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	241 times ranked	4487 citing authors

#	Article	IF	Citations
1	Direct assessment of overnight parent-child proximity in children with behavioral insomnia: Extending models of operant and classical conditioning. Behavioral Sleep Medicine, 2023, 21, 254-272.	2.1	3
2	Acquired mild cognitive impairment in adults with Down syndrome: Ageâ€related prevalence derived from single point assessment data normed by degree of intellectual disability. International Journal of Geriatric Psychiatry, 2022, 37, .	2.7	7
3	Executive function, repetitive behaviour and restricted interests in neurodevelopmental disorders. Research in Developmental Disabilities, 2022, 122, 104166.	2.2	3
4	Caregiversâ€™ experience of sleep management in Smithâ€“Magenis syndrome: a mixed-methods study. Orphanet Journal of Rare Diseases, 2022, 17, 35.	2.7	6
5	Distress in people with severe disability: the unmet challenge. Developmental Medicine and Child Neurology, 2022, 64, 401-402.	2.1	0
6	Cornelia de Lange syndrome and the Cohesin complex: Abstracts from the 9th Biennial Scientific and Educational Virtual Symposium 2020. American Journal of Medical Genetics, Part A, 2022, 188, 1005-1014.	1.2	1
7	The behavioural phenotype of SATB2-associated syndrome: a within-group and cross-syndrome analysis. Journal of Neurodevelopmental Disorders, 2022, 14, 25.	3.1	5
8	Prevalence of anxiety symptomatology and diagnosis in syndromic intellectual disability: A systematic review and meta-analysis. Neuroscience and Biobehavioral Reviews, 2022, 138, 104719.	6.1	10
9	Skin Picking in People with Praderâ€“Willi Syndrome: Phenomenology and Management. Journal of Autism and Developmental Disorders, 2021, 51, 286-297.	2.7	2
10	Genetic modifiers in rare disorders: the case of fragile X syndrome. European Journal of Human Genetics, 2021, 29, 173-183.	2.8	9
11	A systematic review of the behaviours associated with depression in people with severeâ€“profound intellectual disability. Journal of Intellectual Disability Research, 2021, 65, 211-229.	2.0	20
12	Sleep disorders in rare genetic syndromes: a meta-analysis of prevalence and profile. Molecular Autism, 2021, 12, 18.	4.9	42
13	Refining the Behavioral Phenotype of Angelman Syndrome: Examining Differences in Motivation for Social Contact Between Genetic Subgroups. Frontiers in Behavioral Neuroscience, 2021, 15, 618271.	2.0	7
14	Anxiety characteristics in individuals with Williams syndrome. Journal of Applied Research in Intellectual Disabilities, 2021, 34, 1098-1107.	2.0	13
15	Low speech rate but high gesture rate during conversational interaction in people with Cornelia de Lange syndrome. Journal of Intellectual Disability Research, 2021, 65, 601-607.	2.0	3
16	The adaptive functioning profile of Pitt-Hopkins syndrome. European Journal of Medical Genetics, 2021, 64, 104279.	1.3	3
17	Behaviour across the lifespan in Cornelia de Lange syndrome. Current Opinion in Psychiatry, 2021, 34, 112-117.	6.3	7
18	The development of early social cognitive skills in neurogenetic syndromes associated with autism: Cornelia de Lange, fragile X and Rubinsteinâ€“Taybi syndromes. Orphanet Journal of Rare Diseases, 2021, 16, 488.	2.7	7

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19	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
20	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
21	Sleep in children with Smithâ€“Magenis syndrome: a caseâ€“control actigraphy study. Sleep, 2020, 43, .	1.1	18
22	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
23	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
24	The Profiles and Correlates of Psychopathology in Adolescents and Adults with Williams, Fragile X and Praderâ€“Willi Syndromes. Journal of Autism and Developmental Disorders, 2020, 50, 893-903.	2.7	7
25	Fragile X syndrome: an overview of cause, characteristics, assessment and management. Paediatrics and Child Health (United Kingdom), 2020, 30, 400-403.	0.4	12
26	The behaviour and wellbeing of children and adults with severe intellectual disability and complex needs: the Be-Well checklist for carers and professionals. Paediatrics and Child Health (United) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 457		0
27	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
28	Scaling of Early Social Cognitive Skills in Typically Developing Infants and Children with Autism Spectrum Disorder. Journal of Autism and Developmental Disorders, 2020, 50, 3988-4000.	2.7	7
29	Cornelia de Lange Syndrome. , 2020, , 129-157.		0
30	Persistence and predictors of self-injurious behaviour in autism: a ten-year prospective cohort study. Molecular Autism, 2020, 11, 8.	4.9	26
31	Sleep problems in autism spectrum disorders: A comparison to sleep in typically developing children using actigraphy, diaries and questionnaires. Research in Autism Spectrum Disorders, 2019, 67, 101439.	1.5	10
32	Behavioural and psychological characteristics in Pitt-Hopkins syndrome: a comparison with Angelman and Cornelia de Lange syndromes. Journal of Neurodevelopmental Disorders, 2019, 11, 24.	3.1	10
33	Temper outbursts in Lowe syndrome: Characteristics, sequence, environmental context and comparison to Praderâ€“Willi syndrome. Journal of Applied Research in Intellectual Disabilities, 2019, 32, 1216-1227.	2.0	8
34	Lifespan trajectory of affect in Cornelia de Lange syndrome: towards a neurobiological hypothesis. Journal of Neurodevelopmental Disorders, 2019, 11, 6.	3.1	10
35	Communication in Angelman syndrome: a scoping review. Developmental Medicine and Child Neurology, 2019, 61, 1266-1274.	2.1	24
36	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

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37	Age-related Behavioural Change in Cornelia de Lange and Cri du Chat Syndromes: A Seven Year Follow-up Study. Journal of Autism and Developmental Disorders, 2019, 49, 2476-2487.	2.7	8
38	Multi-Method Assessment of Sleep in Children With Angelman Syndrome: A Caseâ€"Controlled Study. Frontiers in Psychiatry, 2019, 10, 874.	2.6	13
39	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
40	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
41	Persistence of selfâ€injury, aggression and property destruction in children and adults with tuberous sclerosis complex. Journal of Intellectual Disability Research, 2018, 62, 1058-1071.	2.0	10
42	Differences in the Information Needs of Parents With a Child With a Genetic Syndrome: A Crossâ€Syndrome Comparison. Journal of Policy and Practice in Intellectual Disabilities, 2018, 15, 94-100.	2.7	8
43	Overactivity, impulsivity and repetitive behaviour in males with fragile X syndrome: contrasting developmental trajectories in those with and without elevated autism symptoms. Journal of Intellectual Disability Research, 2018, 62, 672-683.	2.0	10
44	Attenuated behaviour in Cornelia de Lange and fragile X syndromes. Journal of Intellectual Disability Research, 2018, 62, 486-495.	2.0	5
45	Service use and access in young children with an intellectual disability or global developmental delay: Associations with challenging behaviour. Journal of Intellectual and Developmental Disability, 2018, 43, 232-241.	1.6	7
46	Self-injurious behavior. Neuroscience and Biobehavioral Reviews, 2018, 84, 483-491.	6.1	57
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	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
49	Service receipt of adults with rare genetic syndromes who engage in challenging behaviour. Journal of Intellectual and Developmental Disability, 2018, 43, 308-316.	1.6	3
50	Interventions for mental health problems in children and adults with severe intellectual disabilities: a systematic review. BMJ Open, 2018, 8, e021911.	1.9	40
51	Diagnosis and management of Cornelia de Lange syndrome: first international consensus statement. Nature Reviews Genetics, 2018, 19, 649-666.	16.3	223
52	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
53	The behavioural phenotype of Potocki-Lupski syndrome: a cross-syndrome comparison. Journal of Neurodevelopmental Disorders, 2018, 10, 2.	3.1	14
54	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

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55	Brief Report: Repetitive Behaviour Profiles in Williams syndrome: Cross Syndrome Comparisons with Praderâ€“Willi and Down syndromes. Journal of Autism and Developmental Disorders, 2018, 48, 326-331.	2.7	14
56	A Comparison of Two Methods for Recruiting Children with an Intellectual Disability. Journal of Applied Research in Intellectual Disabilities, 2017, 30, 696-704.	2.0	4
57	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
58	Genotypeâ€“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
59	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
60	Self-injurious behaviour in people with intellectual disability and autism spectrum disorder. Current Opinion in Psychiatry, 2017, 30, 97-101.	6.3	27
61	Brief Report: Contrasting Profiles of Everyday Executive Functioning in Smithâ€“Magenis Syndrome and Down Syndrome. Journal of Autism and Developmental Disorders, 2017, 47, 2602-2609.	2.7	11
62	Phenotypes and genotypes in individuals with SMC1A variants. American Journal of Medical Genetics, Part A, 2017, 173, 2108-2125.	1.2	69
63	Diverse Profiles of Anxiety Related Disorders in Fragile X, Cornelia de Lange and Rubinsteinâ€“Taybi Syndromes. Journal of Autism and Developmental Disorders, 2017, 47, 3728-3740.	2.7	40
64	Behaviour in Cornelia de Lange syndrome: a systematic review. Developmental Medicine and Child Neurology, 2017, 59, 361-366.	2.1	20
65	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
66	Sleep in children with Angelman syndrome: Parental concerns and priorities. Research in Developmental Disabilities, 2017, 69, 105-115.	2.2	21
67	Associations between behaviours that challenge in adults with intellectual disability, parental perceptions and parental mental health. British Journal of Clinical Psychology, 2017, 56, 408-430.	3.5	6
68	Signalling changes to individuals who show resistance to change can reduce challenging behaviour. Journal of Behavior Therapy and Experimental Psychiatry, 2017, 54, 58-70.	1.2	9
69	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
70	The importance of knowing when to be precise.. Journal of Intellectual Disability Research, 2017, 61, 1079-1082.	2.0	4
71	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
72	Executive functioning in Cornelia de Lange syndrome: domain asynchrony and age-related performance. Journal of Neurodevelopmental Disorders, 2017, 9, 29.	3.1	17

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73	An experimental study of executive function and social impairment in Cornelia de Lange syndrome. Journal of Neurodevelopmental Disorders, 2017, 9, 33.	3.1	12
74	Prospective study of autism phenomenology and the behavioural phenotype of Phelanâ€“McDermid syndrome: comparison to fragile X syndrome, Down syndrome and idiopathic autism spectrum disorder. Journal of Neurodevelopmental Disorders, 2017, 9, 37.	3.1	25
75	Differences in Social Motivation in Children with Smithâ€“Magenis Syndrome and Down Syndrome. Journal of Autism and Developmental Disorders, 2016, 46, 2148-2159.	2.7	18
76	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
77	Neutrophil function in young and old caregivers. British Journal of Health Psychology, 2016, 21, 173-189.	3.5	9
78	JIDR: 60â€‰years of diversity. Journal of Intellectual Disability Research, 2016, 60, 1137-1139.	2.0	0
79	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
80	Visual preference for social stimuli in individuals with autism or neurodevelopmental disorders: an eye-tracking study. Molecular Autism, 2016, 7, 24.	4.9	41
81	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
82	Ageing in Rett syndrome. Journal of Intellectual Disability Research, 2016, 60, 182-190.	2.0	22
83	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
84	The behavioral characteristics of Sotos syndrome. American Journal of Medical Genetics, Part A, 2015, 167, 2945-2956.	1.2	25
85	Face scanning and spontaneous emotion preference in Cornelia de Lange syndrome and Rubinstein-Taybi syndrome. Journal of Neurodevelopmental Disorders, 2015, 7, 22.	3.1	5
86	Behavioral characteristics associated with 19p13.2 microdeletions. American Journal of Medical Genetics, Part A, 2015, 167, 2334-2343.	1.2	5
87	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
88	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
89	Repetitive Behavior in Rubinsteinâ€“Taybi Syndrome: Parallels with Autism Spectrum Phenomenology. Journal of Autism and Developmental Disorders, 2015, 45, 1238-1253.	2.7	32
90	An Informant Report Behavior Diary for Measuring Temper Outbursts in an Intervention Setting. Journal of Developmental and Physical Disabilities, 2015, 27, 489-504.	1.6	4

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91	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
92	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
93	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
94	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
95	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
96	A national survey of Rett syndrome: behavioural characteristics. Journal of Neurodevelopmental Disorders, 2015, 7, 11.	3.1	38
97	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
98	Anticytomegalovirus antibody titres are not associated with caregiving burden in younger caregivers. British Journal of Health Psychology, 2015, 20, 68-84.	3.5	4
99	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
100	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
101	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
102	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
103	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
104	Temper outbursts in <scp>P</scp>raderâ€“<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
105	Developmental trajectories in behavioural phenotypes. Journal of Intellectual Disability Research, 2014, 58, 881-882.	2.0	1
106	The Motivating Operation and Negatively Reinforced Problem Behavior. Behavior Modification, 2014, 38, 107-159.	1.6	15
107	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
108	Trends and synergies in intellectual disability research. Journal of Intellectual Disability Research, 2014, 58, 1-2.	2.0	0

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109	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
110	The prevalence of aggression in genetic syndromes: A review. Research in Developmental Disabilities, 2014, 35, 1051-1071.	2.2	43
111	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
112	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
113	The nature of social preference and interactions in Smithâ€“Magenis syndrome. Research in Developmental Disabilities, 2013, 34, 4355-4365.	2.2	23
114	Self-injurious, aggressive and destructive behaviour in young children with a moderate to profound intellectual disability. Paediatrics and Child Health (United Kingdom), 2013, 23, 322-324.	0.4	2
115	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
116	Prevalence of autism spectrum disorder symptomatology and related behavioural characteristics in individuals with Down syndrome. Autism, 2013, 17, 390-404.	4.1	81
117	Impact with conviction. Journal of Intellectual Disability Research, 2013, 57, 293-294.	2.0	1
118	Behavioural phenotypes and neurodevelopmental disorders in <scp>A</scp>frica. Journal of Intellectual Disability Research, 2013, 57, 793-795.	2.0	9
119	Use of the structured descriptive assessment to identify possible functions of challenging behaviour exhibited by adults with brain injury. Neuropsychological Rehabilitation, 2013, 23, 501-527.	1.6	8
120	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
121	Causal Models of Clinically Significant Behaviors in Angelman, Cornelia de Lange, Praderâ€“Willi and Smithâ€“Magenis Syndromes. International Review of Research in Developmental Disabilities, 2013, 44, 167-211.	0.8	34
122	Functional analysis and functional communication training in individuals with Angelman syndrome. Developmental Neurorehabilitation, 2012, 15, 91-104.	1.1	17
123	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
124	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
125	Selfâ€injurious behaviour in individuals with autism spectrum disorder and intellectual disability. Journal of Intellectual Disability Research, 2012, 56, 476-489.	2.0	142
126	Selfâ€injurious behaviour in people with intellectual disability. Journal of Intellectual Disability Research, 2012, 56, 421-426.	2.0	9

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127	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
128	A review of defining and measuring sociability in children with intellectual disabilities. Research in Developmental Disabilities, 2011, 32, 11-24.	2.2	36
129	Phenotypeâ€“environment interactions in genetic syndromes associated with severe or profound intellectual disability. Research in Developmental Disabilities, 2011, 32, 404-418.	2.2	34
130	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
131	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
132	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
133	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
134	Effects of adult familiarity on social behaviours in Angelman syndrome. Journal of Intellectual Disability Research, 2011, 55, 339-350.	2.0	29
135	The relationship between specific cognitive impairment and behaviour in Praderâ€“Willi syndrome. Journal of Intellectual Disability Research, 2011, 55, 152-171.	2.0	46
136	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
137	Prevalence, phenomenology, aetiology and predictors of challenging behaviour in Smithâ€Magenis syndrome. Journal of Intellectual Disability Research, 2011, 55, 138-151.	2.0	45
138	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
139	Behavioural phenotypes: from models to intervention. Journal of Intellectual Disability Research, 2011, 55, 945-947.	2.0	1
140	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
141	â€œYou Have to Sit and Explain it All, and Explain Yourself.â€•Mothersâ€™ 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
142	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
143	Further refinement of the nature of the communication impairment in Cornelia de Lange syndrome. Advances in Mental Health and Intellectual Disabilities, 2011, 5, 15-25.	1.1	2
144	Self-injurious behaviour in people with intellectual disability. Current Opinion in Psychiatry, 2010, 23, 412-416.	6.3	24

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145	Neural correlates of task switching in paternal 15q11â€“q13 deletion Praderâ€“Willi syndrome. Brain Research, 2010, 1363, 128-142.	2.2	46
146	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
147	Variability in genetic syndromes: Some lessons from history. Journal of Intellectual Disability Research, 2010, 54, 877-880.	2.0	0
148	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
149	Descriptive analysis of challenging behaviours shown by adults with acquired brain injury. Neuropsychological Rehabilitation, 2010, 20, 212-238.	1.6	14
150	Task-switching deficits and repetitive behaviour in genetic neurodevelopmental disorders: Data from children with Praderâ€“Willi syndrome chromosome 15 q11â€“q13 deletion and boys with Fragile X syndrome. Cognitive Neuropsychology, 2009, 26, 172-194.	1.1	84
151	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
152	Dorsal and ventral stream mediated visual processing in genetic subtypes of Praderâ€“Willi syndrome. Neuropsychologia, 2009, 47, 2367-2373.	1.6	19
153	Associations between repetitive questioning, resistance to change, temper outbursts and anxiety in Praderâ€“Willi and Fragileâ€X syndromes. Journal of Intellectual Disability Research, 2009, 53, 265-278.	2.0	67
154	A specific pathway can be identified between genetic characteristics and behaviour profiles in Praderâ€“Willi syndrome via cognitive, environmental and physiological mechanisms. Journal of Intellectual Disability Research, 2009, 53, 493-500.	2.0	32
155	Selfâ€injurious behaviour in Cornelia de Lange syndrome: 1. Prevalence and phenomenology. Journal of Intellectual Disability Research, 2009, 53, 575-589.	2.0	46
156	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
157	Intellectual disabilities and genetic disorders can lead the way in translational research. Journal of Intellectual Disability Research, 2009, 53, 829-830.	2.0	2
158	The Prevalence and Phenomenology of Repetitive Behavior in Genetic Syndromes. Journal of Autism and Developmental Disorders, 2009, 39, 572-588.	2.7	202
159	Social Anxiety in Cornelia de Lange Syndrome. Journal of Autism and Developmental Disorders, 2009, 39, 1155-1162.	2.7	39
160	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
161	Experimental functional analysis of aggression in children with Angelman syndrome. Research in Developmental Disabilities, 2009, 30, 1095-1106.	2.2	45
162	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

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163	Facial expression of affect in children with Cornelia de Lange syndrome. Journal of Intellectual Disability Research, 2008, 52, 207-215.	2.0	18
164	Behavioural characteristics associated with dementia assessment referrals in adults with Down syndrome. Journal of Intellectual Disability Research, 2008, 52, 358-368.	2.0	33
165	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
166	The behavioural phenotype of Smithâ€“Magenis syndrome: evidence for a geneâ€“environment interaction. Journal of Intellectual Disability Research, 2008, 52, 830-841.	2.0	49
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