Mary L Marazita

List of Publications by Year in descending order

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61857 53109 8,757 161 43 85 citations h-index g-index papers 169 169 169 7078 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Variant analyses of candidate genes in orofacial clefts in multiâ€ethnic populations. Oral Diseases, 2022, 28, 1921-1935.	1.5	3
2	Limb development genes underlie variation in human fingerprint patterns. Cell, 2022, 185, 95-112.e18.	13.5	30
3	Genome-wide Interaction Study Implicates VGLL2 and Alcohol Exposure and PRL and Smoking in Orofacial Cleft Risk. Frontiers in Cell and Developmental Biology, 2022, 10, 621261.	1.8	3
4	Genomeâ€wide association study of multiethnic nonsyndromic orofacial cleft families identifies novel loci specific to family and phenotypic subtypes. Genetic Epidemiology, 2022, , .	0.6	4
5	Exploring Mothers' Perspectives About Why Grandparents in Appalachia Give Their Grandchildren Cariogenic Foods and Beverages: A Qualitative Study. Journal of the Academy of Nutrition and Dietetics, 2022, , .	0.4	2
6	Editorial: Genetic, Environmental and Synergistic Gene-Environment Contributions to Craniofacial Defects. Frontiers in Cell and Developmental Biology, 2022, 10, 887051.	1.8	0
7	Racism in oral healthcare settings: Implications for dental <scp>careâ€related</scp> fear/anxiety and utilization among Black/African American women in Appalachia. Journal of Public Health Dentistry, 2022, 82, 28-35.	0.5	4
8	Effects of Male Facial Masculinity on Perceived Attractiveness. Adaptive Human Behavior and Physiology, 2021, 7, 73-88.	0.6	10
9	Insights into the genetic architecture of the human face. Nature Genetics, 2021, 53, 45-53.	9.4	94
10	Genome-Wide Association Analysis of Longitudinal Bone Mineral Content Data From the Iowa Bone Development Study. Journal of Clinical Densitometry, 2021, 24, 44-54.	0.5	0
11	Impact of low-frequency coding variants on human facial shape. Scientific Reports, 2021, 11, 748.	1.6	3
12	Fluctuating Asymmetry and Sexual Dimorphism in Human Facial Morphology: A Multi-Variate Study. Symmetry, 2021, 13, 304.	1.1	6
13	The Intersection of the Genetic Architectures of Orofacial Clefts and Normal Facial Variation. Frontiers in Genetics, 2021, 12, 626403.	1.1	10
14	Detecting Gene-Environment Interaction for Maternal Exposures Using Case-Parent Trios Ascertained Through a Case With Non-Syndromic Orofacial Cleft. Frontiers in Cell and Developmental Biology, 2021, 9, 621018.	1.8	2
15	Genome-Wide Association Study of Non-syndromic Orofacial Clefts in a Multiethnic Sample of Families and Controls Identifies Novel Regions. Frontiers in Cell and Developmental Biology, 2021, 9, 621482.	1.8	16
16	The PAX1 locus at $20p11$ is a potential genetic modifier for bilateral cleft lip. Human Genetics and Genomics Advances, 2021 , 2 , 100025 .	1.0	9
17	Using the PhenX Toolkit to Select Standard Measurement Protocols for Your Research Study. Current Protocols, 2021, 1, e149.	1.3	16
18	Replication of GWAS significant loci in a sub-Saharan African Cohort with early childhood caries: a pilot study. BMC Oral Health, 2021, 21, 274.	0.8	3

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19	Variant Analyses of Candidate Genes in Orofacial Clefts in Multiâ€Ethnic Populations. FASEB Journal, 2021, 35, .	0.2	0
20	<i>FAT4</i> identified as a potential modifier of orofacial cleft laterality. Genetic Epidemiology, 2021, 45, 721-735.	0.6	14
21	Associations Between Salivary Bacteriome Diversity and Salivary Human Herpesvirus Detection in Early Childhood: A Prospective Cohort Study. Journal of the Pediatric Infectious Diseases Society, 2021, 10, 856-863.	0.6	0
22	Integrative approaches generate insights into the architecture of non-syndromic cleft lip \hat{A}_{\pm} cleft palate. Human Genetics and Genomics Advances, 2021, 2, 100038.	1.0	8
23	Predictors of use of dental care by children in north-central Appalachia in the USA. PLoS ONE, 2021, 16, e0250488.	1.1	4
24	Pleiotropy method reveals genetic overlap between orofacial clefts at multiple novel loci from GWAS of multi-ethnic trios. PLoS Genetics, 2021, 17, e1009584.	1.5	18
25	Genome-Wide Association Study (GWAS) of dental caries in diverse populations. BMC Oral Health, 2021, 21, 377.	0.8	16
26	The Influence of Sex and Ancestry on Three-Dimensional Palate Shape. Journal of Craniofacial Surgery, 2021, 32, 2883-2887.	0.3	1
27	Exploring palatal and dental shape variation with 3D shape analysis and geometric deep learning. Orthodontics and Craniofacial Research, 2021, 24, 134-143.	1.2	12
28	Genome-Wide Scan for Parent-of-Origin Effects in a sub-Saharan African Cohort With Nonsyndromic Cleft Lip and/or Cleft Palate (CL/P). Cleft Palate-Craniofacial Journal, 2021, , 105566562110363.	0.5	1
29	Genome scans of facial features in East Africans and cross-population comparisons reveal novel associations. PLoS Genetics, 2021, 17, e1009695.	1.5	13
30	PRICKLE1 $\tilde{A}-$ FOCAD Interaction Revealed by Genome-Wide vQTL Analysis of Human Facial Traits. Frontiers in Genetics, 2021, 12, 674642.	1.1	6
31	Oral health and related risk indicators in northâ€central Appalachia differ by rurality. Community Dentistry and Oral Epidemiology, 2021, 49, 427-436.	0.9	1
32	Whole genome sequencing of orofacial cleft trios from the Gabriella Miller Kids First Pediatric Research Consortium identifies a new locus on chromosome 21. Human Genetics, 2020, 139, 215-226.	1.8	19
33	FaceBase 3: analytical tools and FAIR resources for craniofacial and dental research. Development (Cambridge), 2020, 147, .	1.2	25
34	Low levels of salivary metals, oral microbiome composition and dental decay. Scientific Reports, 2020, 10, 14640.	1.6	14
35	Expanding the genetic architecture of nicotine dependence and its shared genetics with multiple traits. Nature Communications, 2020, 11, 5562.	5.8	80
36	Co-occurrence of yeast, streptococci, dental decay, and gingivitis in the post-partum period: results of a longitudinal study. Journal of Oral Microbiology, 2020, 12, 1746494.	1.2	5

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37	Cleft lip/palate and educational attainment: cause, consequence or correlation? A Mendelian randomization study. International Journal of Epidemiology, 2020, 49, 1282-1293.	0.9	21
38	Nonâ€random distribution of deleterious mutations in the DNA and proteinâ€binding domains of <i>IRF6</i> are associated with Van Der Woude syndrome. Molecular Genetics & Denomic Medicine, 2020, 8, e1355.	0.6	13
39	Individuals with nonsyndromic orofacial clefts have increased asymmetry of fingerprint patterns. PLoS ONE, 2020, 15, e0230534.	1.1	6
40	A Multivariate Approach to Determine the Dimensionality of Human Facial Asymmetry. Symmetry, 2020, 12, 348.	1.1	9
41	Prevalence of Torus Palatinus and association with dental arch shape in a multi-ethnic cohort. HOMO- Journal of Comparative Human Biology, 2020, 71, 273-280.	0.3	5
42	Detection of de novo copy number deletions from targeted sequencing of trios. Bioinformatics, 2019, 35, 571-578.	1.8	2
43	Transmission of dental fear from parent to adolescent in an Appalachian sample in the USA. International Journal of Paediatric Dentistry, 2019, 29, 720-727.	1.0	14
44	Variants in the fetal genome near pro-inflammatory cytokine genes on 2q13 associate with gestational duration. Nature Communications, 2019, 10, 3927.	5.8	49
45	Genome-wide analysis of dental caries and periodontitis combining clinical and self-reported data. Nature Communications, 2019, 10, 2773.	5.8	183
46	A systematic genetic analysis and visualization of phenotypic heterogeneity among orofacial cleft GWAS signals. Genetic Epidemiology, 2019, 43, 704-716.	0.6	36
47	Predictors of dental care utilization in northâ€eentral Appalachia in the USA. Community Dentistry and Oral Epidemiology, 2019, 47, 283-290.	0.9	17
48	Facial recognition from DNA using face-to-DNA classifiers. Nature Communications, 2019, 10, 2557.	5.8	46
49	Primary teeth microhardness and lead (Pb) levels. Heliyon, 2019, 5, e01551.	1.4	4
50	Speech Phenotyping in Unaffected Family Members of Individuals With Nonsyndromic Cleft Lip With or Without Palate. Cleft Palate-Craniofacial Journal, 2019, 56, 867-876.	0.5	2
51	Association of lowâ€frequency genetic variants in regulatory regions with nonsyndromic orofacial clefts. American Journal of Medical Genetics, Part A, 2019, 179, 467-474.	0.7	18
52	Genetic association and differential expression of PITX2 with acute appendicitis. Human Genetics, 2019, 138, 37-47.	1.8	14
53	Genomic analyses in African populations identify novel risk loci for cleft palate. Human Molecular Genetics, 2019, 28, 1038-1051.	1.4	61
54	Inferring disease risk genes from sequencing data in multiplex pedigrees through sharing of rare variants. Genetic Epidemiology, 2019, 43, 37-49.	0.6	6

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55	Novel genetic loci affecting facial shape variation in humans. ELife, 2019, 8, .	2.8	58
56	Mother's Perceived Social Support and Children's Dental Caries in Northern Appalachia. Pediatric Dentistry (discontinued), 2019, 41, 200-205.	0.4	4
57	Novel <i>GREM1</i> Variations in Sub-Saharan African Patients With Cleft Lip and/or Cleft Palate. Cleft Palate-Craniofacial Journal, 2018, 55, 736-742.	0.5	9
58	Is the Fagerstr \tilde{A} ¶m test for nicotine dependence invariant across secular trends in smoking? A question for cross-birth cohort analysis of nicotine dependence. Drug and Alcohol Dependence, 2018, 185, 127-132.	1.6	3
59	Soft tissue nasal asymmetry as an indicator of orofacial cleft predisposition. American Journal of Medical Genetics, Part A, 2018, 176, 1296-1303.	0.7	4
60	Genome-wide mapping of global-to-local genetic effects on human facial shape. Nature Genetics, 2018, 50, 414-423.	9.4	205
61	Vitamin D metabolic loci and preeclampsia risk in multi-ethnic pregnant women. Physiological Reports, 2018, 6, e13468.	0.7	10
62	Genetic correlation between smoking behaviors and schizophrenia. Schizophrenia Research, 2018, 194, 86-90.	1.1	71
63	Vitamin D metabolic loci and vitamin D status in Black and White pregnant women. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2018, 220, 61-68.	0.5	10
64	Exploring the genomic basis of early childhood caries: a pilot study. International Journal of Paediatric Dentistry, 2018, 28, 217-225.	1.0	24
65	SNPs Associated With Testosterone Levels Influence Human Facial Morphology. Frontiers in Genetics, 2018, 9, 497.	1.1	23
66	Six NSCL/P Loci Show Associations With Normal-Range Craniofacial Variation. Frontiers in Genetics, 2018, 9, 502.	1.1	20
67	Genomeâ€wide interaction studies identify sexâ€specific risk alleles for nonsyndromic orofacial clefts. Genetic Epidemiology, 2018, 42, 664-672.	0.6	15
68	Exploration of genetic factors determining cleft side in a pair of monozygotic twins with mirror-image cleft lip and palate using whole-genome sequencing and comparison of craniofacial morphology. Archives of Oral Biology, 2018, 96, 33-38.	0.8	8
69	Identification of paternal uniparental disomy on chromosome 22 and a <i>de novo</i> deletion on chromosome 18 in individuals with orofacial clefts. Molecular Genetics & Enomic Medicine, 2018, 6, 924-932.	0.6	4
70	GWAS reveals loci associated with velopharyngeal dysfunction. Scientific Reports, 2018, 8, 8470.	1.6	8
71	Consortium-based genome-wide meta-analysis for childhood dental caries traits. Human Molecular Genetics, 2018, 27, 3113-3127.	1.4	32
72	Novel caries loci in children and adults implicated by genome-wide analysis of families. BMC Oral Health, 2018, 18, 98.	0.8	8

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73	Investigating the shared genetics of non-syndromic cleft lip/palate and facial morphology. PLoS Genetics, 2018, 14, e1007501.	1.5	44
74	Mapping genetic variants for cranial vault shape in humans. PLoS ONE, 2018, 13, e0196148.	1.1	11
75	Wholeâ€genome sequencing in a pair of monozygotic twins with discordant cleft lip and palate subtypes. Oral Diseases, 2018, 24, 1303-1309.	1.5	5
76	Genetic variants in pachyonychia congenita-associated keratins increase susceptibility to tooth decay. PLoS Genetics, 2018, 14, e1007168.	1,5	12
77	Identifying genetic risk loci for diabetic complications and showing evidence for heterogeneity of type 1 diabetes based on complications risk. PLoS ONE, 2018, 13, e0192696.	1.1	6
78	Genome-wide meta-analyses of nonsyndromic orofacial clefts identify novel associations between FOXE1 and all orofacial clefts, and TP63 and cleft lip with or without cleft palate. Human Genetics, 2017, 136, 275-286.	1.8	139
79	Genome-wide analyses of non-syndromic cleft lip with palate identify 14 novel loci and genetic heterogeneity. Nature Communications, 2017, 8, 14364.	5. 8	207
80	Variants on chromosome 4q21 near PKD2 and SIBLINGs are associated with dental caries. Journal of Human Genetics, 2017, 62, 491-496.	1.1	11
81	Geneâ€gene interaction of single nucleotide polymorphisms in 16p13.3 may contribute to the risk of nonâ€syndromic cleft lip with or without cleft palate in Chinese caseâ€parent trios. American Journal of Medical Genetics, Part A, 2017, 173, 1489-1494.	0.7	4
82	Association studies of lowâ€frequency coding variants in nonsyndromic cleft lip with or without cleft palate. American Journal of Medical Genetics, Part A, 2017, 173, 1531-1538.	0.7	36
83	Sonic Hedgehog regulation of <i>Foxf2</i> promotes cranial neural crest mesenchyme proliferation and is disrupted in cleft lip morphogenesis. Development (Cambridge), 2017, 144, 2082-2091.	1.2	55
84	Whole exome association of rare deletions in multiplex oral cleft families. Genetic Epidemiology, 2017, 41, 61-69.	0.6	10
85	Identification of common non-coding variants at 1p22 that are functional for non-syndromic orofacial clefting. Nature Communications, 2017, 8, 14759.	5.8	48
86	Evidence for SNP-SNP interaction identified through targeted sequencing of cleft case-parent trios. Genetic Epidemiology, 2017, 41, 244-250.	0.6	24
87	Testing the face shape hypothesis in twins discordant for nonsyndromic orofacial clefting. American Journal of Medical Genetics, Part A, 2017, 173, 2886-2892.	0.7	7
88	Diagnosing subtle palatal anomalies: Validation of video-analysis and assessment protocol for diagnosing occult submucous cleft palate. International Journal of Pediatric Otorhinolaryngology, 2017, 100, 242-246.	0.4	6
89	Identifying Genetic Sources of Phenotypic Heterogeneity in Orofacial Clefts by Targeted Sequencing. Birth Defects Research, 2017, 109, 1030-1038.	0.8	41
90	Multiethnic GWAS Reveals Polygenic Architecture of Earlobe Attachment. American Journal of Human Genetics, 2017, 101, 913-924.	2.6	29

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91	Identification of 16q21 as a modifier of nonsyndromic orofacial cleft phenotypes. Genetic Epidemiology, 2017, 41, 887-897.	0.6	24
92	Toward a genetic understanding of dental fear: evidence of heritability. Community Dentistry and Oral Epidemiology, 2017, 45, 66-73.	0.9	20
93	A Preliminary Genome-Wide Association Study of Pain-Related Fear: Implications for Orofacial Pain. Pain Research and Management, 2017, 2017, 1-12.	0.7	20
94	Periodontal Status and Quality of Life: Impact of Fear of Pain and Dental Fear. Pain Research and Management, 2017, 2017, 1-9.	0.7	19
95	Genome-wide association study of facial morphology reveals novel associations with FREM1 and PARK2. PLoS ONE, 2017, 12, e0176566.	1.1	68
96	Genome-wide analysis of parent-of-origin interaction effects with environmental exposure (PoOxE): An application to European and Asian cleft palate trios. PLoS ONE, 2017, 12, e0184358.	1.1	16
97	ldentifying Genetic Sources of Phenotypic Heterogeneity in Orofacial Clefts by Targeted Sequencing. Birth Defects Research, 2017, , .	0.8	0
98	Genetic factors influencing risk to orofacial clefts: today's challenges and tomorrow's opportunities. F1000Research, 2016, 5, 2800.	0.8	155
99	Genome-Wide Association Study Reveals Multiple Loci Influencing Normal Human Facial Morphology. PLoS Genetics, 2016, 12, e1006149.	1.5	140
100	The 3D Facial Norms Database: Part 1. A Web-Based Craniofacial Anthropometric and Image Repository for the Clinical and Research Community. Cleft Palate-Craniofacial Journal, 2016, 53, 185-197.	0.5	80
101	Effects of Specimen Collection Methodologies and Storage Conditions on the Short-Term Stability of Oral Microbiome Taxonomy. Applied and Environmental Microbiology, 2016, 82, 5519-5529.	1.4	30
102	The effects of family, dentition, and dental caries on the salivary microbiome. Annals of Epidemiology, 2016, 26, 348-354.	0.9	19
103	A multi-ethnic genome-wide association study identifies novel loci for non-syndromic cleft lip with or without cleft palate on 2p24.2, 17q23 and 19q13. Human Molecular Genetics, 2016, 25, ddw104.	1.4	163
104	Genetic Modifiers of Patent Ductus Arteriosus in Term Infants. Journal of Pediatrics, 2016, 176, 57-61.e1.	0.9	12
105	The FaceBase Consortium: A comprehensive resource for craniofacial researchers. Development (Cambridge), 2016, 143, 2677-88.	1.2	62
106	Depression and Rural Environment Are Associated With Poor Oral Health Among Pregnant Women in Northern Appalachia. Behavior Modification, 2016, 40, 325-340.	1.1	20
107	Using the 3D Facial Norms Database to investigate craniofacial sexual dimorphism in healthy children, adolescents, and adults. Biology of Sex Differences, 2016, 7, 23.	1.8	65
108	A Genome-wide Association Study of Nonsyndromic Cleft Palate Identifies an Etiologic Missense Variant in GRHL3. American Journal of Human Genetics, 2016, 98, 744-754.	2.6	146

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109	Genome-wide association study of dental caries in the Hispanic Communities Health Study/Study of Latinos (HCHS/SOL). Human Molecular Genetics, 2016, 25, 807-816.	1.4	29
110	Using the PhenX Toolkit to Add Standard Measures to a Study. Current Protocols in Human Genetics, 2015, 86, 1.21.1-1.21.17.	3.5	33
111	Oral Health in a Sample of Pregnant Women from Northern Appalachia (2011–2015). International Journal of Dentistry, 2015, 2015, 1-12.	0.5	32
112	Caries Experience Differs between Females and Males across Age Groups in Northern Appalachia. International Journal of Dentistry, 2015, 2015, 1-8.	0.5	30
113	Characterization of Large Structural Genetic Mosaicism in Human Autosomes. American Journal of Human Genetics, 2015, 96, 487-497.	2.6	101
114	Symptoms of Attention-Deficit Hyperactivity Disorder, Nonsyndromic Orofacial Cleft Children, and Dopamine Polymorphisms. Biological Research for Nursing, 2015, 17, 257-262.	1.0	5
115	Genetics of Orofacial Cleft Birth Defects. Current Genetic Medicine Reports, 2015, 3, 118-126.	1.9	6
116	Identification of Functional Variants for Cleft Lip with or without Cleft Palate in or near PAX7, FGFR2, and NOG by Targeted Sequencing of GWAS Loci. American Journal of Human Genetics, 2015, 96, 397-411.	2.6	150
117	Geneâ€Gene Interaction Among <i>WNT</i> Genes for Oral Cleft in Trios. Genetic Epidemiology, 2015, 39, 385-394.	0.6	30
118	Replication of $13q31.1$ association in nonsyndromic cleft lip with cleft palate in Europeans. American Journal of Medical Genetics, Part A, 2015, 167, 1054-1060.	0.7	31
119	Aquaporin 5 Interacts with Fluoride and Possibly Protects against Caries. PLoS ONE, 2015, 10, e0143068.	1.1	22
120	Genome Wide Association Study of Dental Arch Form and Occlusal Relationships in the Mixed Dentition Stage. FASEB Journal, 2015, 29, 697.4.	0.2	0
121	Evidence of Geneâ^'Environment Interaction for Two Genes on Chromosome 4 and Environmental Tobacco Smoke in Controlling the Risk of Nonsyndromic Cleft Palate. PLoS ONE, 2014, 9, e88088.	1.1	33
122	Expanding the cleft phenotype: the dental characteristics of unaffected parents of Australian children with nonâ€syndromic cleft lip and palate. International Journal of Paediatric Dentistry, 2014, 24, 286-292.	1.0	15
123	Genome-Wide Association Study of Periodontal Health Measured by Probing Depth in Adults Ages 18â°'49 years. G3: Genes, Genomes, Genetics, 2014, 4, 307-314.	0.8	54
124	Exploring the effect of dentition, dental decay and familiality on oral health using metabolomics. Infection, Genetics and Evolution, 2014, 22, 201-207.	1.0	17
125	Whole Exome Sequencing of Distant Relatives in Multiplex Families Implicates Rare Variants in Candidate Genes for Oral Clefts. Genetics, 2014, 197, 1039-1044.	1.2	79
126	Search for genetic modifiers of IRF6 and genotype–phenotype correlations in Van der Woude and popliteal pterygium syndromes. American Journal of Medical Genetics, Part A, 2013, 161, 2535-2544.	0.7	21

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127	Genetics of cleft lip and cleft palate. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2013, 163, 246-258.	0.7	336
128	Heritability of face shape in twins: a preliminary study using 3D stereophotogrammetry and geometric morphometrics. Dentistry 3000, 2013, 1, 7-11.	0.1	39
129	Shape Analysis of the Facebase 3D Facial Norms Dataset Reveals Sexual Dimorphism in Human Faces in Juveniles, Adolescents and Adults. FASEB Journal, 2013, 27, 519.5.	0.2	0
130	Genome-wide meta-analyses of nonsyndromic cleft lip with or without cleft palate identify six new risk loci. Nature Genetics, 2012, 44, 968-971.	9.4	311
131	The Evolution of Human Genetic Studies of Cleft Lip and Cleft Palate. Annual Review of Genomics and Human Genetics, 2012, 13, 263-283.	2.5	174
132	Effects of Smoking and Genotype on the PSR Index of Periodontal Disease in Adults Aged 18–49. International Journal of Environmental Research and Public Health, 2012, 9, 2839-2850.	1.2	14
133	Rapid Testing of SNPs and Gene–Environment Interactions in Case–Parent Trio Data Based on Exact Analytic Parameter Estimation. Biometrics, 2012, 68, 766-773.	0.8	34
134	Cleft lip and palate: understanding genetic and environmental influences. Nature Reviews Genetics, 2011, 12, 167-178.	7.7	1,435
135	Evidence for gene-environment interaction in a genome wide study of nonsyndromic cleft palate. Genetic Epidemiology, 2011, 35, n/a-n/a.	0.6	145
136	A genome-wide association study of cleft lip with and without cleft palate identifies risk variants near MAFB and ABCA4. Nature Genetics, 2010, 42, 525-529.	9.4	518
137	Genome Scan, Fine-Mapping, and Candidate Gene Analysis of Non-Syndromic Cleft Lip with or without Cleft Palate Reveals Phenotype-Specific Differences in Linkage and Association Results. Human Heredity, 2009, 68, 151-170.	0.4	113
138	FOXE1 association with both isolated cleft lip with or without cleft palate, and isolated cleft palate. Human Molecular Genetics, 2009, 18, 4879-4896.	1.4	136
139	Study protocol of the Center for Oral Health Research in Appalachia (COHRA) etiology study. BMC Oral Health, 2008, 8, 18.	0.8	69
140	Disruption of an AP-2 \hat{i} ± binding site in an IRF6 enhancer is associated with cleft lip. Nature Genetics, 2008, 40, 1341-1347.	9.4	382
141	Oral Health Disparities in Appalachia. Journal of the American Dental Association, 2008, 139, 598-604.	0.7	14
142	Orbicularis oris muscle defects as an expanded phenotypic feature in nonsyndromic cleft lip with or without cleft palate. American Journal of Medical Genetics, Part A, 2007, 143A, 1143-1149.	0.7	69
143	Subclinical features in non-syndromic cleft lip with or without cleft palate (CL/P): review of the evidence that subepithelial orbicularis oris muscle defects are part of an expanded phenotype for CL/P. Orthodontics and Craniofacial Research, 2007, 10, 82-87.	1.2	49
144	Pittsburgh Registry of Infant Multiplets (PRIM): An Update. Twin Research and Human Genetics, 2006, 9, 1006-1008.	0.3	2

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145	Direct Sequencing of Candidate Genes for Nonsyndromic Cleft Lip and Palate. PLoS Genetics, 2005, preprint, e64.	1.5	1
146	Genome-scan for loci involved in cleft lip with or without cleft palate in consanguineous families from Turkey., 2004, 126A, 111-122.		37
147	Current concepts in the embryology and genetics of cleft lip and cleft palate. Clinics in Plastic Surgery, 2004, 31, 125-140.	0.7	112
148	Meta-Analysis of 13 Genome Scans Reveals Multiple Cleft Lip/Palate Genes with Novel Loci on 9q21 and 2q32-35. American Journal of Human Genetics, 2004, 75, 161-173.	2.6	200
149	Genome Scan for Loci Involved in Cleft Lip With or Without Cleft Palate, in Chinese Multiplex Families. American Journal of Human Genetics, 2002, 71, 349-364.	2.6	107
150	Nonsyndromic Cleft Lip with or without Cleft Palate in China: Assessment of Candidate Regions. Cleft Palate-Craniofacial Journal, 2002, 39, 149-156.	0.5	48
151	Case/control family study of autonomic nervous system dysfunction in idiopathic congenital central hypoventilation syndrome. American Journal of Medical Genetics Part A, 2001, 100, 237-245.	2.4	83
152	Genetic segregation analysis of autonomic nervous system dysfunction in families of probands with idiopathic congenital central hypoventilation syndrome. American Journal of Medical Genetics Part A, 2001, 100, 229-236.	2.4	49
153	Segregation analysis of attention deficit hyperactivity disorder. , 1999, 88, 71-78.		31
154	Changing the definition of ?proband? in the new standardized nomenclature for pedigrees. Journal of Genetic Counseling, 1996, 5, 51-52.	0.9	0
155	Evidence for Autosomal Dominant Inheritance and Raceâ€Specific Heterogeneity in Earlyâ€Onset Periodontitis. Journal of Periodontology, 1994, 65, 623-630.	1.7	177
156	Cleft lip with or without cleft palate: Reanalysis of a three-generation family study from England. Genetic Epidemiology, 1986, 3, 335-342.	0.6	45
157	Linkage analysis and multi-point mapping of 11p markers. Genetic Epidemiology, 1986, 3, 159-164.	0.6	0
158	Linkage analysis of G8 and Huntington's disease. Genetic Epidemiology, 1986, 3, 247-250.	0.6	0
159	Genetic analysis of cleft lip with or without cleft palate in Danish kindreds. American Journal of Medical Genetics Part A, 1984, 19, 9-18.	2.4	82
160	Genetic Etiologies of Facial Clefting. , 0, , 147-161.		12
161	Pittsburgh Registry of Infant Multiplets (PRIM). , 0, .		3