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Clinical features of 78 adults with 22q11 Deletion Syndrome

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#	Paper	IF	Citations
359	Velocardiofacial syndrome (chromosome 22q11.2 deletion syndrome) as a model of schizophrenia. 309-327		
358	The 22q11.2 deletion in children: high rate of autistic disorders and early onset of psychotic symptoms. 2006 , 45, 1104-1113		290
357	The role of the psychologist in adult congenital heart disease. 2006 , 24, 607-18, vi		38
356	Neurocognitive profile in 22q11 deletion syndrome and schizophrenia. 2006 , 87, 270-8		145
355	Altered expression of hippocampal dentate granule neuron genes in a mouse model of human 22q11 deletion syndrome. 2006 , 88, 251-9		20
354	The team approach to pregnancy and congenital heart disease. 2006 , 24, 587-605, vi		15
353	PCR screening for 22q11.2 microdeletion: development of a new cost-effective diagnostic tool. 2006 , 369, 78-81		6
352	Identification of DNA copy-number aberrations by array-comparative genomic hybridization in patients with schizophrenia. 2006 , 344, 531-9		42
351	Analyses of the associations between the genes of 22q11 deletion syndrome and schizophrenia. 2006 , 51, 1037-1045		44
350	Schizophrenia in an adult with 6p25 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2006 , 140, 1208-13	2.5	19
349	Metyrosine in psychosis associated with 22q11.2 deletion syndrome: case report. 2007 , 17, 115-20		8
348	Early developmental specification of the thyroid gland depends on hox-expressing surrounding tissue and on FGF signals. 2007 , 134, 2871-9		53
347	Ocular findings in the chromosome 22q11.2 deletion syndrome. 2007 , 11, 179-82		39
346	The 22q11 deletion syndrome candidate gene Tbx1 determines thyroid size and positioning. 2007 , 16, 276-85		60
345	Catechol-O-methyl transferase and expression of schizophrenia in 73 adults with 22q11 deletion syndrome. 2007 , 61, 1135-40		79
344	Fluorescence in situ hybridization (FISH) screening for the 22q11.2 deletion in patients with clinical features of velocardiofacial syndrome but without cardiac anomalies. 2007 , 30, 21-24		7
343	An inherited atypical 1 Mb 22q11.2 deletion within the DGS/VCFS 3 Mb region in a child with obesity and aggressive behavior. <i>American Journal of Medical Genetics, Part A</i> , 2007 , 143A, 1928-32	2.5	25

342	Adult dysmorphology: perspectives on approach to diagnosis and care. 2007 , 145C, 227-9		8
341	Natural history of aging in Cornelia de Lange syndrome. 2007 , 145C, 248-60		99
340	Deletion 22q11: spectrum of associated disorders. 2007 , 14, 136-9		37
339	Molecular characterization of deletion breakpoints in adults with 22q11 deletion syndrome. 2007 , 120, 837-45		43
338	The fate of children with microdeletion 22q11.2 syndrome and congenital heart defect: clinical course and cardiac outcome. 2008 , 29, 76-83		35
337	Advanced imaging of the cervical spine and spinal cord in 22q11.2 deletion syndrome: age-matched, double-cohort, controlled study. 2008 , 2, 333-41		13
336	Schizophrenia and 22q11.2 deletion syndrome. 2008 , 10, 148-57		173
335	Genomic structural variation and schizophrenia. 2008 , 10, 171-7		10
334	Candidate genes and the behavioral phenotype in 22q11.2 deletion syndrome. 2008 , 14, 26-34		58
333	Sclerocornea associated with the chromosome 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2008 , 146A, 904-9	2.5	28
332	22q11 deletion syndrome: is that what they used to call . . . ?. 2008 , 44, 259-66		2
331	Adults with genetic syndromes and cardiovascular abnormalities: clinical history and management. 2008 , 10, 469-94		106
330	[Review of 22 patients with 22q11.2 deletion syndrome: phenotype spectrum]. 2008 , 69, 304-10		2
329	Chromosomes, Genes, and the Thyroid Gland. 2008 , 55-71		
328	Extracardiac features predicting 22q11.2 deletion syndrome in adult congenital heart disease. 2008 , 131, 51-8		42
327	Recognizing a common genetic syndrome: 22q11.2 deletion syndrome. 2008 , 178, 391-3		18
326	Understanding genetic factors in idiopathic scoliosis, a complex disease of childhood. 2008 , 9, 51-9		58
325	Copy number variations and risk for schizophrenia in 22q11.2 deletion syndrome. 2008 , 17, 4045-53		132

324	miRNA and Schizophrenia. 2008 , 267-281		
323	The Case: Hypocalcemia, chronic renal failure and dysmorphism. 2008 , 74, 1495-6		
322	The Value of a Genetic Diagnosis for Individuals with Intellectual Disabilities: Optimising Healthcare and Function Across the Lifespan. 2008 , 54, 69-82		13
321	[22q11.2 deficiency syndrome diagnosed with congestive heart failure]. 2008 , 97, 2785-7		1
320	Síndrome de deleçã 22q11.2: compreendendo o CATCH22. 2009 , 27, 211-220		7
319	Premature death in adults with 22q11.2 deletion syndrome. 2009 , 46, 324-30		66
318	Neuropathologic features in adults with 22q11.2 deletion syndrome. 2009 , 19, 153-64		62
317	Diminished dosage of 22q11 genes disrupts neurogenesis and cortical development in a mouse model of 22q11 deletion/DiGeorge syndrome. 2009 , 106, 16434-45		126
316	Affective disorders and other psychiatric diagnoses in children and adolescents with 22q11.2 Deletion Syndrome. 2009 , 119, 177-80		35
315	The co-occurrence of early onset Parkinson disease and 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2009 , 149A, 525-8	2.5	39
314	Genetic basis in epilepsies caused by malformations of cortical development and in those with structurally normal brain. 2009 , 126, 173-93		47
313	Thymus transplantation in complete DiGeorge anomaly. 2009 , 44, 61-70		48
312	Psychosis in children with velocardiofacial syndrome (22q11.2 deletion syndrome). 2009 , 11, 99-105		26
311	Microdileçã 22q11.2 ríve par une hypocalcémie nã natale avec une dysmorphie faciale mineure. 2009 , 2009, 83-85		
310	De novo copy number variants identify new genes and loci in isolated sporadic tetralogy of Fallot. 2009 , 41, 931-5		325
309	Modeling cognitive endophenotypes of schizophrenia in mice. 2009 , 32, 347-58		118
308	Thyroid gland development and function in the zebrafish model. 2009 , 312, 14-23		144
307	Tetralogy of Fallot. 2009 , 374, 1462-71		314

306	Spinal deformity in patients with DiGeorge syndrome. 2010 , 23, 208-14		6
305	Deletion 22q11.2 (Velo-Cardio-Facial Syndrome/Digeorge Syndrome). 2010 , 263-284		5
304	Evidence of gray matter reduction and dysfunction in chromosome 22q11.2 deletion syndrome. 2010 , 181, 1-8		37
303	Cardiovascular anomalies associated with chromosome 22q11.2 deletion syndrome. 2010 , 105, 1617-24		178
302	Thymus transplantation. 2010 , 135, 236-46		102
301	Secondary immunologic consequences in chromosome 22q11.2 deletion syndrome (DiGeorge syndrome/velocardiofacial syndrome). 2010 , 136, 409-18		61
300	The morphology of the sella turcica in velocardiofacial syndrome suggests involvement of a neural crest developmental field. <i>American Journal of Medical Genetics, Part A</i> , 2010 , 152A, 1450-7	2.5	10
299	Elements of psychocardiology in the psychosocial handling of adults with congenital heart disease. 2010 , 1, 34		7
298	New Genetic Findings in Schizophrenia: Is there Still Room for the Dopamine Hypothesis of Schizophrenia?. 2010 , 4, 23		18
297	De novo rates and selection of large copy number variation. 2010 , 20, 1469-81		221
296	Striatal D ₂ receptor binding in 22q11 deletion syndrome: an [¹¹ C]IBZM SPECT study. 2010 , 24, 1525-31		20
295	Penetrance for copy number variants associated with schizophrenia. 2010 , 19, 3477-81		117
294	ESC Guidelines for the management of grown-up congenital heart disease (new version 2010). 2010 , 31, 2915-57		1708
293	Review of pathological hallmarks of schizophrenia: comparison of genetic models with patients and nongenetic models. 2010 , 36, 301-13		111
292	Cross-disorder genomewide analysis of schizophrenia, bipolar disorder, and depression. 2010 , 167, 1254-63		157
291	Bridging the gene-behavior divide through neuroimaging deletion syndromes: Velocardiofacial (22q11.2 Deletion) and Williams (7q11.23 Deletion) syndromes. 2010 , 53, 857-69		15
290	Genetics of Schizophrenia and Bipolar Affective Disorder. 2010 , 759-776		
289	Psychiatric education in the genomic era. 2010 , 34, 87-9		6

288	Genetics of Congenital Heart Defects. 2011 , 283-304		
287	Patterns of cardiac and extracardiac anomalies in adults with tetralogy of fallot. 2011 , 161, 131-7		25
286	Síndrome de deleço 22q11.2 e cardiopatias congnitas. 2011 , 29, 251-260		5
285	Practical guidelines for managing patients with 22q11.2 deletion syndrome. 2011 , 159, 332-9.e1		381
284	Pain in adults with intellectual disabilities. 2011 , 152, 971-974		25
283	Complex congenital heart disease in unaffected relatives of adults with 22q11.2 deletion syndrome. 2011 , 107, 466-71		19
282	A prospective study of influenza vaccination and a comparison of immunologic parameters in children and adults with chromosome 22q11.2 deletion syndrome (digeorge syndrome/velocardiofacial syndrome). 2011 , 31, 927-35		29
281	Cognitive, behavioural and psychiatric phenotype in 22q11.2 deletion syndrome. 2011 , 41, 403-12		97
280	How might stress contribute to increased risk for schizophrenia in children with chromosome 22q11.2 deletion syndrome?. 2011 , 3, 68-75		36
279	Copy number variants: a new molecular frontier in clinical psychiatry. 2011 , 13, 129-37		9
278	High frequency of known copy number abnormalities and maternal duplication 15q11-q13 in patients with combined schizophrenia and epilepsy. 2011 , 12, 154		48
277	A comprehensive network and pathway analysis of candidate genes in major depressive disorder. 2011 , 5 Suppl 3, S12		79
276	Chromosome 22q11.2 deletion syndrome in African-American patients: a diagnostic challenge. <i>American Journal of Medical Genetics, Part A</i> , 2011 , 155A, 2186-95	2.5	12
275	Association of schizophrenia in 22q11.2 deletion syndrome and gray matter volumetric deficits in the superior temporal gyrus. 2011 , 168, 522-9		45
274	Hypoparathyroidism and autoimmunity in the 22q11.2 deletion syndrome. 2011 , 165, 345-52		39
273	Behavioral and molecular biomarkers in translational animal models for neuropsychiatric disorders. 2011 , 101, 203-38		25
272	Is there a core neuropsychiatric phenotype in 22q11.2 deletion syndrome?. 2012 , 25, 131-7		45
271	Startle reactivity and prepulse inhibition of the acoustic startle response are modulated by catechol-O-methyl-transferase Val(158) Met polymorphism in adults with 22q11 deletion syndrome. 2012 , 26, 1548-60		12

270	Dysregulation of DGCR6 and DGCR6L: psychopathological outcomes in chromosome 22q11.2 deletion syndrome. 2012 , 2, e105		16
269	Age-dependent microRNA control of synaptic plasticity in 22q11 deletion syndrome and schizophrenia. 2012 , 32, 14132-44		84
268	Multiplex ligation-dependent probe amplification (MLPA): a reliable alternative for fetal chromosome analysis?. 2012 , 25, 1383-6		5
267	Antibody deficiency in adults with 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 1934-40	2.5	19
266	Speech and hearing in adults with 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 3071-9	2.5	27
265	22q11.2 deletion syndrome: attitudes towards disclosing the risk of psychiatric illness. 2012 , 21, 825-34		20
264	Palate abnormalities in Chilean patients with chromosome 22q11 microdeletion syndrome. 2012 , 76, 1726-8		8
263	Molecular karyotyping as a relevant diagnostic tool in children with growth retardation with Silver-Russell features. 2012 , 161, 933-42		32
262	A 4.5 Mb terminal deletion of chromosome 12p helps further define a psychosis-associated locus. <i>European Journal of Medical Genetics</i> , 2012 , 55, 573-6	2.6	8
261	Recomendações da ESC para o tratamento da cardiopatia congênita no adulto (nova versão de 2010). 2012 , 31, 541.e1-541.e53		
260	Schizophrenia genetics: progress, at last. 2012 , 22, 238-44		40
259	Chromosome 22 microdeletion in children with syndromic congenital heart disease by fluorescent in situ hybridization (FISH). 2012 , 13, 313-322		
258	Reduced fractional anisotropy and axial diffusivity in white matter in 22q11.2 deletion syndrome: a pilot study. 2012 , 141, 35-9		26
257	Functional outcomes of adults with 22q11.2 deletion syndrome. 2012 , 14, 836-43		63
256	Clinical applications of schizophrenia genetics: genetic diagnosis, risk, and counseling in the molecular era. 2012 , 5, 1-18		29
255	The genetics of schizophrenia. 230-261		1
254	Aging in Prader-Willi syndrome: twelve persons over the age of 50 years. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 1326-36	2.5	24
253	Syndrome-specific growth charts for 22q11.2 deletion syndrome in Caucasian children. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 2665-71	2.5	29

252	Occurrence of affective disorders compared to other psychiatric disorders in children and adolescents with 22q11.2 deletion syndrome. 2012 , 136, 222-8		33
251	Caregiver and adult patient perspectives on the importance of a diagnosis of 22q11.2 deletion syndrome. 2012 , 56, 641-51		38
250	Congenital heart defects in a novel recurrent 22q11.2 deletion harboring the genes CRKL and MAPK1. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 574-80	2.5	31
249	Computerized neurocognitive profile in young people with 22q11.2 deletion syndrome compared to youths with schizophrenia and at-risk for psychosis. 2012 , 159B, 87-93		31
248	Head and neck manifestations of 22q11.2 deletion syndromes. 2012 , 269, 381-7		20
247	Velopharyngoplasty in patients with 22q11.2 microdeletion syndrome: outcomes following the Newcastle protocol. 2013 , 36, 607-618		1
246	Genetic contributions to changes of fiber tracts of ventral visual stream in 22q11.2 deletion syndrome. 2013 , 7, 316-25		20
245	Premorbid adjustment and schizophrenia in individuals with 22q11.2 deletion syndrome. 2013 , 151, 221-5		19
244	Genetics of psychiatric disorders in the GWAS era: an update on schizophrenia. 2013 , 263 Suppl 2, S147-54		38
243	Human-specific endogenous retroviral insert serves as an enhancer for the schizophrenia-linked gene PRODH. 2013 , 110, 19472-7		55
242	Radial microcolumnar cortical architecture: maturational arrest or cortical dysplasia?. 2013 , 48, 259-70		35
241	Discussing the psychiatric manifestations of 22q11.2 deletion syndrome: an exploration of clinical practice among medical geneticists. 2013 , 15, 713-20		12
240	Heterogeneity of schizophrenia: Genetic and symptomatic factors. 2013 , 162B, 648-52		30
239	Diffuse malformations of cortical development. 2013 , 111, 653-65		19
238	The role of modern imaging techniques in the diagnosis of malposition of the branch pulmonary arteries and possible association with microdeletion 22q11.2. 2013 , 23, 181-8		12
237	The diverse clinical features of chromosome 22q11.2 deletion syndrome (DiGeorge syndrome). 2013 , 1, 589-94		35
236	Génétique des schizophrénies : mise en perspective des schizophrénies à début précoce et autres pathologies du développement. 2013 , 61, 317-325		
235	Prenatal diagnosis of 22q11.2 deletion syndrome in twin pregnancy: a case report. 2013 , 41 Suppl 1, 6-9		3

234	Chromosome 22q11.2 deletion syndrome presenting as adult onset hypoparathyroidism: clues to diagnosis from dysmorphic facial features. 2013 , 2013, 802793		3
233	Association between early-onset Parkinson disease and 22q11.2 deletion syndrome: identification of a novel genetic form of Parkinson disease and its clinical implications. 2013 , 70, 1359-66		107
232	Hippocampal malrotation is associated with chromosome 22q11.2 microdeletion. 2013 , 40, 652-6		29
231	The value of the clinical geneticist caring for adults with congenital heart disease: diagnostic yield and patients' perspective. <i>American Journal of Medical Genetics, Part A</i> , 2013 , 161A, 1628-37	2.5	14
230	Principal components of heritability from neurocognitive domains differ between families with schizophrenia and control subjects. 2013 , 39, 464-71		6
229	Genetics of Schizophrenia. 2013 , 42, 5-22		3
228	Estudos traducionais de neuropsiquiatria e esquizofrenia: modelos animais geneticos e de neurodesenvolvimento. 2013 , 40, 41-50		4
227	Characteristic face: a key indicator for direct diagnosis of 22q11.2 deletions in Chinese velocardiofacial syndrome patients. 2013 , 8, e54404		23
226	22q11 deletion syndrome: a review of the neuropsychiatric features and their neurobiological basis. 2013 , 9, 1873-84		26
225	A 32-year-old Woman Diagnosed with 22q11.2 Deletion Syndrome and Complicated by Hypothyroidism. 2014 , 15, 72-75		2
224	Applicability of the nonverbal learning disability paradigm for children with 22q11.2 deletion syndrome. 2014 , 47, 153-66		12
223	Neonatal hypocalcemia, neonatal seizures, and intellectual disability in 22q11.2 deletion syndrome. 2014 , 16, 40-4		57
222	Defects in Thymic Development. 2014 , 221-242		1
221	Perceived burden and neuropsychiatric morbidities in adults with 22q11.2 deletion syndrome. 2014 , 58, 198-210		15
220	Prevalence of hypocalcaemia and its associated features in 22q11.2 deletion syndrome. 2014 , 81, 190-6		49
219	MicroRNAs in Alcohol Abuse and Toxicity. 2014 , 497-521		1
218	Investigation of selected genomic deletions and duplications in a cohort of 338 patients presenting with syndromic obesity by multiplex ligation-dependent probe amplification using synthetic probes. 2014 , 7, 75		7
217	Under-recognition of 22q11.2 deletion in adult Chinese patients with conotruncal anomalies: implications in transitional care. <i>European Journal of Medical Genetics</i> , 2014 , 57, 306-11	2.6	28

216	The 22q11.2 deletion syndrome as a window into complex neuropsychiatric disorders over the lifespan. 2014 , 75, 351-60		121
215	Intergenerational and intrafamilial phenotypic variability in 22q11.2 deletion syndrome subjects. 2014 , 15, 1		28
214	Towards a safety net for management of 22q11.2 deletion syndrome: guidelines for our times. 2014 , 173, 757-65		46
213	A French collaborative survey of 272 fetuses with 22q11.2 deletion: ultrasound findings, fetal autopsies and pregnancy outcomes. 2014 , 34, 424-30		38
212	Evidence that duplications of 22q11.2 protect against schizophrenia. 2014 , 19, 37-40		130
211	Copy number variants in short children born small for gestational age. 2014 , 82, 310-8		17
210	The neural crest: a versatile organ system. 2014 , 102, 275-98		40
209	Cross-Disorder Comparison of Four Neuropsychiatric CNV Loci. 2014 , 2, 151-161		21
208	Further evidence for high rates of schizophrenia in 22q11.2 deletion syndrome. 2014 , 153, 231-6		62
207	Presenting symptoms in adults with the 22q11 deletion syndrome. <i>European Journal of Medical Genetics</i> , 2014 , 57, 157-62	2.6	16
206	Psychopathology and cognition in children with 22q11.2 deletion syndrome. 2014 , 204, 46-54		53
205	A longitudinal study of hearing and middle ear status of individuals with cleft palate with and without additional malformations/syndromes. 2014 , 51, e94-e101		12
204	22q11.2 deletion syndrome. 2015 , 1, 15071		492
203	The Epidemiology of Nonsurgical Hypoparathyroidism in Denmark: A Nationwide Case Finding Study. 2015 , 30, 1738-44		138
202	Case history and genome-wide scans for copy number variants in a family with patient having 15q11.1-q11.2 duplication and 22q11.2 deletion, and schizophrenia. 2015 , 168B, 229-35		
201	Co-occurrence of a de novo Williams and 22q11.2 microdeletion syndromes. <i>American Journal of Medical Genetics, Part A</i> , 2015 , 167A, 1927-31	2.5	3
200	A New Method for Detecting Associations with Rare Copy-Number Variants. 2015 , 11, e1005403		10
199	The First Case Report in Italy of Di George Syndrome Detected by Noninvasive Prenatal Testing. 2015 , 2015, 813104		2

198	Characteristics of 22q 11.2 deletion syndrome undiagnosed until adulthood: an example suggesting the importance of psychiatric manifestations. <i>BMJ Case Reports</i> , 2015 , 2015,	0.9	6
197	Salicylic Acid and Drought Stress Response: Biochemical to Molecular Crosstalk. 2015 , 247-265		4
196	MicroRNA and Posttranscriptional Dysregulation in Psychiatry. 2015 , 78, 231-9		116
195	Morphological changes in gray matter volume correlate with catechol-O-methyl transferase gene Val158Met polymorphism in first-episode treatment-naïve patients with schizophrenia. 2015 , 31, 31-42		6
194	Chromosomal Imbalances in Patients with Congenital Cardiac Defects: A Meta-analysis Reveals Novel Potential Critical Regions Involved in Heart Development. 2015 , 10, 193-208		18
193	Reproductive Health Issues for Adults with a Common Genomic Disorder: 22q11.2 Deletion Syndrome. 2015 , 24, 810-21		18
192	Practical guidelines for managing adults with 22q11.2 deletion syndrome. 2015 , 17, 599-609		154
191	Craniofacial dysmorphology in 22q11.2 deletion syndrome by 3D laser surface imaging and geometric morphometrics: illuminating the developmental relationship to risk for psychosis. <i>American Journal of Medical Genetics, Part A</i> , 2015 , 167A, 529-36	2.5	16
190	Clinical and molecular cytogenetic studies of an unrecognised 22q11.2 deletion in three families. <i>Experimental and Therapeutic Medicine</i> , 2015 , 9, 823-828	2.1	
189	Gene expression analysis in blood of ultra-high risk subjects compared to first-episode of psychosis patients and controls. 2015 , 16, 441-446		13
188	Comparative mapping of the 22q11.2 deletion region and the potential of simple model organisms. 2015 , 7, 18		61
187	22q11.2 deletion status and disease burden in children and adolescents with tetralogy of Fallot. 2015 , 8, 74-81		37
186	Modeling a model: Mouse genetics, 22q11.2 Deletion Syndrome, and disorders of cortical circuit development. <i>Progress in Neurobiology</i> , 2015 , 130, 1-28	10.9	58
185	DiGeorge Syndrome. 2015 , 368-374		
184	Response to clozapine in a clinically identifiable subtype of schizophrenia. 2015 , 206, 484-91		44
183	Aberrant Cortical Morphometry in the 22q11.2 Deletion Syndrome. 2015 , 78, 135-43		53
182	[22q11.2DS Syndrome as a Genetic Subtype of Schizophrenia]. 2015 , 44, 50-60		1
181	The schizophrenia/bipolar disorder candidate gene GNB1L is regulated in human temporal cortex by a cis-acting element located within the 3Rregion. 2015 , 31, 43-52		2

180	Application of high resolution SNP arrays in patients with congenital oral clefts in south China. 2016 , 95, 801-809		3
179	The importance of copy number variation in congenital heart disease. 2016 , 1, 16031		42
178	Two patients with chromosome 22q11.2 deletion presenting with childhood obesity and hyperphagia. <i>European Journal of Medical Genetics</i> , 2016 , 59, 401-3	2.6	5
177	Neural correlates of reward processing in adults with 22q11 deletion syndrome. 2016 , 8, 25		11
176	Cerebral cortex expansion and folding: what have we learned?. 2016 , 35, 1021-44		168
175	Current controversies in prenatal diagnosis 1: should NIPT routinely include microdeletions/microduplications?. 2016 , 36, 10-4		23
174	Performance on a computerized neurocognitive battery in 22q11.2 deletion syndrome: A comparison between US and Israeli cohorts. 2016 , 106, 33-41		14
173	Atypical functional connectivity in resting-state networks of individuals with 22q11.2 deletion syndrome: associations with neurocognitive and psychiatric functioning. 2016 , 8, 2		19
172	Communication of Psychiatric Risk in 22q11.2 Deletion Syndrome: A Pilot Project. 2016 , 25, 6-17		8
171	Schizophrenia Spectrum Disorders in a Danish 22q11.2 Deletion Syndrome Cohort Compared to the Total Danish Population--A Nationwide Register Study. 2016 , 42, 824-31		12
170	Rare loss-of-function variants in SETD1A are associated with schizophrenia and developmental disorders. 2016 , 19, 571-7		284
169	Fetal growth and gestational factors as predictors of schizophrenia in 22q11.2 deletion syndrome. 2016 , 18, 350-5		22
168	A French multicenter study of over 700 patients with 22q11 deletions diagnosed using FISH or aCGH. 2016 , 24, 844-51		28
167	Post-childhood Presentation and Diagnosis of DiGeorge Syndrome. 2016 , 55, 368-73		6
166	The Effectiveness and Safety of Antipsychotic and Antidepressant Medications in Individuals with 22q11.2 Deletion Syndrome. 2017 , 27, 83-90		20
165	Dental treatment of a patient with Opitz G/BBB syndrome. 2017 , 37, 102-106		
164	Genetic Drivers of Kidney Defects in the DiGeorge Syndrome. 2017 , 376, 742-754		83
163	Autism Spectrum and psychosis risk in the 22q11.2 deletion syndrome. Findings from a prospective longitudinal study. 2017 , 188, 59-62		35

162	Signs of dysarthria in adults with 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2017 , 173, 618-626	2.5	7
161	Growth characteristics and endocrine abnormalities in 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2017 , 173, 1301-1308	2.5	8
160	High prevalence of fatigue in adults with a 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2017 , 173, 858-867	2.5	9
159	Enlarged Cavum Septi Pellucidi and Vergae in the Fetus: A Cause for Concern. 2017 , 36, 1657-1668		10
158	Hyposmia, symptoms of rapid eye movement sleep behavior disorder, and parkinsonian motor signs suggest prodromal neurodegeneration in 22q11 deletion syndrome. 2017 , 28, 677-681		7
157	A CNV Catalogue. 2017 , 235-417		0
156	Prevalence of hearing loss and clinical otologic manifestations in patients with 22q11.2 deletion syndrome: A literature review. 2017 , 42, 1319-1328		12
155	Otologic and audiologic findings in 22q11.2 deletion syndrome. 2017 , 274, 765-771		10
154	Mitochondria in complex psychiatric disorders: Lessons from mouse models of 22q11.2 deletion syndrome: Hemizygous deletion of several mitochondrial genes in the 22q11.2 genomic region can lead to symptoms associated with neuropsychiatric disease. 2017 , 39, 1600177		19
153	A neurogenetic model for the study of schizophrenia spectrum disorders: the International 22q11.2 Deletion Syndrome Brain Behavior Consortium. 2017 , 22, 1664-1672		48
152	Machine-learning classification of 22q11.2 deletion syndrome: A diffusion tensor imaging study. 2017 , 15, 832-842		21
151	Obesity in adults with 22q11.2 deletion syndrome. 2017 , 19, 204-208		37
150	Neuropsychiatric aspects of 22q11.2 deletion syndrome: considerations in the prenatal setting. 2017 , 37, 61-69		9
149	Thalamic miR-338-3p mediates auditory thalamocortical disruption and its late onset in models of 22q11.2 microdeletion. 2017 , 23, 39-48		35
148	Treatment of 22q11.2 deletion syndrome-associated schizophrenia with comorbid anxiety and panic disorder. 2017 , 9, 7225		3
147	A surgical approach to the craniofacial defects of Opitz G/BBB syndrome. 2017 , 2017, rjx032		1
146	A case of schizophrenia comorbid for tetralogy of Fallot treated with clozapine: further considerations on a role for 22q.11.2 in the proneness for seizures. 2017 , 13, 2271-2273		1
145	Small RNA Dysregulation in Neurocognitive and Neuropsychiatric Disorders. 2017 , 225-245		1

144	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> , 2018 , 176, 2087-2098	2.5	29
143	Elucidating the diagnostic odyssey of 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2018 , 176, 936-944	2.5	26
142	Assessing Gene Expression in Treatment-Resistant Schizophrenia. 2018 , 55, 7000-7008		17
141	Variable immune deficiency related to deletion size in chromosome 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2018 , 176, 2082-2086	2.5	30
140	Chromosomal microarray analysis in the genetic evaluation of 279 patients with syndromic obesity. 2018 , 11, 14		21
139	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
138	Striatal dopamine release and impaired reinforcement learning in adults with 22q11.2 deletion syndrome. 2018 , 28, 732-742		6
137	Orthopaedic manifestations within the 22q11.2 Deletion syndrome: A systematic review. <i>American Journal of Medical Genetics, Part A</i> , 2018 , 176, 2104-2120	2.5	11
136	Higher adaptive functioning and lower rate of psychotic comorbidity in married versus unmarried individuals with 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2018 , 176, 2365-2374	2.5	2
135	Dental management of a patient with 22q11.2 deletion syndrome (22q11.2DS). <i>BMJ Case Reports</i> , 2018 , 2018,	0.9	1
134	Clozapine-induced myocarditis in an adolescent male with DiGeorge Syndrome. 2018 , 8, 313-316		4
133	Childhood Executive Functioning Predicts Young Adult Outcomes in 22q11.2 Deletion Syndrome. 2018 , 24, 905-916		7
132	unbalanced translocation t(15;22)(q26.2;q12) with velo-cardio-facial syndrome: A case report and review of the literature. <i>Experimental and Therapeutic Medicine</i> , 2018 , 16, 3589-3595	2.1	5
131	Otolaryngological features in a cohort of patients affected with 22q11.2 deletion syndrome: A monocentric survey. <i>American Journal of Medical Genetics, Part A</i> , 2018 , 176, 2128-2134	2.5	6
130	Understanding the pediatric psychiatric phenotype of 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2018 , 176, 2182-2191	2.5	36
129	Neuropsychiatric expression and catatonia in 22q11.2 deletion syndrome: An overview and case series. <i>American Journal of Medical Genetics, Part A</i> , 2018 , 176, 2146-2159	2.5	13
128	A mutation in transcription factor MAFB causes Focal Segmental Glomerulosclerosis with Duane Retraction Syndrome. 2018 , 94, 396-407		33
127	Genome-wide copy number variation analysis identifies novel candidate loci associated with pediatric obesity. 2018 , 26, 1588-1596		15

126	22q11.2 Deletion Syndrome. 2018 , 621-626.e1	3
125	The Role of Clinical Psychology and Peer to Peer Support in the Management of Chronic Medical Conditions - A Practical Example With Adults With Congenital Heart Disease. 2018 , 9, 731	12
124	Application of chromosomal microarray to investigate genetic causes of isolated fetal growth restriction. 2018 , 11, 33	8
123	Typical features of Parkinson disease and diagnostic challenges with microdeletion 22q11.2. 2018 , 90, e2059-e2067	25
122	Attention deficit hyperactivity disorder symptoms as antecedents of later psychotic outcomes in 22q11.2 deletion syndrome. 2019 , 204, 320-325	11
121	2018 AHA/ACC Guideline for the Management of Adults With Congenital Heart Disease: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. 2019 , 139, e698-e800	248
120	Neurocognition and adaptive functioning in a genetic high risk model of schizophrenia. 2019 , 49, 1047-1054	7
119	2018 AHA/ACC Guideline for the Management of Adults With Congenital Heart Disease: Executive Summary: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. 2019 , 73, 1494-1563	203
118	2018 AHA/ACC Guideline for the Management of Adults With Congenital Heart Disease: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. 2019 , 73, e81-e192	279
117	An Overview of Animal Models Related to Schizophrenia. 2019 , 64, 5-17	69
116	Genome-wide association study identifies 14 previously unreported susceptibility loci for adolescent idiopathic scoliosis in Japanese. 2019 , 10, 3685	22
115	Neurocognitive profile and onset of psychosis symptoms in children, adolescents and young adults with 22q11 deletion syndrome: A longitudinal study. 2019 , 208, 76-81	4
114	Long contiguous stretches of homozygosity detected by chromosomal microarrays (CMA) in patients with neurodevelopmental disorders in the South of Brazil. 2019 , 12, 50	7
113	Schizophrenia is a later-onset feature of PCDH19 Girls Clustering Epilepsy. 2019 , 60, 429-440	11
112	Lower cortisol levels and attenuated cortisol reactivity to daily-life stressors in adults with 22q11.2 deletion syndrome. 2019 , 106, 85-94	9
111	All-cause mortality and survival in adults with 22q11.2 deletion syndrome. 2019 , 21, 2328-2335	19
110	Ontogeny of the facial phenotypic variability in Mexican patients with 22q11.2 deletion syndrome. 2019 , 15, 29	0
109	2018 AHA/ACC Guideline for the Management of Adults With Congenital Heart Disease: Executive Summary: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. 2019 , 139, e637-e697	142

108	Chromosome 22q11.2 deletion syndrome and DiGeorge syndrome. 2019 , 287, 186-201		47
107	Hypoparathyroidism and Other Causes of Hypocalcemia. 2019 , 192-203		
106	Downregulation of genes outside the deleted region in individuals with 22q11.2 deletion syndrome. 2019 , 138, 93-103		5
105	22q11.2 Deletion Syndrome-Associated Parkinson Disease. 2019 , 6, 11-16		12
104	22q11 microdeletion syndrome and ultra-high risk for psychosis: The role of neurological soft signs as an independent marker of vulnerability for psychosis. 2019 , 13, 1191-1198		
103	Scoliosis in association with the 22q11.2 deletion syndrome: an observational study. 2019 , 104, 19-24		11
102	Pediatric endocrinology through syndromes. <i>European Journal of Medical Genetics</i> , 2020 , 63, 103614	2.6	5
101	Lower [¹⁸ F]fallypride binding to dopamine D receptors in frontal brain areas in adults with 22q11.2 deletion syndrome: a positron emission tomography study. 2020 , 50, 799-807		1
100	A genetic model for multimorbidity in young adults. 2020 , 22, 132-141		9
99	TBX1 is required for normal stria vascularis and semicircular canal development. 2020 , 457, 91-103		4
98	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
97	22q11.2 deletion syndrome and schizophrenia. 2020 , 52, 1181-1190		5
96	Clinical application of chromosomal microarray analysis for fetuses with craniofacial malformations. 2020 , 13, 38		3
95	Cognitive behavioral therapy in 22q11.2 deletion syndrome: A case study of two young adults with an anxiety disorder. 2021 , 25, 695-704		1
94	Early language measures associated with later psychosis features in 22q11.2 deletion syndrome. 2020 , 183, 392-400		3
93	Autoimmune Thyroid Disease in Specific Genetic Syndromes in Childhood and Adolescence. 2020 , 11, 543		13
92	Age-Related Parkinsonian Signs in Microdeletion 22q11.2. 2020 , 35, 1239-1245		2
91	Treatment of Epilepsy Associated with Common Chromosomal Developmental Diseases. 2020 , 15, 21-29		

90	Vitamin D status and the immune assessment in 22q11.2 deletion syndrome. 2020 , 200, 272-286	5
89	Access to Multidisciplinary Care for Patients With 22q11.2 Deletion Syndrome: Identifying Breakdowns in the Screening Process. 2020 , 31, 428-431	1
88	Complex Reoperation in a Patient With DiGeorge Syndrome. 2020 , 34, 1655-1662	
87	Views of adults with 22q11 deletion syndrome on reproductive choices. <i>American Journal of Medical Genetics, Part A</i> , 2020 , 182, 1284-1287	2.5 4
86	The Genetics and Epigenetics of 22q11.2 Deletion Syndrome. 2019 , 10, 1365	27
85	Defects in thymic development. 2020 , 357-379	1
84	Comprehensive analysis of a novel mouse model of the 22q11.2 deletion syndrome: a model with the most common 3.0-Mb deletion at the human 22q11.2 locus. 2020 , 10, 35	14
83	Meta-analysis of olfactory dysfunction in 22q11.2 deletion syndrome. 2020 , 285, 112783	1
82	The role of 22q11.2 deletion syndrome in the relationship between congenital heart disease and scoliosis. 2020 , 20, 956-963	3
81	Inborn errors of thymic stromal cell development and function. 2021 , 43, 85-100	5
80	2020 ESC Guidelines for the management of adult congenital heart disease. 2021 , 42, 563-645	290
79	Genomic Variation, Evolvability, and the Paradox of Mental Illness. 2020 , 11, 593233	0
78	Social cognition in 22q11.2 deletion syndrome and idiopathic developmental neuropsychiatric disorders. 2021 , 13, 15	1
77	Abnormal spirometry in adults with 22q11.2 microdeletion and congenital heart disease. 2021 , 3, 100085	0
76	Late diagnosed DiGeorge syndrome in a 44-year-old female: a rare cause for recurrent syncope in adulthood-a case report. 2021 , 5, ytab166	0
75	Care of Children with DiGeorge Before and After Cultured Thymus Tissue Implantation. 2021 , 41, 896-905	0
74	Guía ESC 2020 para el tratamiento de las cardiopatías congénitas del adulto. 2021 , 74, 436.e1-436.e79	3
73	ÖFRENİN GENETİK TEMELİNİN TARİHTEKNIK ÖLÇÜME ANLILMASI.	

72	Sexual knowledge and behaviour in 22q11.2 deletion syndrome, a complex care condition. 2021 ,		0
71	Clinical Features in a Large Cohort of Patients With 22q11.2 Deletion Syndrome. 2021 , 238, 215-220.e5		2
70	Evaluation and Maintenance of Behavioral Interventions for 22q11.2 Deletion Syndrome. 2021 , 1-8		
69	A Rare Etiology of Hypocalcemic Seizures in Adulthood: Clues to Diagnosis from Facial Dysmorphism. 2021 , 69, 161-163		0
68	22q11.2 Deletion Syndrome. 2021 , 154-163		0
67	Neuroradiographic findings in 22q11.2 deletion syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2017 , 173, 2158-2165	2.5	14
66	Current Understanding of Genetic Factors in Idiopathic Scoliosis. 2010 , 167-190		3
65	Hypocalcemia. 2014 , 265-278		3
64	22q11.2 microdeletion syndrome is a common cause of renal tract malformations. 2008 , 4, E1		6
63	Speech-Language Disorders in 22q11.2 Deletion Syndrome: Best Practices for Diagnosis and Management. 2019 , 28, 984-999		21
62	Association between catechol-O-methyltransferase Val108/158Met genotype and prefrontal hemodynamic response in schizophrenia. 2009 , 4, e5495		35
61	Therapeutic monoclonal antibodies approved by FDA in 2017. 2018 , 6,		4
60	Genetics of bipolar disorder. 2014 , 7, 33-42		68
59	New findings in the genetics of major psychoses. <i>Dialogues in Clinical Neuroscience</i> , 2010 , 12, 85-93	5.7	53
58	Ocular Findings in Children With 22q11.2 Deletion Syndrome. <i>Journal of Pediatric Ophthalmology and Strabismus</i> , 2016 , 53, 218-22	0.9	6
57	Delayed diagnosis of 22q11 deletion syndrome due to late onset hypocalcemia in a 11-year-old girl with imperforated anus. <i>Annals of Pediatric Endocrinology and Metabolism</i> , 2017 , 22, 133-138	2.9	5
56	Revisiting the Embryology of the Facial Muscles, the Superficial Musculoaponeurotic System, and the Facial Nerve. <i>Neurographics</i> , 2021 , 11, 200-228	0.4	
55	Congenital heart disease in pregnancy. <i>Deutsches A&#x0308;rztblatt International</i> , 2008 , 105, 347-54		11

54 Genetic counseling in schizophrenia. *Medical Psychiatry*, **2009**, 322-332

53 Interrupted Aortic Arch in a Patient with DiGeorge Syndrome. **2010**, 157-161

52 Late Repair of Tetralogy of Fallot. **2010**, 223-227

51 Epilepsies Associated with Malformations of Cortical Development and Inborn Errors of Metabolism. **2010**, 79-93 1

50 Schizophrenia. *Journal of the Nihon University Medical Association*, **2012**, 71, 369-374 0

49 Remédiation cognitive en psychiatrie. **2012**, 210-247

48 Late Sequelae in the Adult Patient with Congenital Heart Disease. **2014**, 2609-2632

47 Psychological Functioning and Life Experiences in Adults with Congenital Heart Disease. **2015**, 85-97

46 Psychosocial Aspects of Adults with Congenital Heart Disease. **2015**, 1-13

45 Autoimmune Hepatitis with Distal Renal Tubular Acidosis and Small Bowel Partial Malrotation. *Euroasian Journal of Hepato-gastroenterology*, **2015**, 5, 107-109 1.6 1

44 Fallot-Tetralogie (S2k). **2015**, M14.1-M14.8

43 Psychosocial Aspects of Adults with Congenital Heart Disease. **2016**, 439-451

42 Hypoparathyroidism. **2017**, 115-127

41 Randomized Study Comparing Pre-Operative Glycemic Profile in Pediatric Cardiac Surgical Patients Administered Oral Carbohydrate Solution Preoperatively versus Those Kept Fasting. *World Journal of Cardiovascular Diseases*, **2018**, 08, 298-306 0 0

40 Fishing for Digeorge Syndrome in A 40-YEAR-OLD Man. *AACE Clinical Case Reports*, **2018**, 4, 94-97 0.7

39 22q11.2 Deletion Syndrome. **2019**, 241-252

38 Family Members with Congenital Heart Disease and Hypogammaglobulinemia. **2019**, 689-693

37 Clinical characteristics, prognosis and outcome of patients with repaired tetralogy of Fallot. **2019**, 10, 91-100 0

36	Clinical and immunophenotypic characteristics of patients with chromosome 22q11.2 deletion syndrome: a single institution's experience. <i>Turk Pediatri Arsivi</i> , 2019 , 54, 28-34	0.7	1
35	Syndromic Hypoparathyroidism Due to DiGeorge Syndrome. 2020 , 45-62		
34	Integration of genetic, transcriptomic, and clinical data provides insight into 16p11.2 and 22q11.2 CNV genes.		
33	Integration of genetic, transcriptomic, and clinical data provides insight into 16p11.2 and 22q11.2 CNV genes. <i>Genome Medicine</i> , 2021 , 13, 172	14.4	1
32	Clinical Features to Predict 22q11.2 Deletion Syndrome Proven by Molecular Genetic Testing.. <i>Journal of Pediatric Genetics</i> , 2022 , 11, 22-27	0.7	1
31	DELETION 22q11.2 (VELO-CARDIO-FACIAL SYNDROME/DIGEORGE SYNDROME). 2021 , 291-316		
30	[Microdeletion syndromes (Williams syndrome and deletion syndrome 22q11) at CHU Hassan II of Fez: report of 3 observations]. <i>Pan African Medical Journal</i> , 2012 , 11, 3	1.2	
29	Social Cognition impairments in 22q11.2DS individuals with and without psychosis: A comparison study with a large population of patients with schizophrenia. <i>Schizophrenia Bulletin Open</i> ,	2.2	1
28	Ocular findings in 22q11.2 deletion syndrome: A systematic literature review and results of a Dutch multicenter study. <i>American Journal of Medical Genetics, Part A</i> , 2021 ,	2.5	0
27	What Every Internist-Endocrinologist Should Know about Rare Genetic Syndromes in Order to Prevent Needless Diagnostics, Missed Diagnoses and Medical Complications: Five Years of Internal Medicine for Rare Genetic Syndromes. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	2
26	Clinical and genomic findings in brain heterotopia: Report of a pediatric patient cohort from Romania.. <i>Experimental and Therapeutic Medicine</i> , 2022 , 23, 101	2.1	
25	Neurological manifestation of 22q11.2 deletion syndrome.. <i>Neurological Sciences</i> , 2022 , 43, 1695	3.5	0
24	Syndromic obesity with neurodevelopmental delay: Opportunities for targeted interventions.. <i>European Journal of Medical Genetics</i> , 2022 , 65, 104443	2.6	0
23	Hypocalcemia. 2022 , 219-230		0
22	Dopaminergic alterations in populations at increased risk for psychosis: A systematic review of imaging findings.. <i>Progress in Neurobiology</i> , 2022 , 213, 102265	10.9	
21	Social behavior in 16p11.2 and 22q11.2 copy number variations: Insights from mice and humans. <i>Genes, Brain and Behavior</i> , 2021 , e12787	3.6	0
20	Surgical insights and management in patients with the 22q11.2 deletion syndrome.. <i>Pediatric Surgery International</i> , 2022 , 38, 899	2.1	0
19	Craniofacial Phenotypes and Genetics of DiGeorge Syndrome. <i>Journal of Developmental Biology</i> , 2022 , 10, 18	3.5	2

18	Late maternal diagnosis of DiGeorge syndrome with congenital hypoparathyroidism following antenatal detection of the same 22q11.2 microdeletion syndrome in the fetus. <i>BMJ Case Reports</i> , 2022 , 15, e250350	0.9
17	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
16	The immune system in 22q11.2 deletion syndrome. 2022 , 102-120	
15	Identification of 22q11.2 deletion in a patient with schizophrenia and clinically diagnosed Rubinstein-Taybi syndrome. 2022 , 1,	
14	Congenital heart disease and cardiovascular abnormalities associated with 22q11.2 deletion syndrome. 2022 , 78-100	
13	Skeletal anomalies associated with 22q11.2 deletion syndrome. 2022 , 238-251	
12	Gastroenterological manifestations associated with 22q11.2 deletion syndrome. 2022 , 182-209	
11	Psychiatric profile in children and youth with 22q11.2 deletion syndrome. 2022 , 302-321	
10	Healthcare transitions for adolescents and adults with 22q11.2 deletion syndrome. 2022 , 350-369	
9	Prevalence and incidence of psychotic disorders in 22q11.2 deletion syndrome: a meta-analysis. 1-13	1
8	Adult Height, 22q11.2 Deletion Extent, and Short Stature in 22q11.2 Deletion Syndrome. 2022 , 13, 2038	
7	Fever of Unknown Origin Reveals a Missed Diagnosis of DiGeorge Syndrome in a 21-Year-Old Female. 2022 ,	
6	Updated clinical practice recommendations for managing adults with 22q11.2 deletion syndrome. 2023 , 25, 100344	
5	Updated clinical practice recommendations for managing children with 22q11.2 deletion syndrome. 2023 , 25, 100338	
4	Adult-onset obstructive sleep apnea and pediatric pharyngoplasty in 22q11.2 deletion syndrome. 2023 , 104, 49-55	
3	The Evolving Role of Animal Models in the Discovery and Development of Novel Treatments for Psychiatric Disorders. 2023 , 37-99	
2	The Relationship between Motor Symptoms, Signs, and Parkinsonism with Facial Emotion Recognition Deficits in Individuals with 22q11.2 Deletion Syndrome at High Genetic Risk for Psychosis. 2023 , 2023, 1-10	
1	Associations between acute and chronic lifetime stressors and psychosis-risk symptoms in individuals with 22q11.2 copy number variants. 1-10	

