Lisa J Martin

List of Publications by Year
in descending order

286 papers	12,380 citations	28190 55 h-index	35952 97 g-index
295 all docs	295 docs citations	295 times ranked	16005 citing authors

#	Article	IF	Citations
1	Quantitative trait loci on chromosomes 3 and 17 influence phenotypes of the metabolic syndrome. Proceedings of the National Academy of Sciences of the United States of America, 2000, 97, 14478-14483.	3.3	584
2	Bicuspid aortic valve is heritable. Journal of the American College of Cardiology, 2004, 44, 138-143.	1.2	560
3	Common variants at 5q22 associate with pediatric eosinophilic esophagitis. Nature Genetics, 2010, 42, 289-291.	9.4	397
4	A comprehensive linkage analysis for myocardial infarction and its related risk factors. Nature Genetics, 2002, 30, 210-214.	9.4	313
5	The Genetic Basis of Plasma Variation in Adiponectin, a Global Endophenotype for Obesity and the Metabolic Syndrome. Journal of Clinical Endocrinology and Metabolism, 2001, 86, 4321-4325.	1.8	273
6	Coordinate Interaction between IL-13 and Epithelial Differentiation Cluster Genes in Eosinophilic Esophagitis. Journal of Immunology, 2010, 184, 4033-4041.	0.4	257
7	Genome-wide association analysis of eosinophilic esophagitis provides insight into the tissue specificity of this allergic disease. Nature Genetics, 2014, 46, 895-900.	9.4	243
8	Variants of thymic stromal lymphopoietin and its receptor associate with eosinophilic esophagitis. Journal of Allergy and Clinical Immunology, 2010, 126, 160-165.e3.	1.5	236
9	Meta-analysis of Genome-wide Association Studies Identifies 1q22 as a Susceptibility Locus for Intracerebral Hemorrhage. American Journal of Human Genetics, 2014, 94, 511-521.	2.6	235
10	Twin and family studies reveal strong environmental and weaker genetic cues explaining heritability of eosinophilic esophagitis. Journal of Allergy and Clinical Immunology, 2014, 134, 1084-1092.e1.	1.5	218
11	Hypoplastic Left Heart Syndrome Is Heritable. Journal of the American College of Cardiology, 2007, 50, 1590-1595.	1.2	216
12	The complex genetics of hypoplastic left heart syndrome. Nature Genetics, 2017, 49, 1152-1159.	9.4	177
13	Gallstones. Annals of Surgery, 2002, 235, 842-849.	2.1	172
14	Evidence in favor of linkage to human chromosomal regions 18q, 5q and 13q for bicuspid aortic valve and associated cardiovascular malformations. Human Genetics, 2007, 121, 275-284.	1.8	167
15	Newly developed and validated eosinophilic esophagitis histology scoring system and evidence that it outperforms peak eosinophil count for disease diagnosis and monitoring. Ecological Management and Restoration, 2016, 30, n/a-n/a.	0.2	154
16	Adiponectin is present in human milk and is associated with maternal factors. American Journal of Clinical Nutrition, 2006, 83, 1106-1111.	2.2	152
17	High prevalence of eosinophilic esophagitis in patients with inherited connective tissue disorders. Journal of Allergy and Clinical Immunology, 2013, 132, 378-386.	1.5	150
18	Functional Variant in the Autophagy-Related 5 Gene Promotor is Associated with Childhood Asthma. PLoS ONE, 2012, 7, e33454.	1.1	148

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19	Race and Unequal Burden of Perioperative Pain and Opioid Related Adverse Effects in Children. Pediatrics, 2012, 129, 832-838.	1.0	145
20	MicroRNA signature in patients with eosinophilic esophagitis, reversibility with glucocorticoids, and assessment as disease biomarkers. Journal of Allergy and Clinical Immunology, 2012, 129, 1064-1075.e9.	1.5	145
21	Complement drives glucosylceramide accumulation and tissue inflammation in Gaucher disease. Nature, 2017, 543, 108-112.	13.7	145
22	Metaâ€Analysis of Genomeâ€wide Linkage Studies in BMI and Obesity. Obesity, 2007, 15, 2263-2275.	1.5	138
23	Hypoplastic Left Heart Syndrome Links to Chromosomes 10q and 6q and Is Genetically Related to Bicuspid Aortic Valve. Journal of the American College of Cardiology, 2009, 53, 1065-1071.	1.2	132
24	Genetic contributions to social impulsivity and aggressiveness in vervet monkeys. Biological Psychiatry, 2004, 55, 642-647.	0.7	126
25	Pediatric Plexiform Neurofibromas: Impact on Morbidity and Mortality in Neurofibromatosis Type 1. Journal of Pediatrics, 2012, 160, 461-467.	0.9	122
26	Genetic determinants of risk in pulmonary arterial hypertension: international genome-wide association studies and meta-analysis. Lancet Respiratory Medicine,the, 2019, 7, 227-238.	5.2	122
27	The effect of minor allele frequency on the likelihood of obtaining false positives. BMC Proceedings, 2009, 3, S41.	1.8	121
28	Pediatric Eosinophilic Esophagitis Symptom Scores (PEESS v2.0) identify histologic and molecular correlates of the key clinical features of disease. Journal of Allergy and Clinical Immunology, 2015, 135, 1519-1528.e8.	1.5	118
29	The Baboon as a Nonhuman Primate Model for the Study of the Genetics of Obesity. Obesity, 2003, 11, 75-80.	4.0	115
30	Common variation in COL4A1/COL4A2 is associated with sporadic cerebral small vessel disease. Neurology, 2015, 84, 918-926.	1.5	106
31	The Relationships of Adiponectin with Insulin and Lipids Are Strengthened with Increasing Adiposity. Journal of Clinical Endocrinology and Metabolism, 2005, 90, 4255-4259.	1.8	95
32	Heritability of obesity-related traits among Nigerians, Jamaicans and US black people. International Journal of Obesity, 2001, 25, 1034-1041.	1.6	93
33	Evidence for linkage on 21q and 7q in a subset of autism characterized by developmental regression. Molecular Psychiatry, 2005, 10, 741-746.	4.1	91
34	Characterization of Congenital Anomalies in Individuals With Choanal Atresia. JAMA Otolaryngology, 2009, 135, 543.	1.5	91
35	Prenatal, intrapartum, and postnatal factors are associated with pediatric eosinophilic esophagitis. Journal of Allergy and Clinical Immunology, 2018, 141, 214-222.	1.5	91
36	Human Milk Adiponectin Is Associated with Infant Growth in Two Independent Cohorts. Breastfeeding Medicine, 2009, 4, 101-109.	0.8	90

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37	Association of Blood Pressure Level With Left Ventricular Mass in Adolescents. Hypertension, 2019, 74, 590-596.	1.3	87
38	Predicting the pain continuum after adolescent idiopathic scoliosis surgery: A prospective cohort study. European Journal of Pain, 2017, 21, 1252-1265.	1.4	86
39	Novel risk genes and mechanisms implicated by exome sequencing of 2572 individuals with pulmonary arterial hypertension. Genome Medicine, 2019, 11, 69.	3.6	86
40	Insulin resistance and arterial stiffness in healthy adolescents and young adults. Diabetologia, 2012, 55, 625-631.	2.9	81
41	Retrospective Study of Obesity in Children with Down Syndrome. Journal of Pediatrics, 2016, 173, 143-148.	0.9	80
42	Opioid-induced respiratory depression: ABCB1 transporter pharmacogenetics. Pharmacogenomics Journal, 2015, 15, 119-126.	0.9	77
43	Early-life environmental exposures interact with genetic susceptibility variants in pediatric patients with eosinophilic esophagitis. Journal of Allergy and Clinical Immunology, 2018, 141, 632-637.e5.	1.5	76
44	Lifespan in captive baboons is heritable. Mechanisms of Ageing and Development, 2002, 123, 1461-1467.	2.2	75
45	Genetics of pain perception, COMT and postoperative pain management in children. Pharmacogenomics, 2014, 15, 277-284.	0.6	73
46	The antiprotease SPINK7 serves as an inhibitory checkpoint for esophageal epithelial inflammatory responses. Science Translational Medicine, 2018, 10, .	5.8	71
47	Influence of CYP2C19 Metabolizer Status on Escitalopram/Citalopram Tolerability and Response in Youth With Anxiety and Depressive Disorders. Frontiers in Pharmacology, 2019, 10, 99.	1.6	70
48	Regions of homozygosity identified by SNP microarray analysis aid in the diagnosis of autosomal recessive disease and incidentally detect parental blood relationships. Genetics in Medicine, 2013, 15, 70-78.	1.1	69
49	Human Milk Adiponectin Affects Infant Weight Trajectory During the Second Year of Life. Journal of Pediatric Gastroenterology and Nutrition, 2012, 54, 532-539.	0.9	68
50	Comparison of measures of marker informativeness for ancestry and admixture mapping. BMC Genomics, 2011, 12, 622.	1.2	65
51	Searching for genes underlying normal variation in human adiposity. Journal of Molecular Medicine, 2001, 79, 57-70.	1.7	64
52	Differences in Candidate Gene Association between European Ancestry and African American Asthmatic Children. PLoS ONE, 2011, 6, e16522.	1.1	61
53	MiR-375 is downregulated in epithelial cells after IL-13 stimulation and regulates an IL-13-induced epithelial transcriptome. Mucosal Immunology, 2012, 5, 388-396.	2.7	60
54	Population Genomics and the Statistical Values of Race: An Interdisciplinary Perspective on the Biological Classification of Human Populations and Implications for Clinical Genetic Epidemiological Research. Frontiers in Genetics, 2016, 7, 22.	1.1	58

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55	A Pediatric Asthma Risk Score to better predict asthma development in young children. Journal of Allergy and Clinical Immunology, 2019, 143, 1803-1810.e2.	1.5	58
56	Interactions Between Noncontiguous Haplotypes in the Adiponectin Gene ACDC Are Associated With Plasma Adiponectin. Diabetes, 2006, 55, 523-529.	0.3	57
57	Epigenetic modification: a regulatory mechanism in essential hypertension. Hypertension Research, 2019, 42, 1099-1113.	1.5	57
58	International Consensus Recommendations for Eosinophilic Gastrointestinal Disease Nomenclature. Clinical Gastroenterology and Hepatology, 2022, 20, 2474-2484.e3.	2.4	57
59	Variable expression of neurofibromatosis 1 in monozygotic twins. American Journal of Medical Genetics, Part A, 2011, 155, 478-485.	0.7	55
60	Genetics of monoamine metabolites in baboons: overlapping sets of genes influence levels of 5-hydroxyindolacetic acid, 3-hydroxy-4-methoxyphenylglycol, and homovanillic acid. Biological Psychiatry, 2004, 55, 739-744.	0.7	53
61	Pharmacogenetics of antiepileptic drug efficacy in childhood absence epilepsy. Annals of Neurology, 2017, 81, 444-453.	2.8	53
62	Eosinophilic esophagitis (EoE) genetic susceptibility is mediated by synergistic interactions between EoE-specific and general atopic disease loci. Journal of Allergy and Clinical Immunology, 2018, 141, 1690-1698.	1.5	51
63	Genomic architecture of asthma differs by sex. Genomics, 2015, 106, 15-22.	1.3	48
64	The genetic etiology of eosinophilic esophagitis. Journal of Allergy and Clinical Immunology, 2020, 145, 9-15.	1.5	48
65	Cellular sources of interleukin-6 and associations with clinical phenotypes and outcomes in pulmonary arterial hypertension. European Respiratory Journal, 2020, 55, 1901761.	3.1	48
66	Experience with hemihyperplasia and Beckwithâ€“Wiedemann syndrome surveillance protocol. American Journal of Medical Genetics, Part A, 2009, 149A, 1691-1697.	0.7	47
67	Does Breastfeeding Protect Against Childhood Obesity? Moving Beyond Observational Evidence. Current Obesity Reports, 2015, 4, 207-216.	3.5	47
68	Identification of KIF3A as a Novel Candidate Gene for Childhood Asthma Using RNA Expression and Population Allelic Frequencies Differences. PLoS ONE, 2011, 6, e23714.	1.1	46
69	Rank-based genome-wide analysis reveals the association of Ryanodine receptor-2 gene variants with childhood asthma among human populations. Human Genomics, 2013, 7, 16.	1.4	46
70	EMR-linked GWAS study: investigation of variation landscape of loci for body mass index in children. Frontiers in Genetics, 2013, 4, 268.	1.1	46
71	Variation in menstrual cycle length and cessation of menstruation in captive raised baboons. Mechanisms of Ageing and Development, 2003, 124, 865-871.	2.2	45
72	Major Quantitative Trait Locus for Resting Heart Rate Maps to a Region on Chromosome 4. Hypertension, 2004, 43, 1146-1151.	1.3	44

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73	Identification of factors predicting scar outcome after burn in adults: A prospective caseâ€“control study. Burns, 2017, 43, 1271-1283.	1.1	44
74	The Genetic Landscape of Hypoplastic Left Heart Syndrome. Pediatric Cardiology, 2018, 39, 1069-1081.	0.6	44
75	Adolescent Sex Differences in Adiponectin Are Conditional on Pubertal Development and Adiposity. Obesity, 2005, 13, 2095-2101.	4.0	43
76	Vanin-1 expression and methylation discriminate pediatric asthma corticosteroid treatment response. Journal of Allergy and Clinical Immunology, 2015, 136, 923-931.e3.	1.5	43
77	Risk factors for aortic valve disease in bicuspid aortic valve: A familyâ€based study. American Journal of Medical Genetics, Part A, 2011, 155, 1015-1020.	0.7	42
78	OCT1 genetic variants are associated with postoperative morphine-related adverse effects in children. Pharmacogenomics, 2017, 18, 621-629.	0.6	42
79	Leptin's Sexual Dimorphism Results from Genotype by Sex Interactions Mediated by Testosterone. Obesity, 2002, 10, 14-21.	4.0	41
80	Whole Exome Sequencing for Familial Bicuspid Aortic Valve Identifies Putative Variants. Circulation: Cardiovascular Genetics, 2014, 7, 677-683.	5.1	41
81	Increased Prevalence of Eosinophilic Gastrointestinal Disorders in Pediatric PTEN Hamartoma Tumor Syndromes. Journal of Pediatric Gastroenterology and Nutrition, 2014, 58, 553-560.	0.9	41
82	Association of FTO Gene Variants With Adiposity in Africanâ€American Adolescents. Obesity, 2010, 18, 1959-1963.	1.5	40
83	DNA methylation at the mu-1 opioid receptor gene (OPRM1) promoter predicts preoperative, acute, and chronic postsurgical pain after spine fusion. Pharmacogenomics and Personalized Medicine, 2017, Volume 10, 157-168.	0.4	40
84	SHIP-AHOY (Study of High Blood Pressure in Pediatrics: Adult Hypertension Onset in Youth). Hypertension, 2018, 72, 625-631.	1.3	40
85	Impaired immune function in children with Fanconi anaemia. British Journal of Haematology, 2011, 154, 234-240.	1.2	38
86	Codeineâ€related adverse drug reactions in children following tonsillectomy: A prospective study. Laryngoscope, 2014, 124, 1242-1250.	1.1	38
87	Subclinical Systolic and Diastolic Dysfunction Is Evident in Youth With Elevated Blood Pressure. Hypertension, 2020, 75, 1551-1556.	1.3	38
88	Quality assessment of buccal versus blood genomic DNA using the Affymetrix 500 K GeneChip. BMC Genetics, 2007, 8, 79.	2.7	37
89	Genetics and outcomes after traumatic brain injury (TBI): What do we know about pediatric TBI?. Journal of Pediatric Rehabilitation Medicine, 2012, 5, 217-231.	0.3	37
90	Epistasis between serine protease inhibitor Kazal-type 5 (SPINK5) and thymic stromal lymphopoietin (TSLP) genes contributes to childhood asthma. Journal of Allergy and Clinical Immunology, 2014, 134, 891-899.e3.	1.5	37

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91	Esophageal type 2 cytokine expression heterogeneity in eosinophilic esophagitis in a multisite cohort. Journal of Allergy and Clinical Immunology, 2020, 145, 1629-1640.e4.	1.5	37
92	Association of OPRM1 A118G variant with risk of morphine-induced respiratory depression following spine fusion in adolescents. Pharmacogenomics Journal, 2015, 15, 255-262.	0.9	36
93	Pediatric and Adult Ambulatory Blood Pressure Thresholds and Blood Pressure Load as Predictors of Left Ventricular Hypertrophy in Adolescents. Hypertension, 2021, 78, 30-37.	1.3	36
94	Polymorphisms in adiponectin receptor genes ADIPOR1 and ADIPOR2 and insulin resistance. Obesity Reviews, 2007, 8, 419-423.	3.1	35
95	INSIG1 influences obesity-related hypertriglyceridemia in humans. Journal of Lipid Research, 2010, 51, 701-708.	2.0	34
96	The role of mitochondrial genome in essential hypertension in a Chinese Han population. European Journal of Human Genetics, 2009, 17, 1501-1506.	1.4	33
97	Genetic variants in SCN5A promoter are associated with arrhythmia phenotype severity in patients with heterozygous loss-of-function mutation. Heart Rhythm, 2012, 9, 1090-1096.	0.3	33
98	Biofilm propensity of Staphylococcus aureus skin isolates is associated with increased atopic dermatitis severity and barrier dysfunction in the MPAACH pediatric cohort. Allergy: European Journal of Allergy and Clinical Immunology, 2021, 76, 302-313.	2.7	33
99	Genetic determinants of obesity-related lipid traits. Journal of Lipid Research, 2004, 45, 610-615.	2.0	32
100	Opioid-Related Adverse Effects in Children Undergoing Surgery: Unequal Burden on Younger Girls with Higher Doses of Opioids. Pain Medicine, 2015, 16, 985-997.	0.9	32
101	Heterogeneity in Asthma Care in a Statewide Collaborative: the Ohio Pediatric Asthma Repository. Pediatrics, 2015, 135, 271-279.	1.0	32
102	Pharmacogenetics of Sertraline Tolerability and Response in Pediatric Anxiety and Depressive Disorders. Journal of Child and Adolescent Psychopharmacology, 2019, 29, 348-361.	0.7	32
103	Eosinophilic Esophagitis Histology Remission Score. Journal of Pediatric Gastroenterology and Nutrition, 2020, 70, 598-603.	0.9	32
104	Value of an Additional Review for Eosinophil Quantification in Esophageal Biopsies. Journal of Pediatric Gastroenterology and Nutrition, 2015, 61, 65-68.	0.9	32
105	Novel associations between FAAH genetic variants and postoperative central opioid-related adverse effects. Pharmacogenomics Journal, 2015, 15, 436-442.	0.9	31
106	Genetics of Hypoplastic Left Heart Syndrome. Journal of Pediatrics, 2016, 173, 25-31.	0.9	31
107	Barriers to and Motivations for Physician Referral of Patients to Cancer Genetics Clinics. Journal of Genetic Counseling, 2012, 21, 305-325.	0.9	30
108	Identification of factors predicting scar outcome after burn injury in children: a prospective case-control study. Burns and Trauma, 2017, 5, 19.	2.3	30

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109	Genetic variants at the 16p13 locus confer risk for eosinophilic esophagitis. Genes and Immunity, 2019, 20, 281-292.	2.2	30
110	The Genomics Research and Innovation Network: creating an interoperable, federated, genomics learning system. Genetics in Medicine, 2020, 22, 371-380.	1.1	30
111	Genetic Causes of Cardiomyopathy in Children: First Results From the Pediatric Cardiomyopathy Genes Study. Journal of the American Heart Association, 2021, 10, e017731.	1.6	29
112	Pediatric asthma incidence rates in the United States from 1980 to 2017. Journal of Allergy and Clinical Immunology, 2021, 148, 1270-1280.	1.5	28
113	Population structure analysis using rare and common functional variants. BMC Proceedings, 2011, 5, S8.	1.8	27
114	ABCC3 genetic variants are associated with postoperative morphine-induced respiratory depression and morphine pharmacokinetics in children. Pharmacogenomics Journal, 2017, 17, 162-169.	0.9	27
115	A Quantitative Trait Locus Influencing Type 2 Diabetes Susceptibility Maps to a Region on 5q in an Extended French Family. Diabetes, 2002, 51, 3568-3572.	0.3	26
116	Aorta Measurements are Heritable and Influenced by Bicuspid Aortic Valve. Frontiers in Genetics, 2011, 2, 61.	1.1	26
117	Genetic risk signatures of opioid-induced respiratory depression following pediatric tonsillectomy. Pharmacogenomics, 2014, 15, 1749-1762.	0.6	26
118	Adolescentsâ€™ preferences regarding disclosure of incidental findings in genomic sequencing that are not medically actionable in childhood. American Journal of Medical Genetics, Part A, 2016, 170, 2083-2088.	0.7	26
119	Influence of Dopamine-Related Genes on Neurobehavioral Recovery after Traumatic Brain Injury during Early Childhood. Journal of Neurotrauma, 2017, 34, 1919-1931.	1.7	26
120	Very early onset eosinophilic esophagitis is common, responds to standard therapy, and demonstrates enrichment for CAPN14 genetic variants. Journal of Allergy and Clinical Immunology, 2021, 147, 244-254.e6.	1.5	26
121	Personalized Medicine, Availability, and Group Disparity: An Inquiry into How Physicians Perceive and Rate the Elements and Barriers of Personalized Medicine. Public Health Genomics, 2014, 17, 209-220.	0.6	25
122	Applying Systems Biology Methodology To Identify Genetic Factors Possibly Associated with Recovery after Traumatic Brain Injury. Journal of Neurotrauma, 2017, 34, 2280-2290.	1.7	25
123	Longitudinal atopic dermatitis endotypes: An atopic march paradigm that includes Black children. Journal of Allergy and Clinical Immunology, 2022, 149, 1702-1710.e4.	1.5	25
124	Familial resemblance of body composition in prepubertal girls and their biological parents. American Journal of Clinical Nutrition, 2001, 74, 529-533.	2.2	24
125	Genetic Control of Coordinated Changes in HDL and LDL Size Phenotypes. Arteriosclerosis, Thrombosis, and Vascular Biology, 2001, 21, 1829-1833.	1.1	24
126	Genotype by smoking interaction for leptin levels in the San Antonio family heart study. Genetic Epidemiology, 2002, 22, 105-115.	0.6	24

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127	Phenotypic, genetic, and genome-wide structure in the metabolic syndrome. BMC Genetics, 2003, 4, S95.	2.7	24
128	Ambulatory blood pressure monitoring tolerability and blood pressure status in adolescents. Blood Pressure Monitoring, 2019, 24, 12-17.	0.4	24
129	Identification of anoctamin 1 (ANO1) as a key driver of esophageal epithelial proliferation in eosinophilic esophagitis. Journal of Allergy and Clinical Immunology, 2020, 145, 239-254.e2.	1.5	24
130	Disease-associated KIF3A variants alter gene methylation and expression impacting skin barrier and atopic dermatitis risk. Nature Communications, 2020, 11, 4092.	5.8	24
131	Bivariate Linkage between Acylationâ€Stimulating Protein and BMI and Highâ€Density Lipoproteins. Obesity, 2004, 12, 669-678.	4.0	23
132	Serotonin (5-HT) receptor 5A sequence variants affect human plasma triglyceride levels. Physiological Genomics, 2010, 42, 168-176.	1.0	23
133	Influence of Catechol-O-methyltransferase on Executive Functioning Longitudinally After Early Childhood Traumatic Brain Injury. Journal of Head Trauma Rehabilitation, 2016, 31, E1-E9.	1.0	23
134	Desmoplakin and periplakin genetically and functionally contribute to eosinophilic esophagitis. Nature Communications, 2021, 12, 6795.	5.8	23
135	Genetics of leptin expression in baboons. International Journal of Obesity, 2003, 27, 778-783.	1.6	22
136	Evaluation of Growth in Patients With Isolated Cleft Lip and/or Cleft Palate. Pediatrics, 2010, 125, e543-e549.	1.0	22
137	Loss of Endothelial TSPAN12 Promotes Fibrostenotic Eosinophilic Esophagitis via Endothelial Cellâ€“Fibroblast Crosstalk. Gastroenterology, 2022, 162, 439-453.	0.6	22
138	Evidence for a gene influencing fasting LDL cholesterol and triglyceride levels on chromosome 21q. Atherosclerosis, 2005, 179, 119-125.	0.4	21
139	Stakeholder Buy-In and Physician Education Improve Adherence toÂGuidelines for Down Syndrome. Journal of Pediatrics, 2016, 171, 262-268.e2.	0.9	21
140	Rhinovirus infection results in stronger and more persistent genomic dysregulation: Evidence for altered innate immune response in asthmatics at baseline, early in infection, and during convalescence. PLoS ONE, 2017, 12, e0178096.	1.1	21
141	Disease-Related Predictors of Health-Related Quality of Life in Youth With Eosinophilic Esophagitis. Journal of Pediatric Psychology, 2018, 43, 464-471.	1.1	21
142	The genetic architecture of pediatric cardiomyopathy. American Journal of Human Genetics, 2022, 109, 282-298.	2.6	21
143	Evaluating Eosinophilic Colitis as a Unique Disease Using Colonic Molecular Profiles: A Multi-Site Study. Gastroenterology, 2022, 162, 1635-1649.	0.6	21
144	The Genetics of Obesity in Mexican Americans: The Evidence from Genome Scanning Efforts in the San Antonio Family Heart Study. Human Biology, 2003, 75, 635-646.	0.4	20

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145	Application of genetic/genomic approaches to allergic disorders. Journal of Allergy and Clinical Immunology, 2010, 126, 425-436.	1.5	20
146	Heritability of the Severity of the Metabolic Syndrome in Whites and Blacks in 3 Large Cohorts. Circulation: Cardiovascular Genetics, 2017, 10, .	5.1	20
147	Substantial pain burden in frequency, intensity, interference and chronicity among children and adults with neurofibromatosis Type 1. American Journal of Medical Genetics, Part A, 2019, 179, 602-607.	0.7	20
148	Events in Normal Skin Promote Early-Life Atopic Dermatitisâ€"The MPAACH Cohort. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 2285-2293.e6.	2.0	20
149	Noninvasive Prognostic Biomarkers for Left-Sided Heart Failure as Predictors of Survival in Pulmonary Arterial Hypertension. Chest, 2020, 157, 1606-1616.	0.4	20
150	HDL cholesterol in females in the Framingham Heart Study is linked to a region of chromosome 2q. BMC Genetics, 2003, 4, S98.	2.7	19
151	Prediction of Ambulatory Hypertension Based on Clinic Blood Pressure Percentile in Adolescents. Hypertension, 2018, 72, 955-961.	1.3	19
152	Adolescents' and Parents' Genomic Testing Decisions: Associations With Age, Race, and Sex. Journal of Adolescent Health, 2020, 66, 288-295.	1.2	19
153	Mining the Plasma Proteome for Insights into the Molecular Pathology of Pulmonary Arterial Hypertension. American Journal of Respiratory and Critical Care Medicine, 2022, 205, 1449-1460.	2.5	19
154	Retrospective comparison of patient outcomes after in-person and telephone results disclosure counseling for BRCA1/2 genetic testing. Familial Cancer, 2010, 9, 203-212.	0.9	18
155	Fractures in Children With Neurofibromatosis Type 1 From Two <scp>NF</scp> Clinics. American Journal of Medical Genetics, Part A, 2013, 161, 921-926.	0.7	18
156	KIF3A genetic variation is associated with pediatric asthma in the presence of eczema independent of allergic rhinitis. Journal of Allergy and Clinical Immunology, 2017, 140, 595-598.e5.	1.5	18
157	Early life factors are associated with risk for eosinophilic esophagitis diagnosed in adulthood. Ecological Management and Restoration, 2021, 34, .	0.2	18
158	Genotype-by-smoking interaction for leptin levels in the Metabolic Risk Complications of Obesity Genes project. International Journal of Obesity, 2003, 27, 334-340.	1.6	17
159	Effect of population stratification analysis on false-positive rates for common and rare variants. BMC Proceedings, 2011, 5, S116.	1.8	17
160	Eosinophil progenitor levels are increased in patients with active pediatric eosinophilic esophagitis. Journal of Allergy and Clinical Immunology, 2016, 138, 915-918.e5.	1.5	17
161	Mycophenolate mofetilâ€related leukopenia in children and young adults following kidney transplantation: Influence of genes and drugs. Pediatric Transplantation, 2017, 21, e13033.	0.5	17
162	Advancing patient care through the Consortium of Eosinophilic Gastrointestinal Disease Researchers (CEGIR). Journal of Allergy and Clinical Immunology, 2020, 145, 28-37.	1.5	17

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163	Sensitization to peanut, egg or pets is associated with skin barrier dysfunction in children with atopic dermatitis. Clinical and Experimental Allergy, 2021, 51, 666-673.	1.4	17
164	The Genes Influencing Adiponectin Levels Also Influence Risk Factors for Metabolic Syndrome and Type 2 Diabetes. Human Biology, 2007, 79, 191-200.	0.4	16
165	Association of INSIG2 Polymorphism with Overweight and LDL in Children. PLoS ONE, 2015, 10, e0116340.	1.1	16
166	Genetic Influences on Behavioral Outcomes After Childhood TBI: A Novel Systems Biology-Informed Approach. Frontiers in Genetics, 2019, 10, 481.	1.1	16
167	A quantitative trait locus influencing estrogen levels maps to a region homologous to human chromosome 20. Physiological Genomics, 2001, 5, 75-80.	1.0	15
168	Shared genetic contributions of fruit and vegetable consumption with BMI in families 20 y after sharing a household. American Journal of Clinical Nutrition, 2011, 94, 1138-1143.	2.2	15
169	Analysis of metabolic syndrome phenotypes in Framingham Heart Study families from Genetic Analysis Workshop 13. Genetic Epidemiology, 2003, 25, S78-S89.	0.6	14
170	Multiple testing in the genomics era: Findings from Genetic Analysis Workshop 15, Group 15. Genetic Epidemiology, 2007, 31, S124-S131.	0.6	14
171	The Importance of Geneâ€"Environment Interaction. Sociological Methods and Research, 2008, 37, 164-200.	4.3	14
172	The Impact of Supervision Training on Genetic Counselor Supervisory Identity Development. Journal of Genetic Counseling, 2014, 23, 1056-1065.	0.9	14
173	Multiancestral polygenic risk score for pediatric asthma. Journal of Allergy and Clinical Immunology, 2022, 150, 1086-1096.	1.5	14
174	Using Mendelian inheritance errors as quality control criteria in whole genome sequencing data set. BMC Proceedings, 2014, 8, S21.	1.8	13
175	Asthma as an outcome: Exploring multiple definitions of asthma across birth cohorts in the Environmental influences on Child Health Outcomes Children's Respiratory and Environmental Workgroup. Journal of Allergy and Clinical Immunology, 2019, 144, 866-869.e4.	1.5	13
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