H Eugene Hoyme

List of Publications by Year
in descending order

149 papers	10,945 citations	38742 50 h-index	31849 101 g-index
158 all docs	158 docs citations	158 times ranked	7180 citing authors

#	Article	IF	Citations
1	Characteristic physical traits of firstâ€grade children in the United States with fetal alcohol spectrum disorders (<scp>FASD</scp>) and associated alcohol and drug exposures. American Journal of Medical Genetics, Part A, 2022, 188, 2019-2035.	1.2	3
2	The influence of maternal weight and alcohol exposure on infant physical characteristics and neurodevelopmental outcomes. Current Research in Toxicology, 2022, 3, 100076.	2.7	2
3	The role of novel <scp>COQ8B</scp> mutations in glomerulopathy and related kidney defects. American Journal of Medical Genetics, Part A, 2021, 185, 60-67.	1.2	3
4	The prevalence, child characteristics, and maternal risk factors for the continuum of fetal alcohol spectrum disorders: A sixth population-based study in the same South African community. Drug and Alcohol Dependence, 2021, 218, 108408.	3.2	32
5	Evolution of the Physical Phenotype of Fetal Alcohol Spectrum Disorders from Childhood through Adolescence. Alcoholism: Clinical and Experimental Research, 2021, 45, 395-408.	2.4	20
6	Gestational age and birth growth parameters as early predictors of fetal alcohol spectrum disorders. Alcoholism: Clinical and Experimental Research, 2021, 45, 1624-1638.	2.4	4
7	Estimating the community prevalence, child traits, and maternal risk factors of fetal alcohol spectrum disorders (FASD) from a random sample of school children. Drug and Alcohol Dependence, 2021, 227, 108918.	3.2	18
8	The challenges and pitfalls of fetal alcohol spectrum disorders prevalence studies. Alcoholism: Clinical and Experimental Research, 2021, 45, 2468-2470.	2.4	1
9	Perspectives of Pediatric Providers Regarding Clinical Use of Pharmacogenetics. South Dakota Medicine: the Journal of the South Dakota State Medical Association, 2021, 74, 294-301.	0.2	1
10	Ocular measurements in fetal alcohol spectrum disorders. American Journal of Medical Genetics, Part A, 2020, 182, 2243-2252.	1.2	7
11	Fetal Alcohol Spectrum Disorders in a Southeastern County of the United States: Child Characteristics and Maternal Risk Traits. Alcoholism: Clinical and Experimental Research, 2020, 44, 939-959.	2.4	30
12	Fetal Alcohol Spectrum Disorders in a Midwestern City: Child Characteristics, Maternal Risk Traits, and Prevalence. Alcoholism: Clinical and Experimental Research, 2020, 44, 919-938.	2.4	35
13	Fetal Alcohol Spectrum Disorders in a Rocky Mountain Region City: Child Characteristics, Maternal Risk Traits, and Prevalence. Alcoholism: Clinical and Experimental Research, 2020, 44, 900-918.	2.4	35
14	Early-Life Predictors of Fetal Alcohol Spectrum Disorders. Pediatrics, 2019, 144, .	2.1	26
15	Prevalence of Fetal Alcohol Spectrum Disorders in 4 US Communities. JAMA - Journal of the American Medical Association, 2018, 319, 474.	7.4	562
16	Fetal Alcohol Spectrum Disorders: Diagnostic Considerations for Children with a History of Trauma. , 2018, , 93-122.		0
17	Efficacy of Maternal Choline Supplementation During Pregnancy in Mitigating Adverse Effects of Prenatal Alcohol Exposure on Growth and Cognitive Function: A Randomized, Doubleâ€Blind, Placeboâ€Controlled Clinical Trial. Alcoholism: Clinical and Experimental Research, 2018, 42, 1327-1341.	2.4	109
18	Evaluation of a Formal Pediatric Faculty Mentorship Program. South Dakota Medicine: the Journal of the South Dakota State Medical Association, 2018, 71, 256-262.	0.2	2

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19	Heavy Prenatal Alcohol Exposure is Related to Smaller Corpus Callosum in Newborn <scp>MRI</scp> Scans. Alcoholism: Clinical and Experimental Research, 2017, 41, 965-975.	2.4	62
20	A modified Timeline Followback assessment to capture alcohol exposure in pregnant women: Application in the Safe Passage Study. Alcohol, 2017, 62, 17-27.	1.7	28
21	Who is most affected by prenatal alcohol exposure: Boys or girls?. Drug and Alcohol Dependence, 2017, 177, 258-267.	3.2	39
22	Drinking and smoking patterns during pregnancy: Development of group-based trajectories in the Safe Passage Study. Alcohol, 2017, 62, 49-60.	1.7	45
23	Computer-Aided Recognition of Facial Attributes for Fetal Alcohol Spectrum Disorders. Pediatrics, 2017, 140, e20162028.	2.1	44
24	Facial Curvature Detects and Explicates Ethnic Differences in Effects of Prenatal Alcohol Exposure. Alcoholism: Clinical and Experimental Research, 2017, 41, 1471-1483.	2.4	28
25	Commentary on the decision of the American Board of Medical Genetics and Genomics to create a 24-month specialty of Laboratory Genetics and Genomics. Genetics in Medicine, 2017, 19, 294-296.	2.4	2
26	Replication of High Fetal Alcohol Spectrum Disorders Prevalence Rates, Child Characteristics, and Maternal Risk Factors in a Second Sample of Rural Communities in South Africa. International Journal of Environmental Research and Public Health, 2017, 14, 522.	2.6	43
27	Fetal alcohol spectrum disorders and assessment of maxillary and mandibular arc measurements. American Journal of Medical Genetics, Part A, 2016, 170, 1763-1771.	1.2	8
28	Breastfeeding and maternal alcohol use: Prevalence and effects on child outcomes and fetal alcohol spectrum disorders. Reproductive Toxicology, 2016, 63, 13-21.	2.9	52
29	The continuum of fetal alcohol spectrum disorders in a community in South Africa: Prevalence and characteristics in a fifth sample. Drug and Alcohol Dependence, 2016, 168, 274-286.	3.2	41
30	A Decision Tree to Identify Children Affected by Prenatal Alcohol Exposure. Journal of Pediatrics, 2016, 177, 121-127.e1.	1.8	35
31	Updated Clinical Guidelines for Diagnosing Fetal Alcohol Spectrum Disorders. Pediatrics, 2016, 138, .	2.1	561
32	Alcohol-Related Neurobehavioral Disabilities: Need for Further Definition and Common Terminology. Pediatrics, 2016, 138, .	2.1	2
33	The continuum of fetal alcohol spectrum disorders in four rural communities in south africa: Prevalence and characteristics. Drug and Alcohol Dependence, 2016, 159, 207-218.	3.2	66
34	Maternal nutritional status as a contributing factor for the risk of fetal alcohol spectrum disorders. Reproductive Toxicology, 2016, 59, 101-108.	2.9	48
35	Patient and Physician Perceptions of Genetic Testing in Primary Care. South Dakota Medicine: the Journal of the South Dakota State Medical Association, 2016, 69, 487-493.	0.2	3
36	A South African mixed race lip/philtrum guide for diagnosis of fetal alcohol spectrum disorders. American Journal of Medical Genetics, Part A, 2015, 167, 752-755.	1.2	24

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37	Prevalence and characteristics of fetal alcohol syndrome and partial fetal alcohol syndrome in a Rocky Mountain Region City. Drug and Alcohol Dependence, 2015, 155, 118-127.	3.2	64
38	Epigenetics in Pediatrics. Pediatrics in Review, 2015, 36, 14-21.	0.4	6
39	Upper Gastrointestinal Tract. , 2015, , 697-730.		0
40	Epigenetics in Pediatrics. Pediatrics in Review, 2015, 36, 14-21.	0.4	2
41	Maternal risk factors for fetal alcohol spectrum disorders in a province in Italy. Drug and Alcohol Dependence, 2014, 145, 201-208.	3.2	25
42	Prevalence and Characteristics of Fetal Alcohol Spectrum Disorders. Pediatrics, 2014, 134, 855-866.	2.1	474
43	Dietary intake, nutrition, and fetal alcohol spectrum disorders in the Western Cape Province of South Africa. Reproductive Toxicology, 2014, 46, 31-39.	2.9	67
44	Spontaneous Abortion and Intrauterine Fetal Death. , 2014, , 497-508.		0
45	Maternal alcohol consumption producing fetal alcohol spectrum disorders (FASD): Quantity, frequency, and timing of drinking. Drug and Alcohol Dependence, 2013, 133, 502-512.	3.2	182
46	Facial Dysmorphism Across the Fetal Alcohol Spectrum. Pediatrics, 2013, 131, e779-e788.	2.1	114
47	Approaching the Prevalence of the Full Spectrum of Fetal Alcohol Spectrum Disorders in a <scp>S</scp>outh <scp>A</scp>frican Populationâ€Based Study. Alcoholism: Clinical and Experimental Research, 2013, 37, 818-830.	2.4	198
48	Maternal Factors Predicting Cognitive and Behavioral Characteristics of Children with Fetal Alcohol Spectrum Disorders. Journal of Developmental and Behavioral Pediatrics, 2013, 34, 314-325.	1.1	63
49	Core Concepts: Chromosome Aneuploidies. NeoReviews, 2012, 13, e30-e39.	0.8	3
50	Relation Over Time Between Facial Measurements and Cognitive Outcomes in Fetal Alcoholâ€Exposed Children. Alcoholism: Clinical and Experimental Research, 2012, 36, 1634-1646.	2.4	22
51	Marked variability in the radiographic features of cartilageâ€hair hypoplasia: Case report and review of the literature. American Journal of Medical Genetics, Part A, 2012, 158A, 2911-2916.	1.2	17
52	Advances in Whole-Genome Genetic Testing: From Chromosomes to Microarrays. Current Problems in Pediatric and Adolescent Health Care, 2012, 42, 47-73.	1.7	18
53	Adaptive behaviour in children and adolescents with foetal alcohol spectrum disorders: a comparison with specific learning disability and typical development. European Child and Adolescent Psychiatry, 2012, 21, 221-231.	4.7	47
54	Maternal risk factors predicting child physical characteristics and dysmorphology in fetal alcohol syndrome and partial fetal alcohol syndrome. Drug and Alcohol Dependence, 2011, 119, 18-27.	3.2	47

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55	Prevalence of Children with Severe Fetal Alcohol Spectrum Disorders in Communities Near Rome, Italy: New Estimated Rates Are Higher than Previous Estimates. International Journal of Environmental Research and Public Health, 2011, 8, 2331-2351.	2.6	111
56	Impaired Delay and Trace Eyeblink Conditioning in School-Age Children With Fetal Alcohol Syndrome. Alcoholism: Clinical and Experimental Research, 2011, 35, 250-264.	2.4	84
57	Risk factors for behavioural problems in foetal alcohol spectrum disorders. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 1481-1488.	1.5	38
58	Ectopia lentis as the presenting and primary feature in Marfan syndrome. American Journal of Medical Genetics, Part A, 2011, 155, 2661-2668.	1.2	38
59	Interactive Feature Visualization and Detection for 3D Face Classification. International Journal of Cognitive Informatics and Natural Intelligence, 2011, 5, 1-16.	0.4	2
60	Population Differences in Dysmorphic Features Among Children With Fetal Alcohol Spectrum Disorders. Journal of Developmental and Behavioral Pediatrics, 2010, 31, 304-316.	1.1	23
61	Fetal alcohol spectrum disorders: Extending the range of structural defects. American Journal of Medical Genetics, Part A, 2010, 152A, 2731-2735.	1.2	65
62	Prenatal alcohol exposure alters the patterns of facial asymmetry. Alcohol, 2010, 44, 649-657.	1.7	90
63	An fMRI Study of Number Processing in Children With Fetal Alcohol Syndrome. Alcoholism: Clinical and Experimental Research, 2010, 34, 1450-1464.	2.4	54
64	Interactive feature visualization and detection for 3D face classification. , 2010, , .		0
65	Design, Construction, and Testing of a Stereo-Photogrammetric Tool for the Diagnosis of Fetal Alcohol Syndrome in Infants. IEEE Transactions on Medical Imaging, 2009, 28, 1448-1458.	8.9	19
66	Elements of morphology: Standard terminology for the head and face. American Journal of Medical Genetics, Part A, 2009, 149A, 6-28.	1.2	145
67	Prevalence and epidemiologic characteristics of FASD from various research methods with an emphasis on recent inâ€school studies. Developmental Disabilities Research Reviews, 2009, 15, 176-192.	2.9	775
68	Developmental pathogenesis of short palpebral fissure length in children with fetal alcohol syndrome. Birth Defects Research Part A: Clinical and Molecular Teratology, 2009, 85, 695-699.	1.6	10
69	The Effects of Fetal Alcohol Syndrome on Response Execution and Inhibition: An Eventâ€Related Potential Study. Alcoholism: Clinical and Experimental Research, 2009, 33, 1994-2004.	2.4	49
70	Progressive and symmetric supraorbital hyperostosis with bony and soft tissue overgrowth in an Ethiopian female: A newly recognized overgrowth syndrome?. American Journal of Medical Genetics, Part A, 2008, 146A, 543-547.	1.2	1
71	Impaired Eyeblink Conditioning in Children With Fetal Alcohol Syndrome. Alcoholism: Clinical and Experimental Research, 2008, 32, 365-372.	2.4	160
72	Further Delineation of Deletion 1p36 Syndrome in 60 Patients: A Recognizable Phenotype and Common Cause of Developmental Delay and Mental Retardation. Pediatrics, 2008, 121, 404-410.	2.1	233

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73	The epidemiology of fetal alcohol syndrome and partial FAS in a South African community. Drug and Alcohol Dependence, 2007, 88, 259-271.	3.2	257
74	Relationship between dysmorphic features and general cognitive function in children with fetal alcohol spectrum disorders. American Journal of Medical Genetics, Part A, 2007, 143A, 2916-2923.	1.2	30
75	22q13.3 deletion syndrome: A recognizable malformation syndrome associated with marked speech and language delay. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2007, 145C, 393-398.	1.6	78
76	Clinical delineation of fetal alcohol spectrum disorders (FASD) in Italian children: Comparison and contrast with other racial/ethnic groups and implications for diagnosis and prevention. Neuroscience and Biobehavioral Reviews, 2007, 31, 270-277.	6.1	26
77	Unique Facial Features Distinguish Fetal Alcohol Syndrome Patients and Controls in Diverse Ethnic Populations. Alcoholism: Clinical and Experimental Research, 2007, 31, 1707-1713.	2.4	76
78	Fetal alcohol spectrum disorders: A practical clinical approach to diagnosis. Neuroscience and Biobehavioral Reviews, 2007, 31, 230-238.	6.1	125
79	Epidemiology of FASD in a Province in Italy: Prevalence and Characteristics of Children in a Random Sample of Schools. Alcoholism: Clinical and Experimental Research, 2006, 30, 1562-1575.	2.4	147
80	Comparison of Motor Delays in Young Children With Fetal Alcohol Syndrome to Those With Prenatal Alcohol Exposure and With No Prenatal Alcohol Exposure. Alcoholism: Clinical and Experimental Research, 2006, 30, 2037-2045.	2.4	107
81	Triplication of 8p22-8p23 in a patient with features similar to Kabuki syndrome. American Journal of Medical Genetics, Part A, 2006, 140A, 170-173.	1.2	15
82	Fetal alcohol spectrum disorders in Finland: Clinical delineation of 77 older children and adolescents. American Journal of Medical Genetics, Part A, 2006, 140A, 137-143.	1.2	96
83	Malignancy in chromosome 22q11.2 deletion syndrome (DiGeorge syndrome/velocardiofacial) Tj ETQq1 1 0.784314 rgBT /Overlock 1	1.2	59
84	Nablus mask-like facial syndrome is caused by a microdeletion of 8q detected by array-based comparative genomic hybridization. American Journal of Medical Genetics, Part A, 2006, 140A, 1267-1273.	1.2	31
85	Unilateral Aquagenic Wrinkling of the Palms Associated With Aspirin Intake. Archives of Dermatology, 2006, 142, 1661-2.	1.4	46
86	Systemic Hyalinosis: A Distinctive Early Childhoodâ€“Onset Disorder Characterized by Mutations in the Anthrax Toxin Receptor 2 Gene (ANTRX2). Pediatrics, 2006, 118, e1485-e1492.	2.1	26
87	Fetal alcohol syndrome epidemiology in a South African community: a second study of a very high prevalence area.. Journal of Studies on Alcohol and Drugs, 2005, 66, 593-604.	2.3	176
88	Congenital anomaly rate in offspring of mothers with diabetes treated with insulin lispro during pregnancy. Diabetic Medicine, 2005, 22, 803-807.	2.3	99
89	Marinesco-SjÃ¶gren syndrome in a male with mild dysmorphism. , 2005, 133A, 197-201.		19
90	Autosomal dominant microtia and ocular coloboma: New syndrome or an extension of the oculo-auriculo-vertebral spectrum?. American Journal of Medical Genetics, Part A, 2005, 134A, 359-362.	1.2	21

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91	Terminal deletion of 6p results in a recognizable phenotype. American Journal of Medical Genetics, Part A, 2005, 136A, 162-168.	1.2	49
92	Vascular-type disruptive defects in fetuses with homozygous Î±-thalassemia: report of two cases and review of the literature. Prenatal Diagnosis, 2005, 25, 1088-1096.	2.3	22
93	A Practical Clinical Approach to Diagnosis of Fetal Alcohol Spectrum Disorders: Clarification of the 1996 Institute of Medicine Criteria. Pediatrics, 2005, 115, 39-47.	2.1	779
94	Molecular and Clinical Analyses of Greig Cephalopolysyndactyly and Pallister-Hall Syndromes: Robust Phenotype Prediction from the Type and Position of GLI3 Mutations. American Journal of Human Genetics, 2005, 76, 609-622.	6.2	248
95	The clinical picture of the BÃ¶rjesonâ€“Forssmanâ€“Lehmann syndrome in males and heterozygous females with PHF6 mutations. Clinical Genetics, 2004, 65, 226-232.	2.0	63
96	Neuâ€“Laxova syndrome: Detailed prenatal diagnostic and postâ€mortem findings and literature review. American Journal of Medical Genetics Part A, 2004, 125A, 240-249.	2.4	53
97	Alcohol consumption and other maternal risk factors for fetal alcohol syndrome among three distinct samples of women before, during, and after pregnancy: The risk is relative. American Journal of Medical Genetics Part A, 2004, 127C, 10-20.	2.4	91
98	Rare Autosomal Recessive Cardiac Valvular Form of Ehlers-Danlos Syndrome Results from Mutations in the COL1A2 Gene That Activate the Nonsense-Mediated RNA Decay Pathway. American Journal of Human Genetics, 2004, 74, 917-930.	6.2	147
99	Terminal 22q Deletion Syndrome: A Newly Recognized Cause of Speech and Language Disability in the Autism Spectrum. Pediatrics, 2004, 114, 451-457.	2.1	148
100	Morphogenesis: Re: Clinical, natural history, and imaging information on patients included in reports. American Journal of Medical Genetics Part A, 2003, 119A, 93-93.	2.4	5
101	Athabascan brainstem dysgenesis syndrome. American Journal of Medical Genetics Part A, 2003, 120A, 169-173.	2.4	66
102	Methotrexate/misoprostol embryopathy: Report of four cases resulting from failed medical abortion. American Journal of Medical Genetics Part A, 2003, 123A, 72-78.	2.4	48
103	Follow-up of children of depressed mothers exposed or not exposed to antidepressant drugs during pregnancy. Journal of Pediatrics, 2003, 142, 402-408.	1.8	374
104	The practice of clinical genetics: A survey of practitioners. Genetics in Medicine, 2002, 4, 142-149.	2.4	22
105	Early Neonatal Diagnosis of Long-Chain 3-Hydroxyacyl Coenzyme A Dehydrogenase and Mitochondrial Trifunctional Protein Deficiencies. Molecular Genetics and Metabolism, 2002, 75, 120-127.	1.1	37
106	Mutations in PHF6 are associated with BÃ¶rjesonâ€“ForssmanÂâ€“Lehmann syndrome. Nature Genetics, 2002, 32, 661-665.	21.4	192
107	Reply to correspondence from Winter. American Journal of Medical Genetics Part A, 2002, 111, 458-458.	2.4	0
108	Pharmacogenomics: the future of drug therapy. Clinical Genetics, 2002, 62, 257-264.	2.0	31

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109	Limb/pelvis hypoplasia/aplasia with skull defect (Schinzel phocomelia): Distinctive features and prenatal detection. American Journal of Medical Genetics Part A, 2001, 103, 295-301.	2.4	22
110	A Clinical Genetics and Dysmorphology Approach to Growth Deficiency. Pediatric Annals, 2000, 29, 549-555.	0.8	2
111	Blepharophimosis: A causally heterogeneous malformation frequently associated with developmental disabilities. , 1998, 75, 52-54.		26
112	Isolated hemihyperplasia (hemihypertrophy): Report of a prospective multicenter study of the incidence of neoplasia and review. , 1998, 79, 274-278.		165
113	Six patients with oral-facial-digital syndrome IV: The case for heterogeneity. American Journal of Medical Genetics Part A, 1997, 69, 250-260.	2.4	23
114	Six patients with oralâ€facialâ€digital syndrome IV: The case for heterogeneity. American Journal of Medical Genetics Part A, 1997, 69, 250-260.	2.4	1
115	Sudden death in Williams syndrome: Report of ten cases. Journal of Pediatrics, 1996, 129, 926-931.	1.8	167
116	von Voss-Cherstvoy syndrome: A variable perinatally lethal syndrome of multiple congenital anomalies. American Journal of Medical Genetics Part A, 1994, 52, 272-278.	2.4	22
117	Minor anomalies: diagnostic clues to aberrant human morphogenesis. Contemporary Issues in Genetics and Evolution, 1994, , 309-317.	0.9	9
118	Toluene Embryopathy: Delineation of the Phenotype and Comparison With Fetal Alcohol Syndrome. Pediatrics, 1994, 93, 211-215.	2.1	92
119	Congenital scalp defects and vitreoretinal degeneration: Redefining the Knobloch syndrome. American Journal of Medical Genetics Part A, 1993, 46, 203-208.	2.4	49
120	Further delineation of the epidermal nevus syndrome: Two cases with new findings and literature review. American Journal of Medical Genetics Part A, 1993, 47, 24-30.	2.4	74
121	Minor anomalies: Diagnostic clues to aberrant human morphogenesis. Genetica, 1993, 89, 307-315.	1.1	20
122	Urethral obstruction sequence and lower limb deficiency: Evidence for the vascular disruption hypothesis. Journal of Pediatrics, 1993, 123, 398-405.	1.8	10
123	Cardiac Teratogenicity of Dichloroethylene in a Chick Model. Pediatric Research, 1992, 32, 23-26.	2.3	27
124	Teratology in Pediatric Practice. Pediatric Clinics of North America, 1992, 39, 111-134.	1.8	9
125	Possible common pathogenetic mechanisms for Poland sequence and Adams-Oliver syndrome: An additional clinical observation. American Journal of Medical Genetics Part A, 1992, 42, 398-399.	2.4	24
126	Possible common pathogenetic mechanisms for Poland sequence and Adams-Oliver syndrome. American Journal of Medical Genetics Part A, 1991, 38, 69-73.	2.4	54

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127	Fetal Vascular Disruption With Prenatal Exposure to Cocaine or Methamphetamine. Pediatrics, 1991, 87, 417-418.	2.1	0
128	Identification of a balanced translocation carrier by spouse's low maternal serum alpha fetoprotein associated with an aneuploid fetus.. Journal of Medical Genetics, 1990, 27, 728-728.	3.2	0
129	Prenatal Cocaine Exposure and Fetal Vascular Disruption. Pediatrics, 1990, 85, 743-747.	2.1	160
130	The pathogenesis of sirenomelia: An editorial comment. Teratology, 1988, 38, 485-486.	1.6	15
131	Minor Malformations. American Journal of Diseases of Children, 1987, 141, 947.	0.5	13
132	Genetic Disorders and the Fetus: Diagnosis, Prevention, and Treatment. JAMA Pediatrics, 1987, 141, 1062.	3.0	0
133	Congenital Oral Tumor Associated with Neurofibromatosis Detected by Prenatal Ultrasound. Clinical Pediatrics, 1987, 26, 372-374.	0.8	22
134	Vascular pathogenesis of unilateral craniofacial defects. Journal of Pediatrics, 1987, 111, 236-239.	1.8	51
135	Autosomal dominant ectrodactyly and absence of long bones of upper or lower limbs: Further clinical delineation. Journal of Pediatrics, 1987, 111, 538-543.	1.8	24
136	Bronchoscopic evaluation of airway obstruction in campomelic dysplasia. Pediatric Pulmonology, 1987, 3, 364-367.	2.0	8
137	Lymphedema in Noonan syndrome: Clues to pathogenesis and prenatal diagnosis and review of the literature. American Journal of Medical Genetics Part A, 1987, 27, 841-856.	2.4	131
138	Further delineation of Weaver syndrome. Journal of Pediatrics, 1986, 108, 228-235.	1.8	51
139	Trisomy 18 with Ectopia Cordis, Omphalocele, and Ventricular Septal Defect: Case Report. Pediatric Pathology, 1986, 5, 481-483.	0.5	33
140	The Embryology of Birth Defects. , 1983, , 33-65.		4
141	Adjacent 2 translocation involving 13q and 21q.. Journal of Medical Genetics, 1982, 19, 314-315.	3.2	3
142	Vascular pathogenesis of limb defects. I. Radial artery anatomy in radial aplasia. Journal of Pediatrics, 1982, 101, 832-838.	1.8	58
143	Vascular pathogenesis of transverse limb reduction defects. Journal of Pediatrics, 1982, 101, 839-843.	1.8	203
144	The vascular pathogenesis of gastroschisis: Intrauterine interruption of the omphalomesenteric artery. Journal of Pediatrics, 1981, 98, 228-231.	1.8	260

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145	Presentation of mucopolysaccharidosis VII (beta-glucuronidase deficiency) in infancy.. Journal of Medical Genetics, 1981, 18, 237-239.	3.2	19
146	Carbohydrate metabolism in Acanthamoeba castellanii. 1. The activity of key enzymes and [14C]-glucose metabolism. Canadian Journal of Microbiology, 1973, 19, 1131-1136.	1.7	9
147	Fetal and neonatal injury as a consequence of maternal substance abuse. , 0, , 110-126.		0
148	Fetal and Neonatal Injury as a Consequence of Maternal Substance Abuse. , 0, , 135-161.		0
149	Interactive Feature Visualization and Detection for 3D Face Classification. , 0, , 98-110.		0