

Allan Colver

List of Publications by Year in descending order

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Version: 2024-02-01

73
papers

5,155
citations

109137

35
h-index

91712

69
g-index

73
all docs

73
docs citations

73
times ranked

4264
citing authors

#	ARTICLE	IF	CITATIONS
1	Developing the FEEDS toolkit of parent-delivered interventions for eating, drinking and swallowing difficulties in young children with neurodisability: findings from a Delphi survey and stakeholder consultation workshops. <i>BMJ Paediatrics Open</i> , 2022, 6, e001425.	0.6	3
2	Disparity of child/parent-reported quality of life in cerebral palsy persists into adolescence. <i>Developmental Medicine and Child Neurology</i> , 2021, 63, 68-74.	1.1	16
3	Parent-delivered interventions used at home to improve eating, drinking and swallowing in children with neurodisability: the FEEDS mixed-methods study. <i>Health Technology Assessment</i> , 2021, 25, 1-208.	1.3	14
4	Children with neurodisability and feeding difficulties: a UK survey of parent-delivered interventions. <i>BMJ Paediatrics Open</i> , 2021, 5, e001095.	0.6	5
5	Facilitating transition of young people with long-term health conditions from children's to adults' healthcare services – implications of a 5-year research programme. <i>Clinical Medicine</i> , 2020, 20, 74-80.	0.8	31
6	Facilitating the transition of young people with long-term conditions through health services from childhood to adulthood: the Transition research programme. <i>Programme Grants for Applied Research</i> , 2019, 7, 1-244.	0.4	21
7	Falling through the gaps: exploring the role of integrated commissioning in improving transition from children's to adults' services for young people with long-term health conditions in England. <i>Journal of Health Services Research and Policy</i> , 2018, 23, 107-115.	0.8	9
8	Are the health needs of young people with cerebral palsy met during transition from child to adult health care?. <i>Child: Care, Health and Development</i> , 2018, 44, 355-363.	0.8	38
9	Understanding frames: A qualitative study of young people's experiences of using standing frames as part of postural management for cerebral palsy. <i>Child: Care, Health and Development</i> , 2018, 44, 203-211.	0.8	14
10	Understanding frames: A UK survey of parents and professionals regarding the use of standing frames for children with cerebral palsy. <i>Child: Care, Health and Development</i> , 2018, 44, 195-202.	0.8	18
11	A longitudinal, observational study examining the relationships of patient satisfaction with services and mental well-being to their clinical course in young people with Type 1 diabetes mellitus during transition from child to adult health services. <i>Diabetic Medicine</i> , 2018, 35, 1216-1222.	1.2	8
12	A longitudinal, observational study of the features of transitional healthcare associated with better outcomes for young people with long-term conditions. <i>BMC Medicine</i> , 2018, 16, 111.	2.3	91
13	Transition of young people with disability. <i>Paediatrics and Child Health (United Kingdom)</i> , 2018, 28, 374-378.	0.2	1
14	How well do services for young people with long term conditions deliver features proposed to improve transition?. <i>BMC Health Services Research</i> , 2018, 18, 337.	0.9	35
15	Standing frames for children with cerebral palsy: a mixed-methods feasibility study. <i>Health Technology Assessment</i> , 2018, 22, 1-232.	1.3	10
16	Predictors of parent-reported quality of life of adolescents with cerebral palsy: A longitudinal study. <i>Research in Developmental Disabilities</i> , 2017, 62, 259-270.	1.2	33
17	How available to European children and young people with cerebral palsy are features of their environment that they need?. <i>Research in Developmental Disabilities</i> , 2017, 71, 1-10.	1.2	6
18	What constitutes successful commissioning of transition from children's to adults' services for young people with long-term conditions and what are the challenges? An interview study. <i>BMJ Paediatrics Open</i> , 2017, 1, e000085.	0.6	8

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19	Views of Young People With Chronic Conditions on Transition From Pediatric to Adult Health Services. <i>Journal of Adolescent Health</i> , 2016, 59, 345-353.	1.2	34
20	Classification systems of communication for use in epidemiological surveillance of children with cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 285-291.	1.1	37
21	Cerebral palsy. <i>Nature Reviews Disease Primers</i> , 2016, 2, 15082.	18.1	603
22	Characteristics of young people with long term conditions close to transfer to adult health services. <i>BMC Health Services Research</i> , 2015, 15, 435.	0.9	18
23	Predictors of participation of adolescents with cerebral palsy: A European multi-centre longitudinal study. <i>Research in Developmental Disabilities</i> , 2015, 36, 551-564.	1.2	45
24	Self-reported quality of life of adolescents with cerebral palsy: a cross-sectional and longitudinal analysis. <i>Lancet, The</i> , 2015, 385, 705-716.	6.3	148
25	European study of frequency of participation of adolescents with and without cerebral palsy. <i>European Journal of Paediatric Neurology</i> , 2014, 18, 282-294.	0.7	94
26	Cerebral palsy. <i>Lancet, The</i> , 2014, 383, 1240-1249.	6.3	423
27	Stability of motor function and associated impairments between childhood and adolescence in young people with cerebral palsy in Europe. <i>Developmental Medicine and Child Neurology</i> , 2014, 56, 833-838.	1.1	17
28	Development of The Viking Speech Scale to classify the speech of children with cerebral palsy. <i>Research in Developmental Disabilities</i> , 2013, 34, 3202-3210.	1.2	132
29	Pain in young people aged 13 to 17 years with cerebral palsy: cross-sectional, multicentre European study. <i>Archives of Disease in Childhood</i> , 2013, 98, 434-440.	1.0	141
30	New understanding of adolescent brain development: relevance to transitional healthcare for young people with long term conditions. <i>Archives of Disease in Childhood</i> , 2013, 98, 902-907.	1.0	101
31	Risk and causes of death in children with a seizure disorder. <i>Developmental Medicine and Child Neurology</i> , 2012, 54, 612-617.	1.1	16
32	Association Between Participation in Life Situations of Children With Cerebral Palsy and Their Physical, Social, and Attitudinal Environment: A Cross-Sectional Multicenter European Study. <i>Archives of Physical Medicine and Rehabilitation</i> , 2012, 93, 2154-2164.	0.5	86
33	Quantifying the physical, social and attitudinal environment of children with cerebral palsy. <i>Disability and Rehabilitation</i> , 2011, 33, 36-50.	0.9	36
34	Access of children with cerebral palsy to the physical, social and attitudinal environment they need: a cross-sectional European study. <i>Disability and Rehabilitation</i> , 2011, 33, 28-35.	0.9	65
35	How can we help develop the evidence base in paediatric neurodisability?. <i>Child: Care, Health and Development</i> , 2011, 37, 611-612.	0.8	1
36	Parenting stress and children with cerebral palsy: a European cross-sectional survey. <i>Developmental Medicine and Child Neurology</i> , 2011, 53, 815-821.	1.1	118

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37	Cerebral palsy rates by birth weight, gestation and severity in North of England, 1991-2000 singleton births. <i>Archives of Disease in Childhood</i> , 2011, 96, 180-185.	1.0	26
38	Trends in prevalence of cerebral palsy in children born with a birthweight of 2,500 g or over in Europe from 1980 to 1998. <i>European Journal of Epidemiology</i> , 2010, 25, 635-642.	2.5	63
39	Congenital anomalies in children with cerebral palsy: a population-based record linkage study. <i>Developmental Medicine and Child Neurology</i> , 2010, 52, 345-351.	1.1	62
40	Why are children with cerebral palsy more likely to have emotional and behavioural difficulties?. <i>Developmental Medicine and Child Neurology</i> , 2010, 52, 986-986.	1.1	10
41	Variation at local government level in the support for families of severely disabled children and the factors that affect it. <i>Developmental Medicine and Child Neurology</i> , 2010, 52, e259-66.	1.1	8
42	Socio-economic inequalities in cerebral palsy prevalence in the United Kingdom: a register-based study. <i>Paediatric and Perinatal Epidemiology</i> , 2010, 24, 149-155.	0.8	26
43	Daslne: the challenge of developing a regional database for autism spectrum disorder. <i>Archives of Disease in Childhood</i> , 2009, 94, 38-41.	1.0	29
44	Psychological problems in children with hemiplegia: a European multicentre survey. <i>Archives of Disease in Childhood</i> , 2009, 94, 429-433.	1.0	31
45	Participation in life situations of 8-12 year old children with cerebral palsy: cross sectional European study. <i>BMJ: British Medical Journal</i> , 2009, 338, b1458-b1458.	2.4	203
46	Frequency of participation of 8-12-year-old children with cerebral palsy: A multi-centre cross-sectional European study. <i>European Journal of Paediatric Neurology</i> , 2009, 13, 165-177.	0.7	116
47	Quality of life and participation. <i>Developmental Medicine and Child Neurology</i> , 2009, 51, 656-659.	1.1	45
48	Rasch Measurement Properties of the KIDSCREEN Quality of Life Instrument in Children with Cerebral Palsy and Differential Item Functioning between Children with and without Cerebral Palsy. <i>Value in Health</i> , 2009, 12, 782-792.	0.1	37
49	Parent and professional reports of the quality of life of children with cerebral palsy and associated intellectual impairment. <i>Developmental Medicine and Child Neurology</i> , 2008, 50, 618-624.	1.1	61
50	Psychological problems in children with cerebral palsy: a cross-sectional European study. <i>Journal of Child Psychology and Psychiatry and Allied Disciplines</i> , 2008, 49, 405-413.	3.1	173
51	Validity and reliability of the guidelines of the Surveillance of Cerebral Palsy in Europe for the classification of cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2008, 50, 828-831.	1.1	78
52	Health status of children with cerebral palsy living in Europe: a multi-centre study. <i>Child: Care, Health and Development</i> , 2008, 34, 806-814.	0.8	38
53	Parent-Reported Quality of Life of Children With Cerebral Palsy in Europe. <i>Pediatrics</i> , 2008, 121, 54-64.	1.0	156
54	Reliability and Validity of the Child Health Questionnaire-PF-50 for European Children with Cerebral Palsy. <i>Journal of Pediatric Psychology</i> , 2007, 34, 41-50.	1.1	22

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55	Self-reported quality of life of 12-year-old children with cerebral palsy: a cross-sectional European study. <i>Lancet, The</i> , 2007, 369, 2171-2178.	6.3	398
56	Determinants of Child-Parent Agreement in Quality-of-Life Reports: A European Study of Children With Cerebral Palsy. <i>Pediatrics</i> , 2007, 120, e804-e814.	1.0	192
57	Participation of young severely disabled children is influenced by their intrinsic impairments and environment. <i>Developmental Medicine and Child Neurology</i> , 2007, 49, 345-349.	1.1	79
58	Classification of cerebral palsy: paediatric perspective. <i>Developmental Medicine and Child Neurology</i> , 2007, 49, 15-16.	1.1	4
59	Recommendations from the SCPE collaborative group for defining and classifying cerebral palsy. <i>Developmental Medicine and Child Neurology</i> , 2007, 49, 35-38.	1.1	213
60	Study protocol: SPARCLE – a multi-centre European study of the relationship of environment to participation and quality of life in children with cerebral palsy. <i>BMC Public Health</i> , 2006, 6, 105.	1.2	117
61	What are we trying to do for disabled children?. <i>Current Paediatrics</i> , 2006, 16, 501-505.	0.2	14
62	How might districts identify local barriers to participation for children with cerebral palsy?. <i>Public Health</i> , 2006, 120, 167-175.	1.4	42
63	Discussion groups with parents of children with cerebral palsy in Europe designed to assist development of a relevant measure of environment. <i>Child: Care, Health and Development</i> , 2006, 32, 185-192.	0.8	42
64	Assessment of data quality in a multi-centre cross-sectional study of participation and quality of life of children with cerebral palsy. <i>BMC Public Health</i> , 2006, 6, 273.	1.2	36
65	A qualitative study of the physical, social and attitudinal environments influencing the participation of children with cerebral palsy in northeast England. <i>Developmental Neurorehabilitation</i> , 2006, 9, 219-228.	1.1	88
66	Are the dangers of childhood food allergy exaggerated?. <i>BMJ: British Medical Journal</i> , 2006, 333, 494-496.	2.4	18
67	A shared framework and language for childhood disability. <i>Developmental Medicine and Child Neurology</i> , 2005, 47, 780.	1.1	49
68	Subjective quality of life in children with intellectual impairment – how can it be assessed?. <i>Developmental Medicine and Child Neurology</i> , 2005, 47, 281-285.	1.1	77
69	Regional Variation in Survival of People With Cerebral Palsy in the United Kingdom. <i>Pediatrics</i> , 2005, 116, 1383-1390.	1.0	47
70	A qualitative study, using focused interviews, of the information needs of families whose children's names are on a cerebral palsy register. <i>Child: Care, Health and Development</i> , 2003, 29, 465-471.	0.8	9
71	The term diplegia should be abandoned. <i>Archives of Disease in Childhood</i> , 2003, 88, 286-290.	1.0	52
72	Benefits of a population register of children with cerebral palsy. <i>Indian Pediatrics</i> , 2003, 40, 639-44.	0.2	4

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73	Measurement of health status and quality of life in neonatal follow-up studies. <i>Seminars in Fetal and Neonatal Medicine</i> , 2000, 5, 149-157.	2.8	14