

Marwan S Shinawi

List of Publications by Year in descending order

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Version: 2024-02-01

153
papers

8,391
citations

76294

40
h-index

54882

84
g-index

162
all docs

162
docs citations

162
times ranked

14122
citing authors

#	ARTICLE	IF	CITATIONS
1	<i>De novo</i> missense variants in FBXO11 alter its protein expression and subcellular localization. <i>Human Molecular Genetics</i> , 2022, 31, 440-454.	1.4	7
2	Semaphorin-Plexin Signaling: From Axonal Guidance to a New X-Linked Intellectual Disability Syndrome. <i>Pediatric Neurology</i> , 2022, 126, 65-73.	1.0	8
3	<i>DNMT3A</i> overgrowth syndrome is associated with the development of hematopoietic malignancies in children and young adults. <i>Blood</i> , 2022, 139, 461-464.	0.6	9
4	Delineation of the 1q24.3 microdeletion syndrome provides further evidence for the potential role of non-coding RNAs in regulating the skeletal phenotype. <i>Bone</i> , 2021, 142, 115705.	1.4	2
5	De novo variants in SNAP25 cause an early-onset developmental and epileptic encephalopathy. <i>Genetics in Medicine</i> , 2021, 23, 653-660.	1.1	20
6	Wilms tumor in patients with osteopathia striata with cranial sclerosis. <i>European Journal of Human Genetics</i> , 2021, 29, 396-401.	1.4	10
7	Response to Mounts and Besser. <i>Genetics in Medicine</i> , 2021, 23, 240-242.	1.1	1
8	New Cohort of Patients With CEDNIK Syndrome Expands the Phenotypic and Genotypic Spectra. <i>Neurology: Genetics</i> , 2021, 7, e553.	0.9	10
9	Paroxysmal Kinesigenic Dyskinesia in Twins With Chromosome 16p11.2 Duplication Syndrome. <i>Neurology: Genetics</i> , 2021, 7, e549.	0.9	1
10	Biallelic <i>ASCC1</i> variants including a novel intronic variant result in expanded phenotypic spectrum of spinal muscular atrophy with congenital bone fractures 2 (<i>SMABF2</i>). <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 2190-2197.	0.7	4
11	Immunodeficiency and bone marrow failure with mosaic and germline TLR8 gain of function. <i>Blood</i> , 2021, 137, 2450-2462.	0.6	47
12	Loss-of-function and missense variants in NSD2 cause decreased methylation activity and are associated with a distinct developmental phenotype. <i>Genetics in Medicine</i> , 2021, 23, 1474-1483.	1.1	24
13	Biallelic variants in <i>RNU12</i> cause CDAGS syndrome. <i>Human Mutation</i> , 2021, 42, 1042-1052.	1.1	5
14	Truncating variants in the SHANK1 gene are associated with a spectrum of neurodevelopmental disorders. <i>Genetics in Medicine</i> , 2021, 23, 1912-1921.	1.1	5
15	Functional and epigenetic phenotypes of humans and mice with DNMT3A Overgrowth Syndrome. <i>Nature Communications</i> , 2021, 12, 4549.	5.8	21
16	Autosomal Dominant ANO5-Related Disorder Associated With Myopathy and Gnathodiaphyseal Dysplasia. <i>Neurology: Genetics</i> , 2021, 7, e612.	0.9	4
17	A phase 1/2 open label nonrandomized clinical trial of intravenous 2-hydroxypropyl- β -cyclodextrin for acute liver disease in infants with Niemann-Pick C1. <i>Molecular Genetics and Metabolism Reports</i> , 2021, 28, 100772.	0.4	8
18	Novel exon-skipping variant disrupting the basic domain of HCFC1 causes intellectual disability without metabolic abnormalities in both male and female patients. <i>Journal of Human Genetics</i> , 2021, 66, 717-724.	1.1	10

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19	Identification of disease-linked hyperactivating mutations in UBE3A through large-scale functional variant analysis. <i>Nature Communications</i> , 2021, 12, 6809.	5.8	10
20	Intragenic CNTN4 copy number variants associated with a spectrum of neurobehavioral phenotypes. <i>European Journal of Medical Genetics</i> , 2020, 63, 103736.	0.7	11
21	Sorting nexin 27 (<i>SNX27</i>) variants associated with seizures, developmental delay, behavioral disturbance, and subcortical brain abnormalities. <i>Clinical Genetics</i> , 2020, 97, 437-446.	1.0	10
22	Delineation of a Human Mendelian Disorder of the DNA Demethylation Machinery: TET3 Deficiency. <i>American Journal of Human Genetics</i> , 2020, 106, 234-245.	2.6	56
23	Overcoming presynaptic effects of VAMP2 mutations with 4-aminopyridine treatment. <i>Human Mutation</i> , 2020, 41, 1999-2011.	1.1	11
24	A Recurrent Gain-of-Function Mutation in CLCN6, Encoding the Cl ⁻ /H ⁺ -Exchanger, Causes Early-Onset Neurodegeneration. <i>American Journal of Human Genetics</i> , 2020, 107, 1062-1077.	2.6	23
25	De Novo and Bi-allelic Pathogenic Variants in NARS1 Cause Neurodevelopmental Delay Due to Toxic Gain-of-Function and Partial Loss-of-Function Effects. <i>American Journal of Human Genetics</i> , 2020, 107, 311-324.	2.6	32
26	Phenotypic spectrum and transcriptomic profile associated with germline variants in TRAF7. <i>Genetics in Medicine</i> , 2020, 22, 1215-1226.	1.1	22
27	Abnormally increased carotid intima media-thickness and elasticity in patients with Morquio A disease. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 73.	1.2	5
28	Delineation of phenotypes and genotypes related to cohesin structural protein RAD21. <i>Human Genetics</i> , 2020, 139, 575-592.	1.8	24
29	Pathogenic variants in <i>TNRC6B</i> cause a genetic disorder characterised by developmental delay/intellectual disability and a spectrum of neurobehavioural phenotypes including autism and ADHD. <i>Journal of Medical Genetics</i> , 2020, 57, 717-724.	1.5	14
30	Diagnostic testing for uniparental disomy: a points to consider statement from the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2020, 22, 1133-1141.	1.1	89
31	AMPA receptor GluA2 subunit defects are a cause of neurodevelopmental disorders. <i>Nature Communications</i> , 2019, 10, 3094.	5.8	150
32	De novo substitutions of TRPM3 cause intellectual disability and epilepsy. <i>European Journal of Human Genetics</i> , 2019, 27, 1611-1618.	1.4	45
33	CTCF variants in 39 individuals with a variable neurodevelopmental disorder broaden the mutational and clinical spectrum. <i>Genetics in Medicine</i> , 2019, 21, 2723-2733.	1.1	48
34	2-Pyrrolidinone and Succinimide as Clinical Screening Biomarkers for GABA-Transaminase Deficiency: Anti-seizure Medications Impact Accurate Diagnosis. <i>Frontiers in Neuroscience</i> , 2019, 13, 394.	1.4	23
35	A mutation in Site-1 Protease is associated with a complex phenotype that includes episodic hyperCKemia and focal myoedema. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e00733.	0.6	13
36	A pathogenic CtBP1 missense mutation causes altered cofactor binding and transcriptional activity. <i>Neurogenetics</i> , 2019, 20, 129-143.	0.7	16

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37	Phenotype and response to growth hormone therapy in siblings with B4GALT7 deficiency. <i>Bone</i> , 2019, 124, 14-21.	1.4	9
38	Birth Defects Among 788 Children Born to Gulf War Veterans Based on Physical Examination. <i>Journal of Occupational and Environmental Medicine</i> , 2019, 61, 263-270.	0.9	2
39	Variants in DOCK3 cause developmental delay and hypotonia. <i>European Journal of Human Genetics</i> , 2019, 27, 1225-1234.	1.4	15
40	Novel parent-of-origin-specific differentially methylated loci on chromosome 16. <i>Clinical Epigenetics</i> , 2019, 11, 60.	1.8	18
41	Pathogenic Variants in GPC4 Cause Keipert Syndrome. <i>American Journal of Human Genetics</i> , 2019, 104, 914-924.	2.6	23
42	Variants in TCF20 in neurodevelopmental disability: description of 27 new patients and review of literature. <i>Genetics in Medicine</i> , 2019, 21, 2036-2042.	1.1	23
43	De Novo Variants Disrupting the HX Repeat Motif of ATN1 Cause a Recognizable Non-Progressive Neurocognitive Syndrome. <i>American Journal of Human Genetics</i> , 2019, 104, 542-552.	2.6	19
44	Special Therapy and Psychosocial Needs Identified in a Multidisciplinary Cancer Predisposition Syndrome Clinic. <i>Journal of Pediatric Hematology/Oncology</i> , 2019, 41, 133-136.	0.3	1
45	Lamin B receptor-related disorder is associated with a spectrum of skeletal dysplasia phenotypes. <i>Bone</i> , 2019, 120, 354-363.	1.4	11
46	Inherited Deletion of 1q, Hyperparathyroidism and Signs of Y-chromosomal Influence in a Patient with Turner Syndrome. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2019, 11, 88-93.	0.4	3
47	WNT Signaling Perturbations Underlie the Genetic Heterogeneity of Robinow Syndrome. <i>American Journal of Human Genetics</i> , 2018, 102, 27-43.	2.6	88
48	Gnathodiaphyseal dysplasia: Severe atypical presentation with novel heterozygous mutation of the anoctamin gene (ANO5). <i>Bone</i> , 2018, 107, 161-171.	1.4	23
49	Child Neurology: Brown-Vialetto-Van Laere syndrome. <i>Neurology</i> , 2018, 91, 938-941.	1.5	10
50	Variable cardiovascular phenotypes associated with <i>SMAD2</i> pathogenic variants. <i>Human Mutation</i> , 2018, 39, 1875-1884.	1.1	23
51	DeSanto-Shinawi Syndrome: First Case in South America. <i>Molecular Syndromology</i> , 2018, 9, 154-158.	0.3	11
52	Mutations in the <i>PH</i> Domain of <i>DNM1L</i> are associated with a nonepileptic phenotype characterized by developmental delay and neurobehavioral abnormalities. <i>Molecular Genetics & Genomic Medicine</i> , 2018, 6, 294-300.	0.6	19
53	Functional characterization of biallelic RTTN variants identified in an infant with microcephaly, simplified gyral pattern, pontocerebellar hypoplasia, and seizures. <i>Pediatric Research</i> , 2018, 84, 435-441.	1.1	11
54	De Novo Variants in the F-Box Protein FBXO11 in 20 Individuals with a Variable Neurodevelopmental Disorder. <i>American Journal of Human Genetics</i> , 2018, 103, 305-316.	2.6	48

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55	BCL11B mutations in patients affected by a neurodevelopmental disorder with reduced type 2 innate lymphoid cells. <i>Brain</i> , 2018, 141, 2299-2311.	3.7	81
56	De novo mutation screening in childhood-onset cerebellar atrophy identifies gain-of-function mutations in the CACNA1G calcium channel gene. <i>Brain</i> , 2018, 141, 1998-2013.	3.7	67
57	CpG Island Hypermethylation Mediated by DNMT3A Is a Consequence of AML Progression. <i>Cell</i> , 2017, 168, 801-816.e13.	13.5	177
58	Support for the Diagnosis of CHARGE Syndrome—Reply. <i>JAMA Otolaryngology - Head and Neck Surgery</i> , 2017, 143, 635.	1.2	0
59	The Exome Clinic and the role of medical genetics expertise in the interpretation of exome sequencing results. <i>Genetics in Medicine</i> , 2017, 19, 1040-1048.	1.1	85
60	Adult-onset dystonia with marfanoid features. <i>Neurology: Clinical Practice</i> , 2017, 7, e31-e34.	0.8	1
61	CEDNIK. <i>Child Neurology Open</i> , 2017, 4, 2329048X1773321.	0.5	16
62	The spectrum of <i>DNMT3A</i> variants in Tattonâ€“Brownâ€“Rahman syndrome overlaps with that in hematologic malignancies. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 3022-3028.	0.7	42
63	Survival among children with â€œLethalâ€“congenital contracture syndrome 11 caused by novel mutations in the gliomedin gene (<i>GLDN</i>). <i>Human Mutation</i> , 2017, 38, 1477-1484.	1.1	19
64	Heterozygous variants in <i>ACTL6A</i> , encoding a component of the BAF complex, are associated with intellectual disability. <i>Human Mutation</i> , 2017, 38, 1365-1371.	1.1	27
65	Prevalence of Semicircular Canal Hypoplasia in Patients With CHARGE Syndrome. <i>JAMA Otolaryngology - Head and Neck Surgery</i> , 2017, 143, 168.	1.2	24
66	The Cognitive and Behavioral Phenotypes of Individuals with CHRNA7 Duplications. <i>Journal of Autism and Developmental Disorders</i> , 2017, 47, 549-562.	1.7	68
67	Neurexin 2 nonsense variant associated with anxiety, autism, intellectual disability, hyperphagia, and obesity. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 213-216.	0.7	68
68	<i>FBXL4</i> defects are common in patients with congenital lactic acidemia and encephalomyopathic mitochondrial DNA depletion syndrome. <i>Clinical Genetics</i> , 2017, 91, 634-639.	1.0	18
69	Clinical Characterization of Patients With Autosomal Dominant Short Stature due to Aggrecan Mutations. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017, 102, 460-469.	1.8	95
70	Multigenerational autosomal dominant inheritance of 5p chromosomal deletions. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 583-593.	0.7	21
71	A recurrent de novo CTBP1 mutation is associated with developmental delay, hypotonia, ataxia, and tooth enamel defects. <i>Neurogenetics</i> , 2016, 17, 173-178.	0.7	32
72	TSHZ3 deletion causes an autism syndrome and defects in cortical projection neurons. <i>Nature Genetics</i> , 2016, 48, 1359-1369.	9.4	69

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73	The expanding clinical phenotype of Bosch-Boonstra-Schaaf optic atrophy syndrome: 20 new cases and possible genotype-phenotype correlations. <i>Genetics in Medicine</i> , 2016, 18, 1143-1150.	1.1	64
74	Recurrent Muscle Weakness with Rhabdomyolysis, Metabolic Crises, and Cardiac Arrhythmia Due to Bi-allelic TANGO2 Mutations. <i>American Journal of Human Genetics</i> , 2016, 98, 347-357.	2.6	98
75	<i>STXPB1</i> encephalopathy. <i>Neurology</i> , 2016, 86, 954-962.	1.5	264
76	Digynic triploidy: utility and challenges of noninvasive prenatal testing. <i>Clinical Case Reports (discontinued)</i> , 2015, 3, 406-410.	0.2	8
77	Multi-systemic involvement in NGLY1-related disorder caused by two novel mutations. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 816-820.	0.7	45
78	WACloss-of-function mutations cause a recognisable syndrome characterised by dysmorphic features, developmental delay and hypotonia and recapitulate 10p11.23 microdeletion syndrome. <i>Journal of Medical Genetics</i> , 2015, 52, 754-761.	1.5	41
79	<i>TBX6</i> Null Variants and a Common Hypomorphic Allele in Congenital Scoliosis. <i>New England Journal of Medicine</i> , 2015, 372, 341-350.	13.9	239
80	Autosomal recessive posterior column ataxia with retinitis pigmentosa caused by novel mutations in the <i>FLVCR1</i> gene. <i>International Journal of Neuroscience</i> , 2015, 125, 43-49.	0.8	21
81	FGFR3-related condition: a skeletal dysplasia with similarities to thanatophoric dysplasia and SADDAN due to Lys650Met. <i>Skeletal Radiology</i> , 2015, 44, 441-445.	1.2	5
82	De Novo Mutations in SIK1 Cause a Spectrum of Developmental Epilepsies. <i>American Journal of Human Genetics</i> , 2015, 96, 1009.	2.6	1
83	A Novel Mutation in Isoform 3 of the Plasma Membrane Ca ²⁺ Pump Impairs Cellular Ca ²⁺ Homeostasis in a Patient with Cerebellar Ataxia and Laminin Subunit 11± Mutations. <i>Journal of Biological Chemistry</i> , 2015, 290, 16132-16141.	1.6	41
84	De Novo Mutations in SIK1 Cause a Spectrum of Developmental Epilepsies. <i>American Journal of Human Genetics</i> , 2015, 96, 682-690.	2.6	48
85	A 5-Month-Old Boy with Delay in Growth and Development and Decreased Muscle Tone. <i>Clinical Chemistry</i> , 2015, 61, 50-54.	1.5	0
86	Intragenic <i>CAMTA1</i> deletions are associated with a spectrum of neurobehavioral phenotypes. <i>Clinical Genetics</i> , 2015, 87, 478-482.	1.0	18
87	Mutations in <i>COQ4</i> , an essential component of coenzyme Q biosynthesis, cause lethal neonatal mitochondrial encephalomyopathy. <i>Journal of Medical Genetics</i> , 2015, 52, 627-635.	1.5	48
88	Scoliosis and vertebral anomalies: Additional abnormal phenotypes associated with chromosome 16p11.2 rearrangement. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 1118-1126.	0.7	38
89	Brain MRI abnormalities and spectrum of neurological and clinical findings in three patients with proximal 16p11.2 microduplication. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 2003-2012.	0.7	19
90	Molecular and phenotypic characterization of atypical Williams-Beuren syndrome. <i>Clinical Genetics</i> , 2014, 86, 487-491.	1.0	7

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91	Dosage Changes of a Segment at 17p13.1 Lead to Intellectual Disability and Microcephaly as a Result of Complex Genetic Interaction of Multiple Genes. <i>American Journal of Human Genetics</i> , 2014, 95, 565-578.	2.6	40
92	Heterozygous 24-polyalanine repeats in the <i>PHOX2B</i> gene with different manifestations across three generations. <i>Pediatric Pulmonology</i> , 2014, 49, E13-E16.	1.0	15
93	Two novel RAD21 mutations in patients with mild Cornelia de Lange syndrome-like presentation and report of the first familial case. <i>Gene</i> , 2014, 537, 279-284.	1.0	31
94	<i>NR2F1</i> haploinsufficiency is associated with optic atrophy, dysmorphism and global developmental delay. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 377-381.	0.7	40
95	Duplication of 20p12.3 associated with familial Wolff-Parkinson-White syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 137-144.	0.7	10
96	Haploinsufficiency of <i>ZNF238</i> is associated with corpus callosum abnormalities in 1q44 deletions. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 711-716.	0.7	28
97	Transient Massive Trimethylaminuria Associated with Food Protein-Induced Enterocolitis Syndrome. <i>JIMD Reports</i> , 2013, 12, 11-15.	0.7	13
98	A common X-linked inborn error of carnitine biosynthesis may be a risk factor for nondysmorphic autism. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 7974-7981.	3.3	118
99	Acute Intermittent Porphyrinuria. <i>Journal of Child Neurology</i> , 2012, 27, 917-921.	0.7	14
100	Duplication of <i>OCRL</i> and adjacent genes associated with autism but not Lowe syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2012, 158A, 2602-2605.	0.7	13
101	ADULT syndrome due to an R243W mutation in <i>TP63</i> . <i>International Journal of Dermatology</i> , 2012, 51, 693-696.	0.5	12
102	Recurrent deletions and reciprocal duplications of 10q11.21q11.23 including CHAT and SLC18A3 are likely mediated by complex low-copy repeats. <i>Human Mutation</i> , 2012, 33, 165-179.	1.1	45
103	Early-onset Hepatic Fibrosis in Lysinuric Protein Intolerance. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2011, 53, 695-698.	0.9	6
104	Expanding the clinical spectrum of the 16p11.2 chromosomal rearrangements: three patients with syringomyelia. <i>European Journal of Human Genetics</i> , 2011, 19, 152-156.	1.4	47
105	11p14.1 microdeletions associated with ADHD, autism, developmental delay, and obesity. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 1272-1280.	0.7	84
106	Desmosterolosis: phenotypic and molecular characterization of a third case and review of the literature. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 1597-1604.	0.7	52
107	Cobalamin F Disease Detected by Newborn Screening and Follow-up on a 14-Year-Old Patient. <i>Pediatrics</i> , 2011, 128, e1636-e1640.	1.0	13
108	Copy number gain at Xp22.31 includes complex duplication rearrangements and recurrent triplications. <i>Human Molecular Genetics</i> , 2011, 20, 1975-1988.	1.4	74

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109	Long-term, open-labeled extension study of idursulfase in the treatment of Hunter syndrome. <i>Genetics in Medicine</i> , 2011, 13, 95-101.	1.1	190
110	Known and Possible Roles of Epigenetics in Autism. , 2011, , 737-755.		0
111	McCune-Albright syndrome presenting with unilateral macroorchidism and bilateral testicular masses. <i>Pediatric Radiology</i> , 2010, 40, 16-20.	1.1	6
112	Structures and molecular mechanisms for common 15q13.3 microduplications involving CHRNA7: benign or pathological?. <i>Human Mutation</i> , 2010, 31, 840-850.	1.1	111
113	Mixed gonadal dysgenesis in a child with isodicentric y chromosome: Does the relative proportion of the 45,X line really matter?. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 1832-1837.	0.7	23
114	Recurrent reciprocal 16p11.2 rearrangements associated with global developmental delay, behavioural problems, dysmorphism, epilepsy, and abnormal head size. <i>Journal of Medical Genetics</i> , 2010, 47, 332-341.	1.5	447
115	Increased Homocysteine in a Patient Diagnosed with Marfan Syndrome. <i>Clinical Chemistry</i> , 2010, 56, 1665-1668.	1.5	3
116	Progressive Myopathy With Multiple Symmetric Lipomatosis. <i>Archives of Neurology</i> , 2009, 66, 1576-7.	4.9	2
117	Mitochondrial Neurogastrointestinal Encephalopathy Due to Mutations in RRM2B. <i>Archives of Neurology</i> , 2009, 66, 1028-32.	4.9	103
118	Is this the Coffinâ€“Siris syndrome or the BOD syndrome?. <i>American Journal of Medical Genetics, Part A</i> , 2009, 149A, 559-562.	0.7	12
119	The Xp contiguous deletion syndrome and autism. <i>American Journal of Medical Genetics, Part A</i> , 2009, 149A, 1138-1148.	0.7	19
120	Atypical presentation of VLCAD deficiency associated with a novel <i>ACADVL</i> splicing mutation. <i>Muscle and Nerve</i> , 2009, 39, 374-382.	1.0	23
121	Mutations involved in Aicardi-GoutiÃˆres syndrome implicate SAMHD1 as regulator of the innate immune response. <i>Nature Genetics</i> , 2009, 41, 829-832.	9.4	610
122	A small recurrent deletion within 15q13.3 is associated with a range of neurodevelopmental phenotypes. <i>Nature Genetics</i> , 2009, 41, 1269-1271.	9.4	171
123	The <i>MTHFR 677</i> CÃˆT polymorphism and behaviors in children with autism: exploratory genotypeâ€“phenotype correlations. <i>Autism Research</i> , 2009, 2, 98-108.	2.1	57
124	Microdeletions including YWHAE in the Miller-Dieker syndrome region on chromosome 17p13.3 result in facial dysmorphisms, growth restriction, and cognitive impairment. <i>Journal of Medical Genetics</i> , 2009, 46, 825-833.	1.5	112
125	Microdeletion 15q13.3: a locus with incomplete penetrance for autism, mental retardation, and psychiatric disorders. <i>Journal of Medical Genetics</i> , 2009, 46, 382-388.	1.5	213
126	Lowâ€“level mosaicism of trisomy 14: Phenotypic and molecular characterization. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 1395-1405.	0.7	34

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127	Delineation of the proximal 3q microdeletion syndrome. American Journal of Medical Genetics, Part A, 2008, 146A, 1729-1735.	0.7	15
128	Prader-Willi phenotype caused by paternal deficiency for the HBII-85 C/D box small nucleolar RNA cluster. Nature Genetics, 2008, 40, 719-721.	9.4	533
129	Recurrent reciprocal 1q21.1 deletions and duplications associated with microcephaly or macrocephaly and developmental and behavioral abnormalities. Nature Genetics, 2008, 40, 1466-1471.	9.4	535
130	The array CGH and its clinical applications. Drug Discovery Today, 2008, 13, 760-770.	3.2	171
131	15q13q14 deletions: Phenotypic characterization and molecular delineation by comparative genomic hybridization. American Journal of Medical Genetics, Part A, 2008, 146A, 1933-1941.	0.7	13
132	Syndromic thrombocytopenia and predisposition to acute myelogenous leukemia caused by constitutional microdeletions on chromosome 21q. Blood, 2008, 112, 1042-1047.	0.6	74
133	Lymphedema of the Lower Extremity: Is It Genetic or Nongenetic?. Clinical Pediatrics, 2007, 46, 835-841.	0.4	13
134	Preaxial polydactyly in neurofibromatosis 1. Clinical Dysmorphology, 2007, 16, 193-194.	0.1	6
135	Hyperhomocysteinemia and cobalamin disorders. Molecular Genetics and Metabolism, 2007, 90, 113-121.	0.5	14
136	Multiple ganglion cysts (â€ˆcystic ganglionosisâ€™™): an unusual presentation in a child. Scandinavian Journal of Rheumatology, 2007, 36, 145-148.	0.6	19
137	A phase II/III clinical study of enzyme replacement therapy with idursulfase in mucopolysaccharidosis II (Hunter syndrome). Genetics in Medicine, 2006, 8, 465-473.	1.1	499
138	Recognition of Smith-Lemli-Opitz syndrome (RSH) in the fetus: Utility of ultrasonography and biochemical analysis in pregnancies with low maternal serum estriol. American Journal of Medical Genetics, Part A, 2005, 138A, 56-60.	0.7	19
139	Extensive primary cutaneous herpes simplex virus type 1 infection in an infant following acute rotavirus gastroenteritis. European Journal of Pediatrics, 2005, 164, 175-176.	1.3	0
140	Autoantibodies against bactericidal/permeabilityâ€ˆincreasing protein (BPIâ€™ANCA) in cystic fibrosis patients treated with azithromycin. Clinical and Experimental Medicine, 2005, 5, 80-85.	1.9	20
141	Splicing mutation in the fibrillin-1 gene associated with neonatal Marfan syndrome and severe pulmonary emphysema with tracheobronchomalacia. Pediatric Pulmonology, 2005, 39, 374-378.	1.0	19
142	No live individual homozygous for a novel endoglin mutation was found in a consanguineous Arab family with hereditary haemorrhagic telangiectasia. Journal of Medical Genetics, 2004, 41, e119-e119.	1.5	21
143	Pulmonary manifestations and function tests in children genetically diagnosed with FMF. Pediatric Pulmonology, 2003, 35, 452-455.	1.0	11
144	The differential contribution of MEFV mutant alleles to the clinical profile of familial Mediterranean fever. European Journal of Human Genetics, 2002, 10, 145-149.	1.4	116

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145	Familial mediterranean fever: The segregation of four different mutations in 13 individuals from one inbred family: Genotype-phenotype correlation and intrafamilial variability. American Journal of Medical Genetics Part A, 2002, 109, 198-201.	2.4	15
146	Crouzon syndrome: Association with absent pulmonary valve syndrome and severe tracheobronchomalacia. Pediatric Pulmonology, 2002, 34, 478-481.	1.0	12
147	The musculoskeletal manifestations of familial Mediterranean fever in children genetically diagnosed with the disease. Arthritis and Rheumatism, 2001, 44, 1416-1419.	6.7	78
148	Familial Mediterranean fever: prevalence, penetrance and genetic drift. European Journal of Human Genetics, 2001, 9, 634-637.	1.4	146
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