

Andreas Tzschach

List of Publications by Year in descending order

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134
papers

7,974
citations

66343

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136
docs citations

136
times ranked

13112
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#	ARTICLE	IF	CITATIONS
1	Range of genetic mutations associated with severe non-syndromic sporadic intellectual disability: an exome sequencing study. <i>Lancet, The</i> , 2012, 380, 1674-1682.	13.7	940
2	Deep sequencing reveals 50 novel genes for recessive cognitive disorders. <i>Nature</i> , 2011, 478, 57-63.	27.8	805
3	Mutations in the JARID1C Gene, Which Is Involved in Transcriptional Regulation and Chromatin Remodeling, Cause X-Linked Mental Retardation. <i>American Journal of Human Genetics</i> , 2005, 76, 227-236.	6.2	349
4	<i>STXBP1</i> encephalopathy. <i>Neurology</i> , 2016, 86, 954-962.	1.1	264
5	Mutations in NSUN2 Cause Autosomal- Recessive Intellectual Disability. <i>American Journal of Human Genetics</i> , 2012, 90, 847-855.	6.2	243
6	X-exome sequencing of 405 unresolved families identifies seven novel intellectual disability genes. <i>Molecular Psychiatry</i> , 2016, 21, 133-148.	7.9	243
7	Mutations in the Small GTPase Gene RAB39B Are Responsible for X-linked Mental Retardation Associated with Autism, Epilepsy, and Macrocephaly. <i>American Journal of Human Genetics</i> , 2010, 86, 185-195.	6.2	220
8	Cranioectodermal Dysplasia, Sensenbrenner Syndrome, Is a Ciliopathy Caused by Mutations in the IFT122 Gene. <i>American Journal of Human Genetics</i> , 2010, 86, 949-956.	6.2	180
9	A novel X-linked recessive mental retardation syndrome comprising macrocephaly and ciliary dysfunction is allelic to oral-“facial”-digital type I syndrome. <i>Human Genetics</i> , 2006, 120, 171-178.	3.8	166
10	Mutations in the polyglutamine binding protein 1 gene cause X-linked mental retardation. <i>Nature Genetics</i> , 2003, 35, 313-315.	21.4	139
11	A Defect in the Ionotropic Glutamate Receptor 6 Gene (GRIK2) Is Associated with Autosomal Recessive Mental Retardation. <i>American Journal of Human Genetics</i> , 2007, 81, 792-798.	6.2	137
12	Mutation Update for Kabuki Syndrome Genes <i>KMT2D</i> and <i>KDM6A</i> and Further Delineation of X-Linked Kabuki Syndrome Subtype 2. <i>Human Mutation</i> , 2016, 37, 847-864.	2.5	134
13	A balanced chromosomal translocation disrupting <i>ARHGEF9</i> is associated with epilepsy, anxiety, aggression, and mental retardation. <i>Human Mutation</i> , 2009, 30, 61-68.	2.5	131
14	Genetics of intellectual disability in consanguineous families. <i>Molecular Psychiatry</i> , 2019, 24, 1027-1039.	7.9	131
15	Structural variation in Xq28: MECP2 duplications in 1% of patients with unexplained XLMR and in 2% of male patients with severe encephalopathy. <i>European Journal of Human Genetics</i> , 2009, 17, 444-453.	2.8	130
16	A Defect in the TUSC3 Gene Is Associated with Autosomal Recessive Mental Retardation. <i>American Journal of Human Genetics</i> , 2008, 82, 1158-1164.	6.2	127
17	Novel JARID1C/SMCX mutations in patients with X-linked mental retardation. <i>Human Mutation</i> , 2006, 27, 389-389.	2.5	120
18	Identification of Mutations in TRAPPC9, which Encodes the NIK- and IKK- β -Binding Protein, in Nonsyndromic Autosomal-Recessive Mental Retardation. <i>American Journal of Human Genetics</i> , 2009, 85, 909-915.	6.2	120

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19	Mapping translocation breakpoints by next-generation sequencing. <i>Genome Research</i> , 2008, 18, 1143-1149.	5.5	118
20	Mutations in autism susceptibility candidate 2 (AUTS2) in patients with mental retardation. <i>Human Genetics</i> , 2007, 121, 501-509.	3.8	116
21	Next-generation sequencing in X-linked intellectual disability. <i>European Journal of Human Genetics</i> , 2015, 23, 1513-1518.	2.8	112
22	High frequency of submicroscopic genomic aberrations detected by tiling path array comparative genome hybridisation in patients with isolated congenital heart disease. <i>Journal of Medical Genetics</i> , 2008, 45, 704-709.	3.2	110
23	Clinical Presentation of a Complex Neurodevelopmental Disorder Caused by Mutations in ADNP. <i>Biological Psychiatry</i> , 2019, 85, 287-297.	1.3	108
24	Mutation frequencies of X-linked mental retardation genes in families from the EuroMRX consortium. <i>Human Mutation</i> , 2007, 28, 207-208.	2.5	103
25	Homozygosity mapping in consanguineous families reveals extreme heterogeneity of non-syndromic autosomal recessive mental retardation and identifies 8 novel gene loci. <i>Human Genetics</i> , 2007, 121, 43-48.	3.8	92
26	ST3GAL3 Mutations Impair the Development of Higher Cognitive Functions. <i>American Journal of Human Genetics</i> , 2011, 89, 407-414.	6.2	89
27	A Noncoding, Regulatory Mutation Implicates HCFC1 in Nonsyndromic Intellectual Disability. <i>American Journal of Human Genetics</i> , 2012, 91, 694-702.	6.2	89
28	A clinical and molecular genetic study of 112 Iranian families with primary microcephaly. <i>Journal of Medical Genetics</i> , 2010, 47, 823-828.	3.2	87
29	Multiple giant cell lesions in patients with Noonan syndrome and cardio-facio-cutaneous syndrome. <i>European Journal of Human Genetics</i> , 2009, 17, 420-425.	2.8	79
30	Mutation of the conserved polyadenosine RNA binding protein, ZC3H14/dNab2, impairs neural function in <i>Drosophila</i> and humans. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 12390-12395.	7.1	77
31	Mutations in the Alpha 1,2-Mannosidase Gene, MAN1B1, Cause Autosomal-Recessive Intellectual Disability. <i>American Journal of Human Genetics</i> , 2011, 89, 176-182.	6.2	73
32	SNP array-based homozygosity mapping reveals MCPH1 deletion in family with autosomal recessive mental retardation and mild microcephaly. <i>Human Genetics</i> , 2006, 118, 708-715.	3.8	67
33	Chromosome deletions in 13q33-q34: Report of four patients and review of the literature. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 337-342.	1.2	63
34	Breakpoint analysis of balanced chromosome rearrangements by next-generation paired-end sequencing. <i>European Journal of Human Genetics</i> , 2010, 18, 539-543.	2.8	61
35	MCT8 mutation analysis and identification of the first female with Allan-Herndon-Dudley syndrome due to loss of MCT8 expression. <i>European Journal of Human Genetics</i> , 2008, 16, 1029-1037.	2.8	56
36	Characterization of a 5.3 Mb deletion in 15q14 by comparative genomic hybridization using a whole genome tiling path-BAC array in a girl with heart defect, cleft palate, and developmental delay. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 172-178.	1.2	55

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37	Mutations of the aminoacyl-tRNA-synthetases SARS and WARS2 are implicated in the etiology of autosomal recessive intellectual disability. <i>Human Mutation</i> , 2017, 38, 621-636.	2.5	54
38	Next generation sequencing in a family with autosomal recessive Kahrizi syndrome (OMIM 612713) reveals a homozygous frameshift mutation in SRD5A3. <i>European Journal of Human Genetics</i> , 2011, 19, 115-117.	2.8	52
39	Identification of a nonsense mutation in the very low-density lipoprotein receptor gene (VLDLR) in an Iranian family with dysequilibrium syndrome. <i>European Journal of Human Genetics</i> , 2008, 16, 270-273.	2.8	50
40	Mutation screening in 86 known X-linked mental retardation genes by droplet-based multiplex PCR and massive parallel sequencing. <i>The HUGO Journal</i> , 2009, 3, 41-49.	4.1	48
41	Characterisation of de novo MAPK10/JNK3 truncation mutations associated with cognitive disorders in two unrelated patients. <i>Human Genetics</i> , 2013, 132, 461-471.	3.8	48
42	The molecular and phenotypic spectrum of IQSEC2-related epilepsy. <i>Epilepsia</i> , 2016, 57, 1858-1869.	5.1	46
43	Novel missense mutations in the ubiquitination-related gene UBE2A cause a recognizable X-linked mental retardation syndrome. <i>Clinical Genetics</i> , 2010, 77, 541-551.	2.0	45
44	Autosomal recessive mental retardation: homozygosity mapping identifies 27 single linkage intervals, at least 14 novel loci and several mutation hotspots. <i>Human Genetics</i> , 2011, 129, 141-148.	3.8	45
45	Expanded mutational spectrum in Cohen syndrome, tissue expression, and transcript variants of COH1. <i>Human Mutation</i> , 2009, 30, E404-E420.	2.5	44
46	Musculoskeletal Disease in MDA5-Related Type I Interferonopathy: A Mendelian Mimic of Jaccoud's Arthropathy. <i>Arthritis and Rheumatology</i> , 2017, 69, 2081-2091.	5.6	44
47	A novel nonsense mutation in TUSC3 is responsible for non-syndromic autosomal recessive mental retardation in a consanguineous Iranian family. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 1976-1980.	1.2	43
48	KCNC1-related disorders: new de novo variants expand the phenotypic spectrum. <i>Annals of Clinical and Translational Neurology</i> , 2019, 6, 1319-1326.	3.7	43
49	Variants in CUL4B are Associated with Cerebral Malformations. <i>Human Mutation</i> , 2015, 36, 106-117.	2.5	37
50	Diagnostic value of partial exome sequencing in developmental disorders. <i>PLoS ONE</i> , 2018, 13, e0201041.	2.5	36
51	Characterization of interstitial Xp duplications in two families by tiling path array CGH. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 197-203.	1.2	35
52	A distinctive gene expression fingerprint in mentally retarded male patients reflects disease-causing defects in the histone demethylase KDM5C. <i>PathoGenetics</i> , 2010, 3, 2.	5.7	35
53	Identification of a novel CDKL5 exon and pathogenic mutations in patients with severe mental retardation, early-onset seizures and Rett-like features. <i>Neurogenetics</i> , 2011, 12, 165-167.	1.4	34
54	De novo partial deletion in GRID2 presenting with complicated spastic paraplegia. <i>Muscle and Nerve</i> , 2014, 49, 289-292.	2.2	33

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55	<i>PIK3R1</i> mutations in <i>SHORT</i> syndrome. <i>Clinical Genetics</i> , 2014, 86, 292-294.	2.0	31
56	Epilepsy is not a mandatory feature of <i>STXBP1</i> associated ataxia-tremor-retardation syndrome. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 661-665.	1.6	30
57	Radiation-Induced Late Effects in Two Affected Individuals of the Lilo Radiation Accident. <i>Radiation Research</i> , 2007, 167, 615-623.	1.5	29
58	Cohen syndrome diagnosis using whole genome arrays. <i>Journal of Medical Genetics</i> , 2011, 48, 136-140.	3.2	29
59	Novel <i>WDR35</i> mutations in patients with cranioectodermal dysplasia (Sensenbrenner) Tj ETQq1 1 0.784314,rgBT /Overlock 10	2.8	28
60	Mutations in the histamine <i>N</i> -methyltransferase gene, <i>HNMT</i> , are associated with nonsyndromic autosomal recessive intellectual disability. <i>Human Molecular Genetics</i> , 2015, 24, 5697-5710.	2.9	27
61	Transitioning the Molecular Tumor Board from Proof of Concept to Clinical Routine: A German Single-Center Analysis. <i>Cancers</i> , 2021, 13, 1151.	3.7	27
62	Mutation screening of <i>ASMT</i> , the last enzyme of the melatonin pathway, in a large sample of patients with Intellectual Disability. <i>BMC Medical Genetics</i> , 2011, 12, 17.	2.1	25
63	Clinical and functional characterisation of the combined respiratory chain defect in two sisters due to autosomal recessive mutations in <i>MTFMT</i> . <i>Mitochondrion</i> , 2013, 13, 743-748.	3.4	25
64	Craniosynostosis in a patient with 2q37.3 deletion 5q34 duplication: Association of extra copy of <i>MSX2</i> with craniosynostosis. <i>American Journal of Medical Genetics, Part A</i> , 2009, 149A, 1544-1549.	1.2	24
65	Isolated <i>NIBPL</i> missense mutations that cause Cornelia de Lange syndrome alter <i>MAU2</i> interaction. <i>European Journal of Human Genetics</i> , 2012, 20, 271-276.	2.8	24
66	Kohlschütter-Tarnz Syndrome: Mutations in <i>ROGDI</i> and Evidence of Genetic Heterogeneity. <i>Human Mutation</i> , 2013, 34, 296-300.	2.5	24
67	Novel <i>SLC9A6</i> mutations in two families with Christianson syndrome. <i>Clinical Genetics</i> , 2013, 83, 596-597.	2.0	24
68	The power of the Mediator complex-Expanding the genetic architecture and phenotypic spectrum of <i>MED12</i> -related disorders. <i>Clinical Genetics</i> , 2018, 94, 450-456.	2.0	24
69	Chromosome aberrations involving 10q22: report of three overlapping interstitial deletions and a balanced translocation disrupting <i>C10orf11</i> . <i>European Journal of Human Genetics</i> , 2010, 18, 291-295.	2.8	22
70	X-linked intellectual disability type Nascimento is a clinically distinct, probably underdiagnosed entity. <i>Orphanet Journal of Rare Diseases</i> , 2013, 8, 146.	2.7	22
71	Hybridisation-based resequencing of 17 X-linked intellectual disability genes in 135 patients reveals novel mutations in <i>ATRX</i> , <i>SLC6A8</i> and <i>PQBP1</i> . <i>European Journal of Human Genetics</i> , 2011, 19, 717-720.	2.8	21
72	<i>Sema3a</i> plays a role in the pathogenesis of <i>CHARGE</i> syndrome. <i>Human Molecular Genetics</i> , 2018, 27, 1343-1352.	2.9	20

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73	X-linked mental retardation: a comprehensive molecular screen of 47 candidate genes from a 7.4 Mb interval in Xp11. <i>European Journal of Human Genetics</i> , 2007, 15, 68-75.	2.8	19
74	Xq22.3â€“q23 deletion including <i>ACSL4</i> in a patient with intellectual disability. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 860-864.	1.2	19
75	Next-generation panel sequencing identifies NF1 germline mutations in three patients with pheochromocytoma but no clinical diagnosis of neurofibromatosis type 1. <i>European Journal of Endocrinology</i> , 2018, 178, K1-K9.	3.7	19
76	Czech dysplasia: Report of a large family and further delineation of the phenotype. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 1859-1864.	1.2	18
77	Characterization of an interstitial 4q32 deletion in a patient with mental retardation and a complex chromosome rearrangement. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 1008-1012.	1.2	18
78	Floatingâ€“Harbor syndrome: <i>SRCAP</i> mutations are not restricted to exon 34. <i>Clinical Genetics</i> , 2014, 85, 498-499.	2.0	18
79	Variable clinical phenotype in two siblings with Aicardi-GoutiÃˆres syndrome type 6 and a novel mutation in the ADAR gene. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 186-189.	1.6	18
80	BOD1 Is Required for Cognitive Function in Humans and <i>Drosophila</i> . <i>PLoS Genetics</i> , 2016, 12, e1006022.	3.5	18
81	An autosomal recessive syndrome of severe mental retardation, cataract, coloboma and kyphosis maps to the pericentromeric region of chromosome 4. <i>European Journal of Human Genetics</i> , 2009, 17, 125-128.	2.8	17
82	Heterotaxy and cardiac defect in a girl with chromosome translocation t(X;1)(q26;p13.1) and involvement of ZIC3. <i>European Journal of Human Genetics</i> , 2006, 14, 1317-1320.	2.8	16
83	Novel <i>GD11</i> mutation in a large family with nonsyndromic Xâ€“linked intellectual disability. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 3067-3070.	1.2	16
84	A novel <i>ALDH5A1</i> mutation is associated with succinic semialdehyde dehydrogenase deficiency and severe intellectual disability in an Iranian family. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 1915-1922.	1.2	16
85	Deletions in 14q24.1q24.3 are associated with congenital heart defects, brachydactyly, and mild intellectual disability. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 620-626.	1.2	16
86	Characterization of a 16 Mb interstitial chromosome 7q21 deletion by tiling path array CGH. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 333-337.	1.2	15
87	Ready to clone: CNV detection and breakpoint fine-mapping in breast and ovarian cancer susceptibility genes by high-resolution array CGH. <i>Breast Cancer Research and Treatment</i> , 2016, 159, 585-590.	2.5	15
88	A mosaic maternal splice donor mutation in the EHMT1 gene leads to aberrant transcripts and to Kleefstra syndrome in the offspring. <i>European Journal of Human Genetics</i> , 2013, 21, 887-890.	2.8	14
89	Congenital CLN disease in two siblings. <i>Wiener Medizinische Wochenschrift</i> , 2015, 165, 210-213.	1.1	14
90	Interstitial deletion 2p11.2â€“p12: Report of a patient with mental retardation and review of the literature. <i>American Journal of Medical Genetics, Part A</i> , 2009, 149A, 242-245.	1.2	13

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91	Christianson syndrome in a patient with an interstitial Xq26.3 deletion. American Journal of Medical Genetics, Part A, 2011, 155, 2771-2774.	1.2	13
92	Platelet defects in congenital variant of Rett syndrome patients with FOXP1 mutations or reduced expression due to a position effect at 14q12. European Journal of Human Genetics, 2013, 21, 1349-1355.	2.8	13
93	Molecular cytogenetic analysis of a de novo interstitial deletion of 5q23.3q31.2 and its phenotypic consequences. American Journal of Medical Genetics, Part A, 2006, 140A, 496-502.	1.2	12
94	Pierpont syndrome: report of a new patient. Clinical Dysmorphology, 2017, 26, 205-208.	0.3	12
95	Skewed X-chromosome inactivation in a family with DLG3-associated X-linked intellectual disability. American Journal of Medical Genetics, Part A, 2017, 173, 2545-2550.	1.2	12
96	ABSENCE OF Yq MICRODELETIONS IN INFERTILE MEN. Archives of Andrology, 2001, 47, 167-171.	1.0	11
97	Interstitial 3p25.3-p26.1 deletion in a patient with intellectual disability. American Journal of Medical Genetics, Part A, 2012, 158A, 2587-2590.	1.2	11
98	Angelman syndrome and severe infections in a patient with de novo 15q11.2-q13.1 deletion and maternally inherited 2q21.3 microdeletion. Gene, 2013, 512, 453-455.	2.2	11
99	Tentative clinical diagnosis of Lujan-Fryns syndrome? A conglomeration of different genetic entities?. American Journal of Medical Genetics, Part A, 2016, 170, 94-102.	1.2	11
100	Novel PRPS1 gain-of-function mutation in a patient with congenital hyperuricemia and facial anomalies. , 2017, 173, 2736-2742.		11
101	Marfan Syndrome Caused by Disruption of the FBN1 Gene due to A Reciprocal Chromosome Translocation. Genes, 2021, 12, 1836.	2.4	11
102	Autosomal dominant inheritance in a large family with focal facial dermal dysplasia (Brauer-Setleis) Tj ETQq0 0 0 rgBT /Overlock 10 Tf	1.2	10
103	Interstitial 9q34.1-q34.13 deletion in a patient with severe intellectual disability, hydrocephalus, and cleft lip/palate. American Journal of Medical Genetics, Part A, 2012, 158A, 1709-1712.	1.2	10
104	A Novel SLC6A8 Mutation in a Large Family with X-Linked Intellectual Disability: Clinical and Proton Magnetic Resonance Spectroscopy Data of Both Hemizygous Males and Heterozygous Females. JIMD Reports, 2013, 13, 91-99.	1.5	10
105	Novel truncating PPM1D mutation in a patient with intellectual disability. European Journal of Medical Genetics, 2019, 62, 70-72.	1.3	10
106	Skeletal abnormalities are common features in Aym-Grripp syndrome. Clinical Genetics, 2020, 97, 362-369.	2.0	10
107	New Cav1.2 Channelopathy with High-Functioning Autism, Affective Disorder, Severe Dental Enamel Defects, a Short QT Interval, and a Novel CACNA1C Loss-of-Function Mutation. International Journal of Molecular Sciences, 2020, 21, 8611.	4.1	10
108	A Novel MGST2 Non-Synonymous Mutation in a Chinese Pedigree with Psoriasis Vulgaris. Journal of Investigative Dermatology, 2006, 126, 1003-1005.	0.7	9

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109	Molecular cytogenetic analysis of a de novo interstitial chromosome 10q22 deletion. American Journal of Medical Genetics, Part A, 2006, 140A, 1108-1110.	1.2	9
110	Mirror-Image Asymmetry in Monozygotic Twins with Kabuki Syndrome. Molecular Syndromology, 2012, 3, 94-97.	0.8	9
111	Parental Origin of de novo Cytogenetically Balanced Reciprocal Non-Robertsonian Translocations. Cytogenetic and Genome Research, 2012, 136, 242-245.	1.1	9
112	A newly recognized autosomal recessive syndrome affecting neurologic function and vision. American Journal of Medical Genetics, Part A, 2013, 161, 1207-1213.	1.2	9
113	Molecular characterization of a balanced chromosome translocation in psoriasis vulgaris. Clinical Genetics, 2005, 69, 189-193.	2.0	8
114	Novel VPS33B mutation in a patient with autosomal recessive keratodermaâ€¢chthysisâ€¢deafness syndrome. American Journal of Medical Genetics, Part A, 2018, 176, 2862-2866.	1.2	8
115	Interstitial 1q23.3q24.1 deletion in a patient with renal malformation, congenital heart disease, and mild intellectual disability. American Journal of Medical Genetics, Part A, 2016, 170, 2394-2399.	1.2	7
116	PUF60-SCRIB fusion transcript in a patient with 8q24.3 microdeletion and atypical Verheij syndrome. European Journal of Medical Genetics, 2019, 62, 103587.	1.3	7
117	Blepharophimosis-Ptosis-Epicanthus Inversus Syndrome in a Girl with Chromosome Translocation t(2;3)(q33;q23). Ophthalmic Genetics, 2008, 29, 37-40.	1.2	6
118	Interstitial 1p32.1p32.3 deletion in a patient with multiple congenital anomalies. American Journal of Medical Genetics, Part A, 2015, 167, 2406-2410.	1.2	6
119	11q14.1â€¢1q22.1 deletion in a 1â€¢yearâ€¢old male with minor dysmorphic features. American Journal of Medical Genetics, Part A, 2010, 152A, 2651-2655.	1.2	5
120	Novel ADAMTSL2-mutations in a patient with geleophysic dysplasia type I. Clinical Dysmorphology, 2016, 25, 106-109.	0.3	5
121	Alopecia mental retardation syndrome: clinical and molecular characterization of four patients. British Journal of Dermatology, 2008, 159, ???-???.	1.5	4
122	Interstitial duplication of chromosome region 1q25.1q25.3: Report of a patient with mild cognitive deficits, tall stature and facial dysmorphisms. American Journal of Medical Genetics, Part A, 2015, 167, 653-656.	1.2	4
123	Posterior amorphous corneal dystrophy in a patient with 12q21.33 deletion. Ophthalmic Genetics, 2018, 39, 645-647.	1.2	4
124	<i>PIGN</i> encephalopathy: Characterizing the epileptology. Epilepsia, 2022, 63, 974-991.	5.1	4
125	Molecular breakpoint analysis and relevance of variable mosaicism in a woman with short stature, primary amenorrhea, unilateral gonadoblastoma, and a 46,X,del(Y)(q11)/45,X karyotype. American Journal of Medical Genetics Part A, 2002, 112, 51-55.	2.4	3
126	Brachyphalangy, polydactyly and tibial aplasia/hypoplasia syndrome (OMIM 609945): case report and review of the literature. European Journal of Pediatrics, 2010, 169, 1535-1539.	2.7	3

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127	Hypergonadotropic hypogonadism in a patient with inv ins (2;4). Journal of Developmental and Physical Disabilities, 2009, 32, 226-230.	3.6	2
128	12q24.33 deletion: Report of a patient with intellectual disability and review of the literature. American Journal of Medical Genetics, Part A, 2013, 161, 1409-1413.	1.2	2
129	The Importance of Extended Analysis Using Current Molecular Genetic Methods Based on the Example of a Cohort of 228 Patients with Hereditary Breast and Ovarian Cancer Syndrome. Genes, 2021, 12, 1483.	2.4	1
130	Genetik der nichtsyndromalen geistigen Behinderung. Medizinische Genetik, 2009, 21, 231-236.	0.2	0
131	Reply to 5q35 duplication and Hunterâ€™McAlpine syndrome: Missing the link. American Journal of Medical Genetics, Part A, 2010, 152A, 804-804.	1.2	0
132	Chromosome aberration associated with hippocampal impairment. Psychiatry Research - Neuroimaging, 2016, 254, 1-2.	1.8	0
133	Abstract LB-044: Germline mutations in patients with hereditary breast and ovarian cancer establish ERCC2 as a cancer susceptibility gene. , 2015, , .		0
134	Abstract 3391: Next generation sequencing paves the way for personalized medicine in pheochromocytoma and paraganglioma patients and their families. , 2017, , .		0