

# CITATION REPORT

List of articles citing

## Obesity in adults with 22q11.2 deletion syndrome

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Genetics in Medicine, 2017, 19, 204-208.

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#	Paper	IF	Citations
50	Update on the 22q11.2 deletion syndrome and its relevance to schizophrenia. <i>Current Opinion in Psychiatry</i> , <b>2017</b> , 30, 191-196	4.9	61
49	A systematic review of genetic syndromes with obesity. <i>Obesity Reviews</i> , <b>2017</b> , 18, 603-634	10.6	93
48	Genomic Disorders in Psychiatry-What Does the Clinician Need to Know?. <i>Current Psychiatry Reports</i> , <b>2017</b> , 19, 82	9.1	27
47	MYT1L mutations cause intellectual disability and variable obesity by dysregulating gene expression and development of the neuroendocrine hypothalamus. <i>PLoS Genetics</i> , <b>2017</b> , 13, e1006957	6	34
46	Congenital heart diseases and cardiovascular abnormalities in 22q11.2 deletion syndrome: From well-established knowledge to new frontiers. <i>American Journal of Medical Genetics, Part A</i> , <b>2018</b> , 176, 2087-2098	2.5	29
45	Recurrent reciprocal copy number variants: Roles and rules in neurodevelopmental disorders. <i>Developmental Neurobiology</i> , <b>2018</b> , 78, 519-530	3.2	24
44	Chromosomal microarray analysis in the genetic evaluation of 279 patients with syndromic obesity. <i>Molecular Cytogenetics</i> , <b>2018</b> , 11, 14	2	21
43	22q and two: 22q11.2 deletion syndrome and coexisting conditions. <i>American Journal of Medical Genetics, Part A</i> , <b>2018</b> , 176, 2203-2214	2.5	15
42	Copy Number Variation and Risk of Stroke. <i>Stroke</i> , <b>2018</b> , 49, 2549-2554	6.7	3
41	Neuropsychiatric expression and catatonia in 22q11.2 deletion syndrome: An overview and case series. <i>American Journal of Medical Genetics, Part A</i> , <b>2018</b> , 176, 2146-2159	2.5	13
40	Genome-wide copy number variation analysis identifies novel candidate loci associated with pediatric obesity. <i>European Journal of Human Genetics</i> , <b>2018</b> , 26, 1588-1596	5.3	15
39	Copy Number Variation of Immune-Related Genes and Their Association with Iodine in Adults with Autoimmune Thyroid Diseases. <i>International Journal of Endocrinology</i> , <b>2018</b> , 2018, 1705478	2.7	11
38	Phenome-wide Burden of Copy-Number Variation in the UK Biobank. <i>American Journal of Human Genetics</i> , <b>2019</b> , 105, 373-383	11	18
37	Adverse effects of antipsychotic medication in patients with 22q11.2 deletion syndrome: A systematic review. <i>American Journal of Medical Genetics, Part A</i> , <b>2019</b> , 179, 2292-2306	2.5	5
36	All-cause mortality and survival in adults with 22q11.2 deletion syndrome. <i>Genetics in Medicine</i> , <b>2019</b> , 21, 2328-2335	8.1	19
35	22q11.2 Deletion Syndrome-Associated Parkinson's Disease. <i>Movement Disorders Clinical Practice</i> , <b>2019</b> , 6, 11-16	2.2	12
34	A genetic model for multimorbidity in young adults. <i>Genetics in Medicine</i> , <b>2020</b> , 22, 132-141	8.1	9

33	The Role of Cell Growth-Related Gene Copy Number Variation in Autoimmune Thyroid Disease. <i>Biological Trace Element Research</i> , <b>2020</b> , 195, 409-416	4.5	4
32	Autoimmune Thyroid Disease in Specific Genetic Syndromes in Childhood and Adolescence. <i>Frontiers in Endocrinology</i> , <b>2020</b> , 11, 543	5.7	13
31	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> , <b>2020</b> , 146, 967-983	11.5	0
30	22q11.2 microdeletion and increased risk for type 2 diabetes. <i>EClinicalMedicine</i> , <b>2020</b> , 26, 100528	11.3	2
29	Complement Activation in 22q11.2 Deletion Syndrome. <i>Journal of Clinical Immunology</i> , <b>2020</b> , 40, 515-523	3.7	1
28	Analysis of genes within the schizophrenia-linked 22q11.2 deletion identifies interaction of night owl/LZTR1 and NF1 in GABAergic sleep control. <i>PLoS Genetics</i> , <b>2020</b> , 16, e1008727	6	7
27	Mechanisms Underlying the Comorbidity of Schizophrenia and Type 2 Diabetes Mellitus. <i>International Journal of Neuropsychopharmacology</i> , <b>2021</b> , 24, 367-382	5.8	5
26	Endocrinopathies et excès pondéral. <b>2021</b> , 375-379		
25	The copy number variation and stroke (CaNVAS) risk and outcome study. <i>PLoS ONE</i> , <b>2021</b> , 16, e0248791	3.7	
24	Gastrointestinal Manifestations of Immunodeficiency. <b>2021</b> , 429-450.e7		
23	Phenome-wide burden of copy number variation in UK Biobank.		3
22	Analysis of genes within the schizophrenia-linked 22q11.2 deletion identifies interaction of night owl/LZTR1 and NF1 in GABAergic sleep control.		
21	Integration of genetic, transcriptomic, and clinical data provides insight into 16p11.2 and 22q11.2 CNV genes.		
20	Integration of genetic, transcriptomic, and clinical data provides insight into 16p11.2 and 22q11.2 CNV genes. <i>Genome Medicine</i> , <b>2021</b> , 13, 172	14.4	1
19	DELETION 22q11.2 (VELO-CARDIO-FACIAL SYNDROME/DIGEORGE SYNDROME). <b>2021</b> , 291-316		
18	Metabolic effects of the schizophrenia-associated 3q29 deletion are sex-specific and uncoupled from behavioral phenotypes.		
17	An Update Evolving View of Copy Number Variations in Autoimmune Diseases.. <i>Frontiers in Genetics</i> , <b>2021</b> , 12, 794348	4.5	1
16	Metabolic effects of the schizophrenia-associated 3q29 deletion.. <i>Translational Psychiatry</i> , <b>2022</b> , 12, 66	8.6	0

15	A case of follow-up of a patient with 22q11.2 distal deletion syndrome and a review of the literature. <i>Journal of Genetic Medicine</i> , <b>2021</b> , 18, 110-116	0.2	
14	Hypertriglyceridemia in young adults with a 22q11.2 microdeletion.. <i>European Journal of Endocrinology</i> , <b>2022</b> , 187, 91-99	6.5	○
13	Gastrointestinal Features of 22Q11.2 Deletion Syndrome Include Chronic Motility Problems from Childhood to Adulthood. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , Publish Ahead of Print,	2.8	○
12	Congenital heart disease and cardiovascular abnormalities associated with 22q11.2 deletion syndrome. <b>2022</b> , 78-100		○
11	Gastroenterological manifestations associated with 22q11.2 deletion syndrome. <b>2022</b> , 182-209		○
10	Mental health in adults with 22q11.2 deletion syndrome. <b>2022</b> , 322-337		○
9	The impact of 22q11.2 copy number variants on human traits in the general population.		○
8	Combinations of genes at the 16p11.2 and 22q11.2 CNVs contribute to neurobehavioral traits.		○
7	Cardiac function in adolescents and young adults with 22q11.2 deletion syndrome without congenital heart disease. <b>2022</b> , 104651		○
6	Environmental Influences on the Relation between the 22q11.2 Deletion Syndrome and Mental Health: A Literature Review. <b>2022</b> , 13, 2003		○
5	Hypoparathyroidism and late-onset hypogonadism in an adult male with familial 22q11.2 deletion syndrome: a case report with 3-year follow-up and review of the literature. <b>2022</b> , 22,		○
4	Tonsillectomy in Children with 22q11.2 Deletion Syndrome. <b>2022</b> , 13, 2187		○
3	The impact of 22q11.2 copy-number variants on human traits in the general population. <b>2023</b> , 110, 300-313		○
2	Updated clinical practice recommendations for managing adults with 22q11.2 deletion syndrome. <b>2023</b> , 25, 100344		○
1	Adult-onset obstructive sleep apnea and pediatric pharyngoplasty in 22q11.2 deletion syndrome. <b>2023</b> , 104, 49-55		○