416.	6.3	24
104	Discrimination training reduces high rate social approach behaviors in Angelman syndrome: Proof of principle. Research in Developmental Disabilities, 2013, 34, 1794-1803.	2.2	24
105	Communication in Angelman syndrome: a scoping review. Developmental Medicine and Child Neurology, 2019, 61, 1266-1274.	2.1	24
106	Profiles of atypical sensory processing in Angelman, Cornelia de Lange and Fragile X syndromes. Journal of Intellectual Disability Research, 2020, 64, 117-130.	2.0	24
107	The nature of social preference and interactions in Smith–Magenis syndrome. Research in Developmental Disabilities, 2013, 34, 4355-4365.	2.2	23
108	Transcutaneous vagus nerve stimulation (t-VNS): A novel effective treatment for temper outbursts in adults with Prader-Willi Syndrome indicated by results from a non-blind study. PLoS ONE, 2019, 14, e0223750.	2.5	23

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109	Prevalence and Risk-Markers of Self-Harm in Autistic Children and Adults. Journal of Autism and Developmental Disorders, 2020, 50, 3561-3574.	2.7	23
110	Relationship Among Challenging, Repetitive, and Communicative Behaviors in Children With Severe Intellectual Disabilities. American Journal on Intellectual and Developmental Disabilities, 2009, 114, 356-368.	1.6	22
111	Ageing in Rett syndrome. Journal of Intellectual Disability Research, 2016, 60, 182-190.	2.0	22
112	Descriptive functional analysis of behavioral excesses shown by adults with Down syndrome and dementia. International Journal of Geriatric Psychiatry, 2003, 18, 844-854.	2.7	21
113	Carer-Reported Contemporary Health Problems in People With Severe and Profound Intellectual Disability and Genetic Syndromes. Journal of Policy and Practice in Intellectual Disabilities, 2007, 4, 120-128.	2.7	21
114	Sleep in children with Angelman syndrome: Parental concerns and priorities. Research in Developmental Disabilities, 2017, 69, 105-115.	2.2	21
115	Allelic variability in D21S11, but not in APP or APOE, is associated with cognitive decline in Down syndrome. NeuroReport, 1997, 8, 1645-1649.	1.2	20
116	Cognitive Deterioration in Adults With Down Syndrome: Effects on the Individual, Caregivers, and Service Use. American Journal on Intellectual and Developmental Disabilites, 2000, 105, 455.	2.4	20
117	Genotype–phenotype correlations in Cornelia de Lange syndrome: Behavioral characteristics and changes with age. American Journal of Medical Genetics, Part A, 2017, 173, 1566-1574.	1.2	20
118	Behaviour in Cornelia de Lange syndrome: a systematic review. Developmental Medicine and Child Neurology, 2017, 59, 361-366.	2.1	20
119	A systematic review of the behaviours associated with depression in people with severe–profound intellectual disability. Journal of Intellectual Disability Research, 2021, 65, 211-229.	2.0	20
120	Aggression and the termination of $\hat{a} \in \infty$ rituals $\hat{a} \in \mathbb{R}$ a new variant of the escape function for challenging behavior?. Research in Developmental Disabilities, 2000, 21, 43-59.	2.2	19
121	Dorsal and ventral stream mediated visual processing in genetic subtypes of Prader–Willi syndrome. Neuropsychologia, 2009, 47, 2367-2373.	1.6	19
122	Selfâ€injurious behaviour in Cornelia de Lange syndrome: 2. association with environmental events. Journal of Intellectual Disability Research, 2009, 53, 590-603.	2.0	19
123	Contrasting age related changes in autism spectrum disorder phenomenology in Cornelia de Lange, Fragile X, and Cri du Chat syndromes: Results from a 2.5 year followâ€up. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2015, 169, 188-197.	1.6	19
124	Chinese Children's Attitudes Toward Mental Retardation. Journal of Developmental and Physical Disabilities, 2000, 12, 73-87.	1.6	18
125	Self-injurious behaviour in individuals with intellectual disabilities. Current Opinion in Psychiatry, 2005, 18, 484-489.	6.3	18
126	Effects of Increasing Task Load on Memory Impairment in Adults With Down Syndrome. American Journal on Intellectual and Developmental Disabilites, 2005, 110, 339.	2.4	18

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127	Facial expression of affect in children with Cornelia de Lange syndrome. Journal of Intellectual Disability Research, 2008, 52, 207-215.	2.0	18
128	Behavioural Excesses and Deficits Associated with Dementia in Adults who have Down Syndrome. Journal of Applied Research in Intellectual Disabilities, 2011, 24, 208-216.	2.0	18
129	Differences in Social Motivation in Children with Smith–Magenis Syndrome and Down Syndrome. Journal of Autism and Developmental Disorders, 2016, 46, 2148-2159.	2.7	18
130	Mental Health and Well-Being in Mothers of Children With Rare Genetic Syndromes Showing Chronic Challenging Behavior: A Cross-Sectional and Longitudinal Study. American Journal on Intellectual and Developmental Disabilities, 2018, 123, 241-253.	1.6	18
131	Sleep in children with Smith–Magenis syndrome: a case–control actigraphy study. Sleep, 2020, 43, .	1.1	18
132	Causal explanations, concern and optimism regarding self-injurious behaviour displayed by individuals with Cornelia de Lange syndrome: the parents' perspective. Journal of Intellectual Disability Research, 2001, 45, 326-334.	2.0	17
133	Functional analysis and functional communication training in individuals with Angelman syndrome. Developmental Neurorehabilitation, 2012, 15, 91-104.	1.1	17
134	The Purported Association Between Depression, Aggression, and Self-Injury in People With Intellectual Disability: A Critical Review of the Literature. American Journal on Intellectual and Developmental Disabilities, 2014, 119, 452-471.	1.6	17
135	Implicit Discrimination of Basic Facial Expressions of Positive/Negative Emotion in Fragile X Syndrome and Autism Spectrum Disorder. American Journal on Intellectual and Developmental Disabilities, 2015, 120, 328-345.	1.6	17
136	Brief Report: A Longitudinal Study of Excessive Smiling and Laughing in Children with Angelman Syndrome. Journal of Autism and Developmental Disorders, 2015, 45, 2624-2627.	2.7	17
137	Executive functioning in Cornelia de Lange syndrome: domain asynchrony and age-related performance. Journal of Neurodevelopmental Disorders, 2017, 9, 29.	3.1	17
138	The Persistence of Self-injurious and Aggressive Behavior in Males with Fragile X Syndrome Over 8 Years: A Longitudinal Study of Prevalence and Predictive Risk Markers. Journal of Autism and Developmental Disorders, 2019, 49, 2913-2922.	2.7	17
139	Phenomenology of Self-Restraint. American Journal on Intellectual and Developmental Disabilites, 2003, 108, 71.	2.4	16
140	A Proactive Psychological Strategy for Determining the Presence of Dementia in Adults with Down Syndrome: Preliminary Description of Service Use and Evaluation. Journal of Policy and Practice in Intellectual Disabilities, 2005, 2, 116-125.	2.7	16
141	Stereotyped behaviour in children with autism and intellectual disability: an examination of the executive dysfunction hypothesis. Journal of Intellectual Disability Research, 2011, 55, 699-709.	2.0	16
142	A Longitudinal Follow-Up Study of Affect in Children and Adults With Cornelia de Lange Syndrome. American Journal on Intellectual and Developmental Disabilities, 2014, 119, 235-252.	1.6	16
143	The Motivating Operation and Negatively Reinforced Problem Behavior. Behavior Modification, 2014, 38, 107-159.	1.6	15
144	Correlates of self-injurious, aggressive and destructive behaviour in children under five who are at risk of developmental delay. Research in Developmental Disabilities, 2014, 35, 36-45.	2.2	15

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145	Self-injury and aggression in adults with tuberous sclerosis complex: Frequency, associated person characteristics, and implications for assessment. Research in Developmental Disabilities, 2017, 64, 119-130.	2.2	15
146	Using Bayesian methodology to explore the profile of mental health and well-being in 646 mothers of children with 13 rare genetic syndromes in relation to mothers of children with autism. Orphanet Journal of Rare Diseases, 2018, 13, 185.	2.7	15
147	Descriptive analysis of challenging behaviours shown by adults with acquired brain injury. Neuropsychological Rehabilitation, 2010, 20, 212-238.	1.6	14
148	The behavioural phenotype of Potocki-Lupski syndrome: a cross-syndrome comparison. Journal of Neurodevelopmental Disorders, 2018, 10, 2.	3.1	14
149	Brief Report: Repetitive Behaviour Profiles in Williams syndrome: Cross Syndrome Comparisons with Prader–Willi and Down syndromes. Journal of Autism and Developmental Disorders, 2018, 48, 326-331.	2.7	14
150	Are Angelman and Praderâ€Willi syndromes more similar than we thought? Foodâ€related behavior problems in Angelman, Cornelia de Lange, Fragile X, Praderâ€Willi and 1p36 deletion syndromes. American Journal of Medical Genetics, Part A, 2015, 167, 572-578.	1.2	13
151	Multi-Method Assessment of Sleep in Children With Angelman Syndrome: A Case–Controlled Study. Frontiers in Psychiatry, 2019, 10, 874.	2.6	13
152	Sleep disorders in children with Angelman and Smith-Magenis syndromes: The assessment of potential causes of disrupted settling and night time waking. Research in Developmental Disabilities, 2020, 97, 103555.	2.2	13
153	Anxiety characteristics in individuals with Williams syndrome. Journal of Applied Research in Intellectual Disabilities, 2021, 34, 1098-1107.	2.0	13
154	A graphical method to aid the sequential analysis of observational data. Behavior Research Methods, 1997, 29, 563-573.	1.3	12
155	Auditory P300 response in the assessment of Alzheimer's disease in Down's syndrome: a 2â€year followâ€up study. Journal of Intellectual Disability Research, 1988, 32, 455-463.	2.0	12
156	Coping with challenges to memory in people with mild to moderate Alzheimer's disease: Observation of behaviour in response to analogues of everyday situations. Aging and Mental Health, 2009, 13, 46-53.	2.8	12
157	Dissociation of Cross-Sectional Trajectories for Verbal and Visuo-Spatial Working Memory Development in Rubinstein-Taybi Syndrome. Journal of Autism and Developmental Disorders, 2016, 46, 2064-2071.	2.7	12
158	Differential effects of anxiety and autism on social scene scanning in males with fragile X syndrome. Journal of Neurodevelopmental Disorders, 2017, 9, 9.	3.1	12
159	An experimental study of executive function and social impairment in Cornelia de Lange syndrome. Journal of Neurodevelopmental Disorders, 2017, 9, 33.	3.1	12
160	Coping Strategies in Mothers of Children with Intellectual Disabilities Showing Multiple Forms of Challenging Behaviour: Associations with Maternal Mental Health. Behavioural and Cognitive Psychotherapy, 2018, 46, 257-275.	1.2	12
161	Fragile X syndrome: an overview of cause, characteristics, assessment and management. Paediatrics and Child Health (United Kingdom), 2020, 30, 400-403.	0.4	12
162	An Observational Study of Social Interaction Skills and Behaviors in Cornelia de Lange, Fragile X and Rubinstein-Taybi Syndromes. Journal of Autism and Developmental Disorders, 2020, 50, 4001-4010.	2.7	12

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163	The Relationship between Components of the Behavioural Phenotype in Praderâ€Willi Syndrome. Journal of Applied Research in Intellectual Disabilities, 2009, 22, 403-407.	2.0	11
164	Increased Exposure to Rigid Routines can Lead to Increased Challenging Behavior Following Changes to Those Routines. Journal of Autism and Developmental Disorders, 2015, 45, 1569-1578.	2.7	11
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