

# CITATION REPORT

List of articles citing

**Consensus statement: chromosomal microarray is a first-tier clinical diagnostic test for individuals with developmental disabilities or congenital anomalies**

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**American Journal of Human Genetics, 2010, 86, 749-64.**

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#	Paper	IF	Citations
2128	Molecular genetics and diagnostic techniques. <b>2000</b> , 481-497		
2127	Prenatal genetic screening and diagnosis for pediatricians. <b>2010</b> , 22, 809-13		1
2126	Deletion 17q12 is a recurrent copy number variant that confers high risk of autism and schizophrenia. <i>American Journal of Human Genetics</i> , <b>2010</b> , 87, 618-30	11	237
2125	Clinically detectable copy number variations in a Canadian catchment population of schizophrenia. <b>2010</b> , 44, 1005-9		58
2124	An introduction to standardized clinical nomenclature for dysmorphic features: the Elements of Morphology project. <b>2010</b> , 8, 56		
2123	Pathogenesis, neuroimaging and management in children with cerebral palsy born preterm. <b>2010</b> , 16, 302-12		13
2122	Mosaic Down syndrome in a patient with low-level mosaicism detected by microarray. <b>2010</b> , 152A, 3154-6		15
2121	Comparative analysis of copy number detection by whole-genome BAC and oligonucleotide array CGH. <b>2010</b> , 3, 11		49
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2118	Fragile X and autism: Intertwined at the molecular level leading to targeted treatments. <b>2010</b> , 1, 12		173
2117	Diagnostic utility of array-based comparative genomic hybridization (aCGH) in a prenatal setting. <b>2010</b> , 30, 1131-7		62
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1487	Low Rates of Genetic Testing in Children With Developmental Delays, Intellectual Disability, and Autism Spectrum Disorders. <b>2015</b> , 2, 2333794X15623717	10
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1437	Diagnóstico de las anomalías cromosómicas en patología constitucional. <b>2015</b> , 19, 1-8		
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1429	Human Structural Variation: Mechanisms of Chromosome Rearrangements. <b>2015</b> , 31, 587-599		118
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1426	Fetal CNAPS DNA/RNA. <b>2015</b> , 165-212		
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1422	The clustering of functionally related genes contributes to CNV-mediated disease. <b>2015</b> , 25, 802-13		24
1421	Lymphovascular invasion and histologic grade are associated with specific genomic profiles in invasive carcinomas of the breast. <b>2015</b> , 36, 1835-48		19
1420	Phelan-McDermid Syndrome and SHANK3: Implications for Treatment. <b>2015</b> , 12, 620-30		40
1419	Genetics of Neuromuscular Disorders. <b>2015</b> , 17-31		1
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1416	p600/UBR4 in the central nervous system. <b>2015</b> , 72, 1149-60		20
1415	Connecting the CNTNAP2 Networks with Neurodevelopmental Disorders. <b>2015</b> , 6, 7-22		63
1414	Gene discovery and functional assessment of rare copy-number variants in neurodevelopmental disorders. <b>2015</b> , 14, 315-28		19
1413	Metabolic and Genetic Causes of Autism. <b>2015</b> , 209-217		
1412	Array CGH as a first-tier test for neonates with congenital heart disease. <b>2015</b> , 25, 115-22		11
1411	Variants of unknown significance on chromosomal microarray analysis: parental perspectives. <b>2015</b> , 6, 343-9		23
1410	Navigating the current landscape of clinical genetic testing for inherited retinal dystrophies. <b>2015</b> , 17, 245-52		44

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1408	Genomic imbalances in pediatric patients with chronic kidney disease. <b>2015</b> , 125, 2171-8	55
1407	Progestogens in Obstetrics and Gynecology. <b>2015</b> ,	1
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1403	Detection of Chromosomal Aberrations in Clinical Practice: From Karyotype to Genome Sequence. <b>2015</b> , 16, 309-26	40
1402	A 5-month-old boy with delay in growth and development and decreased muscle tone. <b>2015</b> , 61, 50-4	
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1373	Asymmetric aneuploidy in mesenchymal stromal cells detected by in situ karyotyping and fluorescence in situ hybridization: suggestions for reference values for stem cells. <b>2015</b> , 24, 77-92	16
1372	Recent advances of genomic testing in perinatal medicine. <b>2015</b> , 39, 44-54	27
1371	FISH for 22q11.2 deletion not cost-effective for infants with congenital heart disease with microarray. <b>2015</b> , 36, 531-6	11
1370	A new syndrome of intellectual disability with dysmorphism due to TBL1XR1 deletion. <b>2015</b> , 167A, 164-8	30
1369	Circulating Nucleic Acids in Early Diagnosis, Prognosis and Treatment Monitoring. <b>2015</b> ,	7
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1362	Delineating the 15q13.3 microdeletion phenotype: a case series and comprehensive review of the literature. <b>2015</b> , 17, 149-57	80
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1360	Detecting somatic mosaicism: considerations and clinical implications. <b>2015</b> , 87, 554-62	32
1359	Genomic Applications in Pathology. <b>2015</b> ,	
1358	Genomewide Array Comparative Genomic Hybridization in 55 Japanese Normokaryotypic Patients with Non-Syndromic Intellectual Disability. <b>2016</b> , 02,	1
1357	Genetics of Parkinson Disease: The Role of Copy Number Variations. <b>2016</b> ,	1
1356	Cytogenetics at the University of Cape Town: A 45-year journey. <b>2016</b> , 106, S29-32	

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1353	Recent advances in prenatal genetic screening and testing. <b>2016</b> , 5, 2591	44
1352	Experimental Tools for the Identification of Specific Genes in Autism Spectrum Disorders and Intellectual Disability. <b>2016</b> , 3-12	1
1351	What Do Parents Think about Chromosomal Microarray Testing? A Qualitative Report from Parents of Children with Autism Spectrum Disorders. <b>2016</b> , 2016, 6852539	5
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1347	Analysis of chromosomal abnormalities by CGH-array in patients with dysmorphic and intellectual disability with normal karyotype. <b>2016</b> , 14, 30-4	5
1346	Fluorescence In situ Hybridization: Cell-Based Genetic Diagnostic and Research Applications. <b>2016</b> , 4, 89	86
1345	Genetic Evaluation and Use of Chromosome Microarray in Patients with Isolated Heart Defects: Benefits and Challenges of a New Model in Cardiovascular Care. <b>2016</b> , 3, 19	9
1344	The Current Landscape of Genetic Testing in Cardiovascular Malformations: Opportunities and Challenges. <b>2016</b> , 3, 22	15
1343	Chromosomal Microarray Analysis of Consecutive Individuals with Autism Spectrum Disorders Using an Ultra-High Resolution Chromosomal Microarray Optimized for Neurodevelopmental Disorders. <b>2016</b> , 17,	40
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1341	Clinical Utility of a Comprehensive, Whole Genome CMA Testing Platform in Pediatrics: A Prospective Randomized Controlled Trial of Simulated Patients in Physician Practices. <b>2016</b> , 11, e0169064	4
1340	Array-CGH analysis in patients with intellectual disability and/or congenital malformations in Brazil. <b>2016</b> , 15,	7
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1336	Detecting Copy Number Variation via Next Generation Technology. <b>2016</b> , 4, 74-85	14
1335	Clinical exome sequencing in neurogenetic and neuropsychiatric disorders. <b>2016</b> , 1366, 49-60	20
1334	What results to disclose, when, and who decides? Healthcare professionals' views on prenatal chromosomal microarray analysis. <b>2016</b> , 36, 252-9	24
1333	DICER1 deletion and 14q32 microdeletion syndrome: an additional case and a review of the literature. <b>2016</b> , 25, 37-40	6
1332	Is one diagnosis the whole story? patients with double diagnoses. <b>2016</b> , 170, 2338-48	16
1331	Autism and chromosome abnormalities-A review. <b>2016</b> , 29, 620-7	8
1330	Novel copy number variants and major limb reduction malformation: Report of three cases. <b>2016</b> , 170A, 1245-50	7
1329	Detection of 1p36 deletion by clinical exome-first diagnostic approach. <b>2016</b> , 3, 16006	18
1328	Clinical utility of array comparative genomic hybridisation in prenatal setting. <b>2016</b> , 17, 81	11
1327	Whole Genome Sequencing Expands Diagnostic Utility and Improves Clinical Management in Pediatric Medicine. <b>2016</b> , 1,	208
1326	Diagnostic yield of array CGH in patients with autism spectrum disorder in Hong Kong. <b>2016</b> , 5, 18	6
1325	Chromosome 10q26 deletion syndrome: Two new cases and a review of the literature. <b>2016</b> , 14, 5134-5140	16
1324	Assessment of copy number variations in 120 patients with Poland syndrome. <b>2016</b> , 17, 89	14
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1322	Mutations in HECW2 are associated with intellectual disability and epilepsy. <b>2016</b> , 53, 697-704	39
1321	CNV analysis in Chinese children of mental retardation highlights a sex differentiation in parental contribution to de novo and inherited mutational burdens. <b>2016</b> , 6, 25954	13
1320	The importance of copy number variation in congenital heart disease. <b>2016</b> , 1, 16031	42

1319	Prenatal diagnosis of fetal multicystic dysplastic kidney via high-resolution whole-genome array. <b>2016</b> , 31, 1693-8	22
1318	Autism Spectrum Disorder. <b>2016</b> , 1497-1528	
1317	Novel bioinformatic developments for exome sequencing. <b>2016</b> , 135, 603-14	27
1316	Chromosomal microarray in prenatal diagnosis: case studies and clinical challenges. <b>2016</b> , 13, 249-255	4
1315	A homozygous truncating mutation in PUS3 expands the role of tRNA modification in normal cognition. <b>2016</b> , 135, 707-13	78
1314	A null mutation in TNIK defines a novel locus for intellectual disability. <b>2016</b> , 135, 773-8	14
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1312	Chromosomal microarray analysis in clinical evaluation of neurodevelopmental disorders-reporting a novel deletion of SETDB1 and illustration of counseling challenge. <b>2016</b> , 80, 371-81	12
1311	What Is New in Genetics and Genomics?. <b>2016</b> , 703-712	
1310	A Clinician's perspective on clinical exome sequencing. <b>2016</b> , 135, 643-54	28
1309	Noninvasive prenatal screening or advanced diagnostic testing: caveat emptor. <b>2016</b> , 215, 298-305	59
1308	Recommendations for the integration of genomics into clinical practice. <b>2016</b> , 18, 1075-1084	92
1307	Síndrome de deleción 22q11: bases embriológicas y algoritmo diagnóstico. <b>2016</b> , 23, 443-452	1
1306	Idiopathic focal epilepsies: the "lost tribe". <b>2016</b> , 18, 252-88	42
1305	An 8.4-Mb 3q26.33-3q28 microdeletion in a patient with blepharophimosis-intellectual disability syndrome and a review of the literature. <b>2016</b> , 4, 824-30	3
1304	Chromosome microarray proficiency testing and analysis of quality metric data trends through an external quality assessment program for Australasian laboratories. <b>2016</b> , 48, 586-96	
1303	Copy number variations in 375 patients with oesophageal atresia and/or tracheoesophageal fistula. <b>2016</b> , 24, 1715-1723	17
1302	Validation of a Chromosomal Microarray for Prenatal Diagnosis Using a Prospective Cohort of Pregnancies with Increased Risk for Chromosome Abnormalities. <b>2016</b> , 20, 791-798	2

1301	Molecular Genetics Methods for Developmental Scientists. <b>2016</b> , 1-38	1
1300	The use of chromosomal microarray for 'prenatal' diagnosis. <b>2016</b> , 215, B2-9	88
1299	Inherited ichthyosis: Non-syndromic forms. <b>2016</b> , 43, 242-51	92
1298	Tentative clinical diagnosis of Lujan-Fryns syndrome--A conglomeration of different genetic entities?. <b>2016</b> , 170A, 94-102	10
1297	Genetic Counseling for Autism Spectrum Disorder in an Evolving Theoretical Landscape. <b>2016</b> , 4, 147-153	10
1296	Preclinical development and qualification of ZFN-mediated CCR5 disruption in human hematopoietic stem/progenitor cells. <b>2016</b> , 3, 16067	73
1295	Genetic Approach to Diagnosis of Intellectual Disability. <b>2016</b> , 83, 1141-9	6
1294	Interstitial Chromosome 3p14.1 Deletion due to a Maternal Insertion: Phenotype and Association with Balanced Parental Rearrangement. <b>2016</b> , 7, 43-8	3
1293	X-Linked Candidate Genes for a Ciliopathy-Like Disorder. <b>2016</b> , 7, 37-42	
1292	ANALYTIC PROCEDURES. <b>2016</b> , 739-753	
1291	POSTANALYTIC PROCEDURES. <b>2016</b> , 755-779	
1290	Somatic Mosaicism and Neurological Diseases. <b>2016</b> , 179-199	3
1289	Neurodevelopmental Disorders, Causes, and Consequences. <b>2016</b> , 587-599	
1288	3Disease Browser: A Web server for integrating 3D genome and disease-associated chromosome rearrangement data. <b>2016</b> , 6, 34651	22
1287	. <b>2016</b> ,	
1286	Genetics of Combined Pituitary Hormone Deficiency: Roadmap into the Genome Era. <b>2016</b> , 37, 636-675	106
1285	Subtelomeric Copy Number Variations: The Importance of 4p/4q Deletions in Patients with Congenital Anomalies and Developmental Disability. <b>2016</b> , 149, 241-246	5
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1281	Associated anomalies in cases with anotia and microtia. <b>2016</b> , 59, 607-614	10
1280	Indexing Effects of Copy Number Variation on Genes Involved in Developmental Delay. <b>2016</b> , 6, 28663	26
1279	RYR2, PTDSS1 and AREG genes are implicated in a Lebanese population-based study of copy number variation in autism. <b>2016</b> , 6, 19088	15
1278	Intragenic CNVs for epigenetic regulatory genes in intellectual disability: Survey identifies pathogenic and benign single exon changes. <b>2016</b> , 170, 2916-2926	11
1277	Testing the Complex Child: CGH Array, WES, Clinical Exome, WGS. <b>2016</b> , 4, 155-163	
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