John M Opitz

List of Publications by Citations

214 papers	9,083 citations	47 h-index	90 g-index
238 ext. papers	10,060 ext. citations	4.9 avg, IF	5.51 L-index

#	Paper	IF	Citations
214	Disruptions of topological chromatin domains cause pathogenic rewiring of gene-enhancer interactions. *Cell*, **2015**, 161, 1012-1025	56.2	1207
213	A NEWLY RECOGNIZED SYNDROME OF MULTIPLE CONGENITAL ANOMALIES. *Journal of Pediatrics*, **1964**, 64, 210-7	3.6	579
212	Resynthesizing evolutionary and developmental biology. *Developmental Biology*, **1996**, 173, 357-72	3.1	502
211	Noonan syndrome: a review. *American Journal of Medical Genetics Part A*, **1985**, 21, 493-506		367
210	Opitz G/BBB syndrome, a defect of midline development, is due to mutations in a new RING finger gene on Xp22. *Nature Genetics*, **1997**, 17, 285-91	36.3	296
209	Evaluation of mental retardation: recommendations of a Consensus Conference: American College of Medical Genetics. *American Journal of Medical Genetics Part A*, **1997**, 72, 468-77		272
208	CNS anomalies and the midline as a "developmental field". *American Journal of Medical Genetics Part A*, **1982**, 12, 443-55		240
207	Mutations in mitochondrial histidyl tRNA synthetase HARS2 cause ovarian dysgenesis and sensorineural hearing loss of Perrault syndrome. *Proceedings of the National Academy of Sciences of the United States of America*, **2011**, 108, 6543-8	11.5	200
206	Smith-Lemli-Opitz syndrome-type II: multiple congenital anomalies with male pseudohermaphroditism and frequent early lethality. *American Journal of Medical Genetics Part A*, **1987**, 26, 45-57		200
205	Germline mutations in DIS3L2 cause the Perlman syndrome of overgrowth and Wilms tumor susceptibility. *Nature Genetics*, **2012**, 44, 277-84	36.3	173
204	A recurrent mutation in MED12 leading to R961W causes Opitz-Kaveggia syndrome. *Nature Genetics*, **2007**, 39, 451-3	36.3	157
203	The DiGeorge anomaly as a developmental field defect. *American Journal of Medical Genetics Part A* , **1986**, 2, 113-27		154
202	The Brachmann-de Lange syndrome. *American Journal of Medical Genetics Part A*, **1985**, 22, 89-102		141
201	Isochromosome 12p mosaicism (Pallister mosaic aneuploidy or Pallister-Killian syndrome): report of 11 cases. *American Journal of Medical Genetics Part A*, **1987**, 27, 257-74		123
200	Studies of malformation syndromes of man 33: the FG syndrome. An X-linked recessive syndrome of multiple congenital anomalies and mental retardation. *European Journal of Pediatrics*, **1974**, 117, 1-18	4.1	122
199	Rett syndrome at an institution for the developmentally disabled. *American Journal of Medical Genetics Part A*, **1986**, 1, 85-97		120
198	Fibular a/hypoplasia: review and documentation of the fibular developmental field. *American Journal of Medical Genetics Part A*, **1986**, 2, 215-38		117

197	The developmental field concept. *American Journal of Medical Genetics Part A*, **1985**, 21, 1-11		108
196	Errors of morphogenesis and developmental field theory **1998**, 76, 291-296		105
195	Defects of blastogenesis. *American Journal of Medical Genetics Part A*, **2002**, 115, 269-86		95
194	Borderline normal intelligence in the Smith-Lemli-Opitz (RSH) syndrome. *American Journal of Medical Genetics Part A*, **1980**, 5, 137-43		91
193	The syndromes of Sotos and Weaver: reports and review. *American Journal of Medical Genetics Part A*, **1998**, 79, 294-304		90
192	RSH/SLO ("Smith-Lemli-Opitz") syndrome: historical, genetic, and developmental considerations. *American Journal of Medical Genetics Part A*, **1994**, 50, 344-6		79
191	The Poland syndrome-clinical and genealogical data, dermatoglyphic analysis, and incidence. *Human Heredity*, **1973**, 23, 97-104	1.1	76
190	Amnion rupture sequence in previable fetuses. *American Journal of Medical Genetics Part A*, **1988**, 31, 63-73		74
189	G syndrome (hypertelorism with esophageal abnormality and hypospadias, or hypospadias-dysphagia, or "Opitz-Frias" or "Opitz-G" syndrome)--perspective in 1987 and bibliography. *American Journal of Medical Genetics Part A*, **1987**, 28, 275-85		73
188	Ovine ceroid-lipofuscinosis II: Pathologic changes interpreted in light of biochemical observations. *American Journal of Medical Genetics Part A*, **1988**, 5, 159-70		70
187	Implications of malformations not due to amniotic bands in the amniotic band sequence. *American Journal of Medical Genetics Part A*, **1986**, 24, 691-700		69
186	A new autosomal dominant acrofacial dysostosis syndrome. *American Journal of Medical Genetics Part A*, **1986**, 2, 143-50		68
185	Opitz G/BBB syndrome: clinical comparisons of families linked to Xp22 and 22q, and a review of the literature. *American Journal of Medical Genetics Part A*, **1996**, 62, 305-17		66
184	Dubowitz syndrome: review of 141 cases including 36 previously unreported patients. *American Journal of Medical Genetics Part A*, **1996**, 63, 277-89		64
183	The Perrault syndrome: autosomal recessive ovarian dysgenesis with facultative, non-sex-limited sensorineural deafness. *American Journal of Medical Genetics Part A*, **1979**, 4, 239-46		64
182	Differential diagnosis of Nager acrofacial dysostosis syndrome: report of four patients with Nager syndrome and discussion of other related syndromes. *American Journal of Medical Genetics Part A*, **1983**, 14, 209-24		61
181	Autosomal recessive syndrome of cerebellar ataxia and hypogonadotropic hypogonadism. *Clinical Genetics*, **1975**, 7, 426-34	4	59
180	Identification and characterization of a missense mutation in the -linked □acetylglucosamine (-GlcNAc) transferase gene that segregates with X-linked intellectual disability. *Journal of Biological Chemistry*, **2017**, 292, 8948-8963	5.4	58

179	Meier-Gorlin syndrome: report of eight additional cases and review. *American Journal of Medical Genetics Part A*, **2001**, 102, 115-24		58
178	Studies of malformation syndromes of man XXIX: the Wiedemann-Beckwith syndrome. Clinical, genetic and pathogenetic studies of 12 cases. *European Journal of Nuclear Medicine and Molecular Imaging*, **1976**, 123, 139-66		57
177	The Perlman syndrome: familial renal dysplasia with Wilms tumor, fetal gigantism and multiple congenital anomalies. *American Journal of Medical Genetics Part A*, **1984**, 19, 195-207		56
176	Polytopic anomalies with agenesis of the lower vertebral column. *American Journal of Medical Genetics Part A*, **1999**, 87, 99-114		54
175	FG syndrome update 1988: note of 5 new patients and bibliography. *American Journal of Medical Genetics Part A*, **1988**, 30, 309-28		53
174	A novel germline PIGA mutation in Ferro-Cerebro-Cutaneous syndrome: a neurodegenerative X-linked epileptic encephalopathy with systemic iron-overload. *American Journal of Medical Genetics, Part A*, **2014**, 164A, 17-28	2.5	51
173	I-Cell disease, mucolipidosis II. *European Journal of Pediatrics*, **1973**, 114, 259-292	4.1	51
172	Further delineation of the C (trigonocephaly) syndrome. *American Journal of Medical Genetics Part A*, **1981**, 9, 147-63		50
171	Severe end of Opitz trigonocephaly (C) syndrome or new syndrome?. *American Journal of Medical Genetics Part A*, **1999**, 85, 438-46		48
170	Dermatoglyphic traits as possible markers of developmental processes in humans. *American Journal of Medical Genetics Part A*, **1987**, 26, 111-22		46
169	Mutations in CD96, a member of the immunoglobulin superfamily, cause a form of the C (Opitz trigonocephaly) syndrome. *American Journal of Human Genetics*, **2007**, 81, 835-41	11	44
168	A new X-linked mental retardation syndrome. *American Journal of Medical Genetics Part A*, **1985**, 21, 697-705		43
167	Prenatal diagnosis of perinatally lethal osteogenesis imperfecta. *American Journal of Medical Genetics Part A*, **1983**, 14, 353-9		39
166	Tandem dup (1p) within the short arm of chromosome 1 in a child with ambiguous genitalia and multiple congenital anomalies. *American Journal of Medical Genetics Part A*, **1984**, 17, 723-30		38
165	Evidence for the □midline□ hypothesis in associated defects of laterality formation and multiple midline anomalies. *American Journal of Medical Genetics Part A*, **2001**, 101, 382-387		36
164	Editorial comment on the paper by de la Monte and Hutchins on familial polyasplenia. *American Journal of Medical Genetics Part A*, **1985**, 21, 175-176		36
163	CHOLESTEROL AND DEVELOPMENT: THE RSH ("SMITH-LEMLI-OPITZ") SYNDROME AND RELATED CONDITIONS. *Fetal and Pediatric Pathology*, **2002**, 21, 153-181		34
162	Vaginal atresia (von Mayer-Rokitansky-K□ster or MRK anomaly) in hereditary renal adysplasia (HRA). *American Journal of Medical Genetics Part A*, **1987**, 26, 873-6		34

161	Studies of malformation syndromes of humans XXXIIIC: the FG syndrome - further studies on three affected individuals from the FG family. *American Journal of Medical Genetics Part A*, **1982**, 12, 147-54		34
160	Syndrome of mental retardation, seizures, hypotonic cerebral palsy and megalocorneae, recessively inherited. *European Journal of Pediatrics*, **1975**, 120, 1-18	4.1	33
159	Autosomal recessive syndrome of pseudogliomantous blindness, osteoporosis and mild mental retardation. *Clinical Genetics*, **1976**, 9, 324-32	4	32
158	Specific congenital heart defects in RSH/Smith-Lemli-Opitz syndrome: postulated involvement of the sonic hedgehog pathway in syndromes with postaxial polydactyly or heterotaxia. *Birth Defects Research Part A: Clinical and Molecular Teratology*, **2003**, 67, 149-53		32
157	Klippel-Feil anomaly with Sprengel anomaly, omovertebral bone, thumb abnormalities, and flexion-crease changes: Novel association or syndrome?. *American Journal of Medical Genetics Part A*, **2001**, 101, 158-162		32
156	The Noonan syndrome. *American Journal of Medical Genetics Part A*, **1985**, 21, 515-8		32
155	Studies of malformation syndromes of man XXXIX: a craniosynostosis-craniofacial dysostosis syndrome with mental retardation and other malformations: "craniofacial dyssynostosis". *European Journal of Nuclear Medicine and Molecular Imaging*, **1976**, 123, 15-28		31
154	Multiple meningiomas, craniofacial hyperostosis and retinal abnormalities in Proteus syndrome. *American Journal of Medical Genetics Part A*, **2000**, 93, 234-40		30
153	Brief historical note: the concept of "gonadal dysgenesis". *American Journal of Medical Genetics Part A*, **1979**, 4, 333-43		30
152	Two sporadic cases of amelia/phocomelia with similar phenotype: rare and unusually symmetrical form of FFU dysostosis or separate entity?. *American Journal of Medical Genetics Part A*, **1986**, 2, 239-45		29
151	Smith-Lemli-Opitz (RSH) syndrome bibliography. *American Journal of Medical Genetics Part A*, **1987**, 28, 745-50		28
150	Simpson-Golabi-Behmel syndrome: follow-up of the Michigan family. *American Journal of Medical Genetics Part A*, **1988**, 30, 301-8		28
149	Autosomal dominant and sporadic radio-ulnar synostosis **1997**, 68, 127-134		27
148	Studies of malformation syndromes of man XXXXIA: anatomical studies in the Hanhart syndrome--a pathogenetic hypothesis. *European Journal of Nuclear Medicine and Molecular Imaging*, **1976**, 122, 1-17		27
147	An Xq22.3 duplication detected by comparative genomic hybridization microarray (Array-CGH) defines a new locus (FGS5) for FG syndrome. *American Journal of Medical Genetics, Part A*, **2005**, 139, 221-6	2.5	26
146	Cholesterol and development: the RSH ("Smith-Lemli-Opitz") syndrome and related conditions. *Fetal and Pediatric Pathology*, **2002**, 21, 153-81		26
145	Smith-Lemli-Opitz (RSH) syndrome bibliography: 1964-1993. *American Journal of Medical Genetics Part A*, **1994**, 50, 339-43		26
144	An X-linked recessive basal ganglia disorder with mental retardation. *American Journal of Medical Genetics Part A*, **1985**, 21, 681-9		26

143	Cervical ribs are more prevalent in stillborn fetuses than in live-born infants and are strongly associated with fetal aneuploidy. *Pediatric and Developmental Pathology*, **2011**, 14, 431-7	2.2	25
142	Meckel on developmental pathology. *American Journal of Medical Genetics, Part A*, **2006**, 140, 115-28	2.5	25
141	A De Novo Nonsense Mutation in MAGEL2 in a Patient Initially Diagnosed as Opitz-C: Similarities Between Schaaf-Yang and Opitz-C Syndromes. *Scientific Reports*, **2017**, 7, 44138	4.9	24
140	Comments on biological asymmetry. *American Journal of Medical Genetics Part A*, **2001**, 101, 359-369		24
139	Proximal femoral focal deficiency (PFFD) and fibular A/hypoplasia (FA/H): a model of a developmental field defect. *American Journal of Medical Genetics Part A*, **1995**, 55, 427-32		24
138	Nager "syndrome" versus "anomaly" and its nosology with the postaxial acrofacial dysostosis syndrome of Genß and Wiedemann. *American Journal of Medical Genetics Part A*, **1987**, 27, 959-63		24
137	Perrault syndrome in sisters. *American Journal of Medical Genetics Part A*, **1985**, 22, 629-31		23
136	Hypertrichosis lanuginosa in a mother and son. *Clinical Genetics*, **1976**, 10, 303-6	4	22
135	FG syndrome: linkage analysis in two families supporting a new gene localization at Xp22.3 [FGS3]. *American Journal of Medical Genetics Part A*, **2002**, 112, 6-11		22
134	Brief clinical report: unilateral partial tibia defect with preaxial polydactyly, general micromelia, and trigonomacrocephaly with a note on "developmental resistance". *American Journal of Medical Genetics Part A*, **1983**, 14, 467-71		22
133	A fetus with upper limb amelia, "caudal regression" and Dandy-Walker defect with an insulin-dependent diabetic mother. *European Journal of Pediatrics*, **1980**, 134, 139-43	4.1	22
132	An evolutionary and developmental biology approach to gastroschisis. *Birth Defects Research*, **2019**, 111, 294-311	2.9	22
131	Kniest dysplasia: radiologic, histopathological, and scanning electronmicroscopic findings. *American Journal of Medical Genetics Part A*, **1996**, 63, 34-45		21
130	Ulnar ray a/hypoplasia: evidence for a developmental field defect on the basis of genetic heterogeneity. Report of three Brazilian families. *American Journal of Medical Genetics Part A*, **1986**, 2, 195-206		21
129	The Genß-Wiedemann syndrome, an acrofacial dysostosis--further observation. *American Journal of Medical Genetics Part A*, **1987**, 27, 971-5		21
128	Studies of malformation syndromes of man XXXXIIB: mother and son affected with the ulnar-mammary syndrome type Pallister. *European Journal of Nuclear Medicine and Molecular Imaging*, **1976**, 123, 225-35		21
127	Down syndrome: comments and reflections on the 50th anniversary of Lejeune's discovery. *American Journal of Medical Genetics, Part A*, **2009**, 149A, 2647-54	2.5	20
126	Blaschkolinear malformation syndrome in complex trisomy-7 mosaicism. *American Journal of Medical Genetics Part A*, **1999**, 87, 375-83		20

125	Severe case of Al Awadi/Raas-Rothschild syndrome or new, possibly autosomal recessive facio-skeleto-genital syndrome. *American Journal of Medical Genetics Part A*, **1995**, 56, 168-72		20
124	Developmental terms -- some proposals: first report of an international working group. *American Journal of Medical Genetics Part A*, **1979**, 3, 297-302		20
123	Heart development: An introduction. *American Journal of Medical Genetics Part A*, **2000**, 97, 238-247		19
122	Microcephaly, lymphedema, and chorioretinal dysplasia: report of two additional cases. *American Journal of Medical Genetics Part A*, **1994**, 53, 99-101		18
121	Sedaghatian congenital lethal metaphyseal chondrodysplasia--observations in a second Iranian family and histopathological studies. *American Journal of Medical Genetics Part A*, **1987**, 26, 583-90		18
120	Developmental field theory and observations□ccidental progress?. *American Journal of Medical Genetics Part A*, **1986**, 25, 1-8		17
119	Lincoln vs. Douglas again; comments on the papers by Curry et al, Greenberg et al, and Belmont et al. *American Journal of Medical Genetics Part A*, **1987**, 26, 69-71		17
118	Hutterite cerebro-osteo-nephrodysplasia: autosomal recessive trait in a Lehrerleut Hutterite family from Montana. *American Journal of Medical Genetics Part A*, **1985**, 22, 521-9		17
117	Mortality and pathological findings in C (Opitz trigonocephaly) syndrome. *Fetal and Pediatric Pathology*, **2006**, 25, 211-31	1.7	16
116	Megacystis-microcolon-intestinal hypoperistalsis syndrome and aganglionosis in trisomy 18. *American Journal of Medical Genetics Part A*, **2001**, 102, 293-296		16
115	Behavior phenotype of FG syndrome: cognition, personality, and behavior in eleven affected boys. *American Journal of Medical Genetics Part A*, **2000**, 97, 112-8		16
114	Noonan's syndrome in the male. *Journal of Pediatrics*, **1965**, 67, 936	3.6	16
113	Prenatal death in Smith-Lemli-Opitz/RSH syndrome. *American Journal of Medical Genetics, Part A*, **2005**, 138, 61-5	2.5	15
112	Generalized gangliosidosis type II (juvenile GM1 gangliosidosis). A pathological, histochemical and ultrastructural study. *European Journal of Pediatrics*, **1975**, 120, 151-80	4.1	15
111	An unusual dysplasia-malformation-cancer syndrome in two patients. *American Journal of Medical Genetics Part A*, **1978**, 1, 279-89		15
110	Smith-Lemli-Opitz syndrome in Japan. *American Journal of Medical Genetics Part A*, **1998**, 75, 118-119		14
109	The evaluation of infants with the Zellweger (cerebro-hepato-renal) syndrome. *Clinical Genetics*, **1975**, 7, 413-6	4	14
108	Twenty-seven-year follow-up in the Wolf-Hirschhorn syndrome. *American Journal of Medical Genetics Part A*, **1995**, 55, 459-61		14

107	Virilism as a late manifestation in the Bardet-Biedl syndrome. *American Journal of Medical Genetics Part A*, **1980**, 7, 279-92		14
106	Delayed Mutation as a Cause of Genetic Disease in Man: Achondroplasia and the Wiedemann-Beckwith Syndrome **1977**, 65-79		14
105	The neurofaciodigitorenal (NFDR) syndrome. *American Journal of Medical Genetics Part A*, **1982**, 11, 329-36		13
104	The FG syndromes (Online Mendelian Inheritance in Man 305450): perspective in 2008. *Advances in Pediatrics*, **2008**, 55, 123-70	2.2	12
103	Prenatal death in Fraser syndrome. *Fetal and Pediatric Pathology*, **2005**, 24, 223-38	1.7	12
102	Disequilibrium syndrome in Montana Hutterites. *American Journal of Medical Genetics Part A*, **1985**, 22, 567-9		12
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100	Invited editorial comment: further reflections on gastroschisis. *American Journal of Medical Genetics, Part C: Seminars in Medical Genetics*, **2008**, 148C, 192-8	3.1	11
99	The genetics of quantifiable homeostasis: I. The general issues. *American Journal of Medical Genetics Part A*, **1986**, 24, 159-69		11
98	Clinicopathological conference: a 29-yr-old man with recurrent episodes of fever, abdominal pain, and vomiting. *American Journal of Medical Genetics Part A*, **1984**, 18, 249-64		11
97	Heterogeneity of nonlethal severe short-limbed dwarfism. *Journal of Pediatrics*, **1977**, 91, 918-23	3.6	11
96	Screening of CD96 and ASXL1 in 11 patients with Opitz C or Bohring-Opitz syndromes. *American Journal of Medical Genetics, Part A*, **2016**, 170A, 24-31	2.5	11
95	Craniocervical anomalies in Dubowitz syndrome. Three cases and a literature review. *Pediatric Neurosurgery*, **2003**, 38, 238-43	0.9	10
94	Developmental field theory and the molecular analysis of morphogenesis: A comment on Dr. Slavkin's observations. *American Journal of Medical Genetics Part A*, **1993**, 47, 687-688		10
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92	The Montana Fetal Genetic Pathology Program and a review of prenatal death in humans. *American Journal of Medical Genetics Part A*, **1987**, 3, 93-112		10
91	Autopsy findings in a stillborn female infant with the Osebold-Remondini syndrome. *American Journal of Medical Genetics Part A*, **1985**, 22, 811-9		10
90	ADAM "sequence" part II: hypothesis and speculation. *American Journal of Medical Genetics, Part A*, **2015**, 167A, 478-503	2.5	9

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88	Reflections on the pathogenesis of Down syndrome. *American Journal of Medical Genetics Part A*, **1990**, 7, 38-51		9
87	Sixty years of X-linked mental retardation: a historical footnote. *American Journal of Medical Genetics Part A*, **2000**, 97, 228-33		9
86	Book reviewISCN 2013: An International System for Human Cytogenetic Nomenclature (2013). Shaffer LG, McGowan-Jordan J, Schmid M, editors. Published in collaboration with Cytogenetic and Genome Research by Karger, Basel. VI and 140 pp. plus fold-out, 11 Fig. 4 tables. **2013**, 161, 2108-2109		8
85	2011 William Allan Award: development and evolution. *American Journal of Human Genetics*, **2012**, 90, 392-404	11	8
84	Opitz trigonocephaly syndrome presenting with sudden unexplained death in the operating room: a case report. *Journal of Medical Case Reports*, **2011**, 5, 222	1.2	8
83	Another "new" form, the palagonia type of acrofacial dysostosis in a Sicilian family. *American Journal of Medical Genetics Part A*, **1997**, 69, 388-94		8
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81	Development: clinical and evolutionary considerations. *American Journal of Medical Genetics, Part A*, **2007**, 143A, 2853-61	2.5	8
80	Heterogeneity and minor anomalies. *American Journal of Medical Genetics Part A*, **2000**, 91, 254-5		8
79	Sudden death in childhood in a case of the G syndrome. *American Journal of Medical Genetics Part A*, **1987**, 28, 293-6		8
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77	Lethal Forms of Chondrodysplastic Dwarfism. *Pediatrics*, **1974**, 53, 76-85	7.4	7
76	Annals of morphology THEODOR BOVERI (1862–1915) To commemorate the centenary of his death and contributions to the Sutton–Boveri hypothesis **2016**, 170, 2803-2829		6
75	MENDEL: Morphologist and Mathematician Founder of Genetics - To Begin a Celebration of the 2015 Sesquicentennial of Mendel's Presentation in 1865 of his Versuche über Pflanzenhybriden. *Molecular Genetics & Genomic Medicine*, **2015**, 3, 1-7	2.3	6
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70	Mandibulofacial dysostosis or bilateral hemifacial microsomia with hearing loss, telecanthus, tetramelic postaxial hexadactyly, congenital hypotonia and lymphedema with joint hypermobility, and pigmentary dysplasia: a new syndrome?. *American Journal of Medical Genetics Part A*, **1989**, 33, 433-5		6
69	Johann Friedrich Meckel the Younger (1781-1833). *American Journal of Medical Genetics Part A*, **1984**, 18, 571-86		6
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67	A De Novo FOXP1 Truncating Mutation in a Patient Originally Diagnosed as C Syndrome. *Scientific Reports*, **2018**, 8, 694	4.9	5
66	Phenotypes, pleiotropy, and phylogeny. *American Journal of Medical Genetics, Part C: Seminars in Medical Genetics*, **2017**, 175, 329-340	3.1	5
65	Sudden infant death "syndrome"-Insights and future directions from a Utah population database analysis. *American Journal of Medical Genetics, Part A*, **2017**, 173, 177-182	2.5	5
64	Genitourinary anomalies of pediatric FG syndrome. *Journal of Urology*, **2007**, 178, 656-9	2.5	5
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62	Human anotocephaly (aprosopus, acrania-synotia) in the Vilnius anatomical collection. *American Journal of Medical Genetics Part A*, **2001**, 101, 163-71		4
61	Behold the CHILD. *American Journal of Medical Genetics Part A*, **2000**, 90, 336-8		4
60	Heterogeneity of cardio-facio-cutaneous syndrome. *American Journal of Medical Genetics Part A*, **2000**, 95, 144		4
59	Encomium: Robert J. Gorlin. *American Journal of Medical Genetics Part A*, **1993**, 46, 316-316		4
58	Renal failure with hypercalcemia, renal stones, multiple pathologic fractures, and growth failure. *American Journal of Medical Genetics Part A*, **1983**, 14, 169-79		4
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56	Genetic caring. the professionalization of genetic services in the USA. *American Journal of Medical Genetics Part A*, **1979**, 3, 1-5		4
55	Serendipity or prepared mind? Recollections of the KOP translocation (1967) and of one form of Perrault syndrome. *American Journal of Medical Genetics, Part C: Seminars in Medical Genetics*, **2014**, 166C, 387-96	3.1	3
54	The RSH/"Smith-Lemli-Opitz" syndrome: historical footnote. *American Journal of Medical Genetics, Part C: Seminars in Medical Genetics*, **2012**, 160C, 242-9	3.1	3

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51	David Klein: an appreciation. *American Journal of Medical Genetics Part A*, **1990**, 37, 320-4		3
50	Bibliography on X-linked mental retardation, the fragile X and related subjects IV (1988). *American Journal of Medical Genetics Part A*, **1988**, 30, 31-60		3
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46	Arthur G. Steinberg: an appreciation. *American Journal of Medical Genetics Part A*, **1995**, 59, 245-9		2
45	Biographical note--Laurence H. Snyder. *American Journal of Medical Genetics Part A*, **1981**, 8, 447-8		2
44	Comments on biological asymmetry **2001**, 101, 359		2
43	Arno G. Motulsky, 1923-2018, Luck and Service. *American Journal of Medical Genetics, Part A*, **2018**, 176, 1285-1288	2.5	1
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