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Childhood cognitive development in 22q11.2 deletion syndrome: case-control study

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British Journal of Psychiatry, 2017, 211, 223-230.

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#	Paper	IF	Citations
30	Neurodevelopmental outcome in 22q11.2 deletion syndrome and management. <i>American Journal of Medical Genetics, Part A</i> , 2018 , 176, 2160-2166	2.5	17
29	Childhood Executive Functioning Predicts Young Adult Outcomes in 22q11.2 Deletion Syndrome. <i>Journal of the International Neuropsychological Society</i> , 2018 , 24, 905-916	3.1	7
28	Exploratory study on cognitive abilities and social responsiveness in children with 22q11.2 deletion syndrome (22q11DS) and children with idiopathic intellectual disability (IID). <i>Research in Developmental Disabilities</i> , 2018 , 81, 89-102	2.7	9
27	Attention deficit hyperactivity disorder symptoms as antecedents of later psychotic outcomes in 22q11.2 deletion syndrome. <i>Schizophrenia Research</i> , 2019 , 204, 320-325	3.6	11
26	Exploratory case study of monozygotic twins with 22q11.2DS provides further clues to circumscribe neurocognitive markers of psychotic symptoms. <i>NeuroImage: Clinical</i> , 2019 , 24, 101987	5.3	1
25	Genotype-phenotype associations in children with copy number variants associated with high neuropsychiatric risk in the UK (IMAGINE-ID): a case-control cohort study. <i>Lancet Psychiatry</i> , 2019 , 6, 493-505	23.3	41
24	Epilepsy and seizures in young people with 22q11.2 deletion syndrome: Prevalence and links with other neurodevelopmental disorders. <i>Epilepsia</i> , 2019 , 60, 818-829	6.4	18
23	The emergence of psychotic experiences in the early adolescence of 22q11.2 Deletion Syndrome. <i>Journal of Psychiatric Research</i> , 2019 , 109, 10-17	5.2	11
22	Co-creating a knowledge base in the "22q11.2 deletion syndrome" community. <i>Journal of Community Genetics</i> , 2020 , 11, 101-111	2.5	3
21	Sleep problems and associations with psychopathology and cognition in young people with 22q11.2 deletion syndrome (22q11.2DS). <i>Psychological Medicine</i> , 2020 , 50, 1191-1202	6.9	10
20	Pan-european landscape of research into neurodevelopmental copy number variants: A survey by the MINDDS consortium. <i>European Journal of Medical Genetics</i> , 2020 , 63, 104093	2.6	0
19	Consensus of the Italian Primary Immunodeficiency Network on transition management from pediatric to adult care in patients affected with childhood-onset inborn errors of immunity. <i>Journal of Allergy and Clinical Immunology</i> , 2020 , 146, 967-983	11.5	0
18	Using common genetic variation to examine phenotypic expression and risk prediction in 22q11.2 deletion syndrome. <i>Nature Medicine</i> , 2020 , 26, 1912-1918	50.5	35
17	Cognitive deficits in childhood, adolescence and adulthood in 22q11.2 deletion syndrome and association with psychopathology. <i>Translational Psychiatry</i> , 2020 , 10, 53	8.6	12
16	ENIGMA-DTI: Translating reproducible white matter deficits into personalized vulnerability metrics in cross-diagnostic psychiatric research. <i>Human Brain Mapping</i> , 2020 ,	5.9	23
15	22q11.2 Deletion Syndrome. 2021 , 163-194		
14	Neurodevelopmental Trajectories and Psychiatric Morbidity: Lessons Learned From the 22q11.2 Deletion Syndrome. <i>Current Psychiatry Reports</i> , 2021 , 23, 13	9.1	7

13	Longitudinal Psychiatric and Developmental Outcomes in 22q11.2 Deletion Syndrome: A Systematic Review. <i>Journal of Developmental and Behavioral Pediatrics</i> , 2021 , 42, 415-427	2.4	0
12	A normative chart for cognitive development in a genetically selected population. <i>Neuropsychopharmacology</i> , 2021 ,	8.7	4
11	Social cognition in 22q11.2 deletion syndrome and idiopathic developmental neuropsychiatric disorders. <i>Journal of Neurodevelopmental Disorders</i> , 2021 , 13, 15	4.6	1
10	Age-Related Improvements in Executive Functions and Focal Attention in 22q11.2 Deletion Syndrome Vary Across Domain and Task. <i>Journal of the International Neuropsychological Society</i> , 2021 , 1-14	3.1	1
9	Learning from atypical development: A systematic review of executive functioning in children and adolescents with the 22q11.2 deletion syndrome. <i>Developmental Review</i> , 2021 , 60, 100962	7.4	0
8	A cross-comparison of cognitive ability across 8 genomic disorders. <i>Current Opinion in Genetics and Development</i> , 2021 , 68, 106-116	4.9	0
7	Annual Research Review: Anterior Modifiers in the Emergence of Neurodevelopmental Disorders (AMEND)-a systems neuroscience approach to common developmental disorders. <i>Journal of Child Psychology and Psychiatry and Allied Disciplines</i> , 2021 , 62, 610-630	7.9	14
6	DRAGON-Data: A platform and protocol for integrating genomic and phenotypic data across large psychiatric cohorts.		0
5	Altered developmental trajectories of verbal learning skills in 22q11.2DS: associations with hippocampal development and psychosis. <i>Psychological Medicine</i> , 1-10	6.9	
4	Neurodevelopmental outcome, developmental trajectories, and management in 22q11.2 deletion syndrome. 2022 , 270-284		0
3	DRAGON-Data: a platform and protocol for integrating genomic and phenotypic data across large psychiatric cohorts. 2023 , 9,		0
2	Updated clinical practice recommendations for managing children with 22q11.2 deletion syndrome. 2023 , 25, 100338		0
1	Executive functioning in preschoolers with 22q11.2 deletion syndrome and the impact of congenital heart defects. 2023 , 15,		0