

Chris Oliver

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

217
papers

6,466
citations

43
h-index

69
g-index

241
ext. papers

7,324
ext. citations

3.7
avg, IF

6.07
L-index

#	Paper	IF	Citations
217	Caregivers' experience of sleep management in Smith-Magenis syndrome: a mixed-methods study.. <i>Orphanet Journal of Rare Diseases</i> , 2022 , 17, 35	4.2	2
216	Executive function, repetitive behaviour and restricted interests in neurodevelopmental disorders.. <i>Research in Developmental Disabilities</i> , 2022 , 122, 104166	2.7	
215	The behavioural phenotype of SATB2-associated syndrome: a within-group and cross-syndrome analysis.. <i>Journal of Neurodevelopmental Disorders</i> , 2022 , 14, 25	4.6	
214	Prevalence of anxiety symptomatology and diagnosis in syndromic intellectual disability: a systematic review and meta-analysis. <i>Neuroscience and Biobehavioral Reviews</i> , 2022 , 104719	9	1
213	The development of early social cognitive skills in neurogenetic syndromes associated with autism: Cornelia de Lange, fragile X and Rubinstein-Taybi syndromes. <i>Orphanet Journal of Rare Diseases</i> , 2021 , 16, 488	4.2	0
212	Behaviour across the lifespan in Cornelia de Lange syndrome. <i>Current Opinion in Psychiatry</i> , 2021 , 34, 112-117	4.9	2
211	The adaptive functioning profile of Pitt-Hopkins syndrome. <i>European Journal of Medical Genetics</i> , 2021 , 64, 104279	2.6	
210	Skin Picking in People with Prader-Willi Syndrome: Phenomenology and Management. <i>Journal of Autism and Developmental Disorders</i> , 2021 , 51, 286-297	4.6	2
209	Genetic modifiers in rare disorders: the case of fragile X syndrome. <i>European Journal of Human Genetics</i> , 2021 , 29, 173-183	5.3	4
208	A systematic review of the behaviours associated with depression in people with severe-profound intellectual disability. <i>Journal of Intellectual Disability Research</i> , 2021 , 65, 211-229	3.2	3
207	Sleep disorders in rare genetic syndromes: a meta-analysis of prevalence and profile. <i>Molecular Autism</i> , 2021 , 12, 18	6.5	13
206	Refining the Behavioral Phenotype of Angelman Syndrome: Examining Differences in Motivation for Social Contact Between Genetic Subgroups. <i>Frontiers in Behavioral Neuroscience</i> , 2021 , 15, 618271	3.5	3
205	Anxiety characteristics in individuals with Williams syndrome. <i>Journal of Applied Research in Intellectual Disabilities</i> , 2021 , 34, 1098-1107	2.2	3
204	Low speech rate but high gesture rate during conversational interaction in people with Cornelia de Lange syndrome. <i>Journal of Intellectual Disability Research</i> , 2021 , 65, 601-607	3.2	1
203	An Observational Study of Social Interaction Skills and Behaviors in Cornelia de Lange, Fragile X and Rubinstein-Taybi Syndromes. <i>Journal of Autism and Developmental Disorders</i> , 2020 , 50, 4001-4010	4.6	5
202	Scaling of Early Social Cognitive Skills in Typically Developing Infants and Children with Autism Spectrum Disorder. <i>Journal of Autism and Developmental Disorders</i> , 2020 , 50, 3988-4000	4.6	3
201	Cornelia de Lange Syndrome 2020 , 129-157		

200	Persistence and predictors of self-injurious behaviour in autism: a ten-year prospective cohort study. <i>Molecular Autism</i> , 2020 , 11, 8	6.5	7
199	Prevalence and Risk-Markers of Self-Harm in Autistic Children and Adults. <i>Journal of Autism and Developmental Disorders</i> , 2020 , 50, 3561-3574	4.6	6
198	Sleep in children with Smith-Magenis syndrome: a case-control actigraphy study. <i>Sleep</i> , 2020 , 43,	1.1	8
197	Profiles of atypical sensory processing in Angelman, Cornelia de Lange and Fragile X syndromes. <i>Journal of Intellectual Disability Research</i> , 2020 , 64, 117-130	3.2	8
196	Sleep disorders in children with Angelman and Smith-Magenis syndromes: The assessment of potential causes of disrupted settling and night time waking. <i>Research in Developmental Disabilities</i> , 2020 , 97, 103555	2.7	8
195	The Profiles and Correlates of Psychopathology in Adolescents and Adults with Williams, Fragile X and Prader-Willi Syndromes. <i>Journal of Autism and Developmental Disorders</i> , 2020 , 50, 893-903	4.6	5
194	Fragile X syndrome: an overview of cause, characteristics, assessment and management. <i>Paediatrics and Child Health (United Kingdom)</i> , 2020 , 30, 400-403	0.6	1
193	The behaviour and wellbeing of children and adults with severe intellectual disability and complex needs: the Be-Well checklist for carers and professionals. <i>Paediatrics and Child Health (United Kingdom)</i> , 2020 , 30, 416-424	0.6	6
192	A Behavioural Assessment of Social Anxiety and Social Motivation in Fragile X, Cornelia de Lange and Rubinstein-Taybi Syndromes. <i>Journal of Autism and Developmental Disorders</i> , 2020 , 50, 127-144	4.6	16
191	Sleep problems in autism spectrum disorders: A comparison to sleep in typically developing children using actigraphy, diaries and questionnaires. <i>Research in Autism Spectrum Disorders</i> , 2019 , 67, 101439	3	5
190	Behavioural and psychological characteristics in Pitt-Hopkins syndrome: a comparison with Angelman and Cornelia de Lange syndromes. <i>Journal of Neurodevelopmental Disorders</i> , 2019 , 11, 24	4.6	5
189	Temper outbursts in Lowe syndrome: Characteristics, sequence, environmental context and comparison to Prader-Willi syndrome. <i>Journal of Applied Research in Intellectual Disabilities</i> , 2019 , 32, 1216-1227	2.2	4
188	Lifespan trajectory of affect in Cornelia de Lange syndrome: towards a neurobiological hypothesis. <i>Journal of Neurodevelopmental Disorders</i> , 2019 , 11, 6	4.6	5
187	Communication in Angelman syndrome: a scoping review. <i>Developmental Medicine and Child Neurology</i> , 2019 , 61, 1266-1274	3.3	8
186	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. <i>Journal of Autism and Developmental Disorders</i> , 2019 , 49, 2913-2922	4.6	10
185	Age-related Behavioural Change in Cornelia de Lange and Cri du Chat Syndromes: A Seven Year Follow-up Study. <i>Journal of Autism and Developmental Disorders</i> , 2019 , 49, 2476-2487	4.6	5
184	Multi-Method Assessment of Sleep in Children With Angelman Syndrome: A Case-Controlled Study. <i>Frontiers in Psychiatry</i> , 2019 , 10, 874	5	6
183	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. <i>PLoS ONE</i> , 2019 , 14, e0223750	3.7	10

182	Coping Strategies in Mothers of Children with Intellectual Disabilities Showing Multiple Forms of Challenging Behaviour: Associations with Maternal Mental Health. <i>Behavioural and Cognitive Psychotherapy</i> , 2018 , 46, 257-275	2.1	8
181	Persistence of self-injury, aggression and property destruction in children and adults with tuberous sclerosis complex. <i>Journal of Intellectual Disability Research</i> , 2018 , 62, 1058-1071	3.2	5
180	Differences in the Information Needs of Parents With a Child With a Genetic Syndrome: A Cross-Syndrome Comparison. <i>Journal of Policy and Practice in Intellectual Disabilities</i> , 2018 , 15, 94-100	1.8	5
179	Overactivity, impulsivity and repetitive behaviour in males with fragile X syndrome: contrasting developmental trajectories in those with and without elevated autism symptoms. <i>Journal of Intellectual Disability Research</i> , 2018 , 62, 672-683	3.2	7
178	Attenuated behaviour in Cornelia de Lange and fragile X syndromes. <i>Journal of Intellectual Disability Research</i> , 2018 , 62, 486-495	3.2	4
177	Service use and access in young children with an intellectual disability or global developmental delay: Associations with challenging behaviour. <i>Journal of Intellectual and Developmental Disability</i> , 2018 , 43, 232-241	1.9	5
176	Self-injurious behavior. <i>Neuroscience and Biobehavioral Reviews</i> , 2018 , 84, 483-491	9	36
175	Interventions for mental health problems in children and adults with severe intellectual disabilities: a systematic review. <i>BMJ Open</i> , 2018 , 8, e021911	3	19
174	Diagnosis and management of Cornelia de Lange syndrome: first international consensus statement. <i>Nature Reviews Genetics</i> , 2018 , 19, 649-666	30.1	123
173	A cross-syndrome cohort comparison of sleep disturbance in children with Smith-Magenis syndrome, Angelman syndrome, autism spectrum disorder and tuberous sclerosis complex. <i>Journal of Neurodevelopmental Disorders</i> , 2018 , 10, 9	4.6	39
172	The behavioural phenotype of Potocki-Lupski syndrome: a cross-syndrome comparison. <i>Journal of Neurodevelopmental Disorders</i> , 2018 , 10, 2	4.6	9
171	Mental Health and Well-Being in Mothers of Children With Rare Genetic Syndromes Showing Chronic Challenging Behavior: A Cross-Sectional and Longitudinal Study. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2018 , 123, 241-253	2.2	8
170	Brief Report: Repetitive Behaviour Profiles in Williams syndrome: Cross Syndrome Comparisons with Prader-Willi and Down syndromes. <i>Journal of Autism and Developmental Disorders</i> , 2018 , 48, 326-331	4.6	9
169	Sleep duration and sleep quality in people with and without intellectual disability: A meta-analysis. <i>Sleep Medicine Reviews</i> , 2018 , 40, 135-150	10.2	32
168	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. <i>Orphanet Journal of Rare Diseases</i> , 2018 , 13, 185	4.2	11
167	Service receipt of adults with rare genetic syndromes who engage in challenging behaviour. <i>Journal of Intellectual and Developmental Disability</i> , 2018 , 43, 308-316	1.9	1
166	A Comparison of Two Methods for Recruiting Children with an Intellectual Disability. <i>Journal of Applied Research in Intellectual Disabilities</i> , 2017 , 30, 696-704	2.2	4
165	Predictors of Self-Injurious Behavior and Self-Restraint in Autism Spectrum Disorder: Towards a Hypothesis of Impaired Behavioral Control. <i>Journal of Autism and Developmental Disorders</i> , 2017 , 47, 701-713	4.6	22

164	Genotype-phenotype correlations in Cornelia de Lange syndrome: Behavioral characteristics and changes with age. <i>American Journal of Medical Genetics, Part A</i> , 2017 , 173, 1566-1574	2.5	16
163	Self-injury and aggression in adults with tuberous sclerosis complex: Frequency, associated person characteristics, and implications for assessment. <i>Research in Developmental Disabilities</i> , 2017 , 64, 119-130	2.7	10
162	Self-injurious behaviour in people with intellectual disability and autism spectrum disorder. <i>Current Opinion in Psychiatry</i> , 2017 , 30, 97-101	4.9	17
161	Brief Report: Contrasting Profiles of Everyday Executive Functioning in Smith-Magenis Syndrome and Down Syndrome. <i>Journal of Autism and Developmental Disorders</i> , 2017 , 47, 2602-2609	4.6	8
160	Phenotypes and genotypes in individuals with SMC1A variants. <i>American Journal of Medical Genetics, Part A</i> , 2017 , 173, 2108-2125	2.5	44
159	Diverse Profiles of Anxiety Related Disorders in Fragile X, Cornelia de Lange and Rubinstein-Taybi Syndromes. <i>Journal of Autism and Developmental Disorders</i> , 2017 , 47, 3728-3740	4.6	24
158	Behaviour in Cornelia de Lange syndrome: a systematic review. <i>Developmental Medicine and Child Neurology</i> , 2017 , 59, 361-366	3.3	16
157	Differential effects of anxiety and autism on social scene scanning in males with fragile X syndrome. <i>Journal of Neurodevelopmental Disorders</i> , 2017 , 9, 9	4.6	6
156	Executive functioning in Cornelia de Lange syndrome: domain asynchrony and age-related performance. <i>Journal of Neurodevelopmental Disorders</i> , 2017 , 9, 29	4.6	15
155	An experimental study of executive function and social impairment in Cornelia de Lange syndrome. <i>Journal of Neurodevelopmental Disorders</i> , 2017 , 9, 33	4.6	8
154	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. <i>Journal of Neurodevelopmental Disorders</i> , 2017 , 9, 37	4.6	16
153	Measurement tools for mental health problems and mental well-being in people with severe or profound intellectual disabilities: A systematic review. <i>Clinical Psychology Review</i> , 2017 , 57, 32-44	10.8	46
152	Sleep in children with Angelman syndrome: Parental concerns and priorities. <i>Research in Developmental Disabilities</i> , 2017 , 69, 105-115	2.7	13
151	Associations between behaviours that challenge in adults with intellectual disability, parental perceptions and parental mental health. <i>British Journal of Clinical Psychology</i> , 2017 , 56, 408-430	3.6	5
150	Signalling changes to individuals who show resistance to change can reduce challenging behaviour. <i>Journal of Behavior Therapy and Experimental Psychiatry</i> , 2017 , 54, 58-70	2.6	8
149	Anxiety Disorders in Williams Syndrome Contrasted with Intellectual Disability and the General Population: A Systematic Review and Meta-Analysis. <i>Journal of Autism and Developmental Disorders</i> , 2017 , 47, 3765-3777	4.6	39
148	A Comparative Study of Sociability in Angelman, Cornelia de Lange, Fragile X, Down and Rubinstein Taybi Syndromes and Autism Spectrum Disorder. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2016 , 121, 465-486	2.2	30
147	Visual preference for social stimuli in individuals with autism or neurodevelopmental disorders: an eye-tracking study. <i>Molecular Autism</i> , 2016 , 7, 24	6.5	25

146	Dissociation of Cross-Sectional Trajectories for Verbal and Visuo-Spatial Working Memory Development in Rubinstein-Taybi Syndrome. <i>Journal of Autism and Developmental Disorders</i> , 2016 , 46, 2064-2071	4.6	8
145	Ageing in Rett syndrome. <i>Journal of Intellectual Disability Research</i> , 2016 , 60, 182-90	3.2	15
144	Self-injury, aggression and destruction in children with severe intellectual disability: Incidence, persistence and novel, predictive behavioural risk markers. <i>Research in Developmental Disabilities</i> , 2016 , 49-50, 291-301	2.7	29
143	Differences in Social Motivation in Children with Smith-Magenis Syndrome and Down Syndrome. <i>Journal of Autism and Developmental Disorders</i> , 2016 , 46, 2148-2159	4.6	13
142	Persistence of self-injurious behaviour in autism spectrum disorder over 3½ years: a prospective cohort study of risk markers. <i>Journal of Neurodevelopmental Disorders</i> , 2016 , 8, 21	4.6	33
141	Neutrophil function in young and old caregivers. <i>British Journal of Health Psychology</i> , 2016 , 21, 173-89	8.3	8
140	Are Angelman and Prader-Willi syndromes more similar than we thought? Food-related behavior problems in Angelman, Cornelia de Lange, fragile X, Prader-Willi and 1p36 deletion syndromes. <i>American Journal of Medical Genetics, Part A</i> , 2015 , 167A, 572-8	2.5	9
139	Increased Exposure to Rigid Routines can Lead to Increased Challenging Behavior Following Changes to Those Routines. <i>Journal of Autism and Developmental Disorders</i> , 2015 , 45, 1569-78	4.6	10
138	Positive impact and its relationship to well-being in parents of children with intellectual disability: a literature review. <i>International Journal of Developmental Disabilities</i> , 2015 , 61, 1-19	1.5	33
137	Brief Report: A Longitudinal Study of Excessive Smiling and Laughing in Children with Angelman Syndrome. <i>Journal of Autism and Developmental Disorders</i> , 2015 , 45, 2624-7	4.6	12
136	A national survey of Rett syndrome: behavioural characteristics. <i>Journal of Neurodevelopmental Disorders</i> , 2015 , 7, 11	4.6	24
135	A national survey of Rett syndrome: age, clinical characteristics, current abilities, and health. <i>American Journal of Medical Genetics, Part A</i> , 2015 , 167, 1493-500	2.5	21
134	Anticytomegalovirus antibody titres are not associated with caregiving burden in younger caregivers. <i>British Journal of Health Psychology</i> , 2015 , 20, 68-84	8.3	2
133	Prevalence of autism spectrum disorder phenomenology in genetic disorders: a systematic review and meta-analysis. <i>Lancet Psychiatry</i> , 2015 , 2, 909-16	23.3	195
132	Self-injurious, aggressive and destructive behaviour in children with severe intellectual disability: Prevalence, service need and service receipt in the UK. <i>Research in Developmental Disabilities</i> , 2015 , 45-46, 307-15	2.7	23
131	The behavioral characteristics of Sotos syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2015 , 167A, 2945-56	2.5	17
130	Face scanning and spontaneous emotion preference in Cornelia de Lange syndrome and Rubinstein-Taybi syndrome. <i>Journal of Neurodevelopmental Disorders</i> , 2015 , 7, 22	4.6	3
129	Behavioral characteristics associated with 19p13.2 microdeletions. <i>American Journal of Medical Genetics, Part A</i> , 2015 , 167A, 2334-43	2.5	5

128	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 follow-up. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2015 , 169, 188-97	3.1	15
127	Practitioner Review: Self-injurious behaviour in children with developmental delay. <i>Journal of Child Psychology and Psychiatry and Allied Disciplines</i> , 2015 , 56, 1042-54	7.9	32
126	Repetitive behavior in Rubinstein-Taybi syndrome: parallels with autism spectrum phenomenology. <i>Journal of Autism and Developmental Disorders</i> , 2015 , 45, 1238-53	4.6	23
125	An Informant Report Behavior Diary for Measuring Temper Outbursts in an Intervention Setting. <i>Journal of Developmental and Physical Disabilities</i> , 2015 , 27, 489-504	1.5	4
124	Implicit Discrimination of Basic Facial Expressions of Positive/Negative Emotion in Fragile X Syndrome and Autism Spectrum Disorder. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2015 , 120, 328-45	2.2	14
123	Correlates of self-injurious, aggressive and destructive behaviour in children under five who are at risk of developmental delay. <i>Research in Developmental Disabilities</i> , 2014 , 35, 36-45	2.7	14
122	Self-injury and aggression in tuberous sclerosis complex: cross syndrome comparison and associated risk markers. <i>Journal of Neurodevelopmental Disorders</i> , 2014 , 6, 10	4.6	29
121	The prevalence of aggression in genetic syndromes: a review. <i>Research in Developmental Disabilities</i> , 2014 , 35, 1051-71	2.7	37
120	A longitudinal follow-up study of affect in children and adults with Cornelia de Lange syndrome. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2014 , 119, 235-52	2.2	13
119	The importance of understanding the behavioural phenotypes of genetic syndromes associated with intellectual disability. <i>Paediatrics and Child Health (United Kingdom)</i> , 2014 , 24, 468-472	0.6	27
118	The purported association between depression, aggression, and self-injury in people with intellectual disability: a critical review of the literature. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2014 , 119, 452-71	2.2	16
117	Temper outbursts in Prader-Willi syndrome: causes, behavioural and emotional sequence and responses by carers. <i>Journal of Intellectual Disability Research</i> , 2014 , 58, 134-50	3.2	33
116	The motivating operation and negatively reinforced problem behavior: a systematic review. <i>Behavior Modification</i> , 2014 , 38, 107-59	2.5	12
115	The age related prevalence of aggression and self-injury in persons with an intellectual disability: a review. <i>Research in Developmental Disabilities</i> , 2013 , 34, 764-75	2.7	44
114	Discrimination training reduces high rate social approach behaviors in Angelman syndrome: proof of principle. <i>Research in Developmental Disabilities</i> , 2013 , 34, 1794-803	2.7	16
113	The nature of social preference and interactions in Smith-Magenis syndrome. <i>Research in Developmental Disabilities</i> , 2013 , 34, 4355-65	2.7	17
112	Self-injurious, aggressive and destructive behaviour in young children with a moderate to profound intellectual disability. <i>Paediatrics and Child Health (United Kingdom)</i> , 2013 , 23, 322-324	0.6	2
111	Delineating the profile of autism spectrum disorder characteristics in Cornelia de Lange and Fragile X syndromes. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2013 , 118, 55-73	2.2	42

110	Prevalence of autism spectrum disorder symptomatology and related behavioural characteristics in individuals with Down syndrome. <i>Autism</i> , 2013 , 17, 390-404	6.6	59
109	Use of the structured descriptive assessment to identify possible functions of challenging behaviour exhibited by adults with brain injury. <i>Neuropsychological Rehabilitation</i> , 2013 , 23, 501-27	3.1	7
108	Social behavior and characteristics of autism spectrum disorder in Angelman, Cornelia de Lange, and Cri du Chat syndromes. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2013 , 118, 262-83	2.2	30
107	Causal Models of Clinically Significant Behaviors in Angelman, Cornelia de Lange, Prader-Willi and Smith-Magenis Syndromes. <i>International Review of Research in Developmental Disabilities</i> , 2013 , 44, 167-211	3.1	30
106	Self-injurious behaviour in individuals with autism spectrum disorder and intellectual disability. <i>Journal of Intellectual Disability Research</i> , 2012 , 56, 476-89	3.2	116
105	Characteristics of autism spectrum disorder in Cornelia de Lange syndrome. <i>Journal of Child Psychology and Psychiatry and Allied Disciplines</i> , 2012 , 53, 883-91	7.9	46
104	The association between repetitive, self-injurious and aggressive behavior in children with severe intellectual disability. <i>Journal of Autism and Developmental Disorders</i> , 2012 , 42, 910-9	4.6	70
103	Functional analysis and functional communication training in individuals with Angelman syndrome. <i>Developmental Neurorehabilitation</i> , 2012 , 15, 91-104	1.8	15
102	The expression and assessment of emotions and internal states in individuals with severe or profound intellectual disabilities. <i>Clinical Psychology Review</i> , 2011 , 31, 293-306	10.8	64
101	A review of defining and measuring sociability in children with intellectual disabilities. <i>Research in Developmental Disabilities</i> , 2011 , 32, 11-24	2.7	29
100	Phenotype-environment interactions in genetic syndromes associated with severe or profound intellectual disability. <i>Research in Developmental Disabilities</i> , 2011 , 32, 404-18	2.7	30
99	The Chronicity of Self-Injurious Behaviour: A Long-Term Follow-Up of a Total Population Study. <i>Journal of Applied Research in Intellectual Disabilities</i> , 2011 , 24, 105-117	2.2	43
98	Behavioural Excesses and Deficits Associated with Dementia in Adults who have Down Syndrome. <i>Journal of Applied Research in Intellectual Disabilities</i> , 2011 , 24, 208-216	2.2	15
97	The prevalence and phenomenology of self-injurious and aggressive behaviour in genetic syndromes. <i>Journal of Intellectual Disability Research</i> , 2011 , 55, 109-20	3.2	167
96	Low mood and challenging behaviour in people with severe and profound intellectual disabilities. <i>Journal of Intellectual Disability Research</i> , 2011 , 55, 182-9	3.2	21
95	Effects of adult familiarity on social behaviours in Angelman syndrome. <i>Journal of Intellectual Disability Research</i> , 2011 , 55, 339-50	3.2	21
94	The relationship between specific cognitive impairment and behaviour in Prader-Willi syndrome. <i>Journal of Intellectual Disability Research</i> , 2011 , 55, 152-71	3.2	39
93	Stereotyped behaviour in children with autism and intellectual disability: an examination of the executive dysfunction hypothesis. <i>Journal of Intellectual Disability Research</i> , 2011 , 55, 699-709	3.2	10

92	Prevalence, phenomenology, aetiology and predictors of challenging behaviour in Smith-Magenis syndrome. <i>Journal of Intellectual Disability Research</i> , 2011 , 55, 138-51	3.2	39
91	Psychological well-being in parents of children with Angelman, Cornelia de Lange and Cri du Chat syndromes. <i>Journal of Intellectual Disability Research</i> , 2011 , 55, 397-410	3.2	52
90	Delineation of behavioral phenotypes in genetic syndromes: characteristics of autism spectrum disorder, affect and hyperactivity. <i>Journal of Autism and Developmental Disorders</i> , 2011 , 41, 1019-32	4.6	70
89	"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. <i>Journal of Genetic Counseling</i> , 2011 , 20, 165-77	2.5	30
88	Age related change in social behavior in children with Angelman syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2011 , 155A, 1290-7	2.5	20
87	Further refinement of the nature of the communication impairment in Cornelia de Lange syndrome. <i>Advances in Mental Health and Intellectual Disabilities</i> , 2011 , 5, 15-25	0.5	2
86	The Assessment and Presentation of Autism Spectrum Disorder and Associated Characteristics in Individuals with Severe Intellectual Disability and Genetic Syndromes 2011 ,		9
85	Descriptive analysis of challenging behaviours shown by adults with acquired brain injury. <i>Neuropsychological Rehabilitation</i> , 2010 , 20, 212-38	3.1	12
84	Self-injurious behaviour in people with intellectual disability. <i>Current Opinion in Psychiatry</i> , 2010 , 23, 412-69	4.9	21
83	Neural correlates of task switching in paternal 15q11-q13 deletion Prader-Willi syndrome. <i>Brain Research</i> , 2010 , 1363, 128-42	3.7	40
82	The relationship between acquired impairments of executive function and behaviour change in adults with Down syndrome. <i>Journal of Intellectual Disability Research</i> , 2010 , 54, 393-405	3.2	55
81	The association between repetitive behaviours, impulsivity and hyperactivity in people with intellectual disability. <i>Journal of Intellectual Disability Research</i> , 2010 , 54, 1078-92	3.2	46
80	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. <i>Cognitive Neuropsychology</i> , 2009 , 26, 172-94	2.3	75
79	Coping with challenges to memory in people with mild to moderate Alzheimer's disease: observation of behaviour in response to analogues of everyday situations. <i>Aging and Mental Health</i> , 2009 , 13, 46-53	3.5	10
78	Dorsal and ventral stream mediated visual processing in genetic subtypes of Prader-Willi syndrome. <i>Neuropsychologia</i> , 2009 , 47, 2367-73	3.2	19
77	Associations between repetitive questioning, resistance to change, temper outbursts and anxiety in Prader-Willi and Fragile-X syndromes. <i>Journal of Intellectual Disability Research</i> , 2009 , 53, 265-78	3.2	57
76	A specific pathway can be identified between genetic characteristics and behaviour profiles in Prader-Willi syndrome via cognitive, environmental and physiological mechanisms. <i>Journal of Intellectual Disability Research</i> , 2009 , 53, 493-500	3.2	29
75	Self-injurious behaviour in Cornelia de Lange syndrome: 1. Prevalence and phenomenology. <i>Journal of Intellectual Disability Research</i> , 2009 , 53, 575-89	3.2	42

74	Self-injurious behaviour in Cornelia de Lange syndrome: 2. Association with environmental events. <i>Journal of Intellectual Disability Research</i> , 2009 , 53, 590-603	3.2	18
73	The prevalence and phenomenology of repetitive behavior in genetic syndromes. <i>Journal of Autism and Developmental Disorders</i> , 2009 , 39, 572-88	4.6	167
72	Social anxiety in Cornelia de Lange syndrome. <i>Journal of Autism and Developmental Disorders</i> , 2009 , 39, 1155-62	4.6	33
71	The Relationship between Components of the Behavioural Phenotype in Prader-Willi Syndrome. <i>Journal of Applied Research in Intellectual Disabilities</i> , 2009 , 22, 403-407	2.2	9
70	Experimental functional analysis of aggression in children with Angelman syndrome. <i>Research in Developmental Disabilities</i> , 2009 , 30, 1095-106	2.7	39
69	Relationship among challenging, repetitive, and communicative behaviors in children with severe intellectual disabilities. <i>American Journal on Intellectual and Developmental Disabilities</i> , 2009 , 114, 356-68	2.2	18
68	Facial expression of affect in children with Cornelia de Lange syndrome. <i>Journal of Intellectual Disability Research</i> , 2008 , 52, 207-15	3.2	15
67	Behavioural characteristics associated with dementia assessment referrals in adults with Down syndrome. <i>Journal of Intellectual Disability Research</i> , 2008 , 52, 358-68	3.2	28
66	Health and sleep problems in Cornelia de Lange Syndrome: a case control study. <i>Journal of Intellectual Disability Research</i> , 2008 , 52, 458-68	3.2	48
65	The behavioural phenotype of Smith-Magenis syndrome: evidence for a gene-environment interaction. <i>Journal of Intellectual Disability Research</i> , 2008 , 52, 830-41	3.2	43
64	The relationship between a deficit in attention switching and specific behaviours in Prader-Willi syndrome. <i>Journal of Intellectual Disability Research</i> , 2008 , 52, 812-812	3.2	0
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