

Alexandre R Vieira

List of Publications by Year in descending order

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344
papers

10,460
citations

41258

49
h-index

49773

87
g-index

364
all docs

364
docs citations

364
times ranked

6994
citing authors

#	ARTICLE	IF	CITATIONS
1	Interferon Regulatory Factor 6 (IRF6) Gene Variants and the Risk of Isolated Cleft Lip or Palate. <i>New England Journal of Medicine</i> , 2004, 351, 769-780.	13.9	534
2	A genome-wide association study of cleft lip with and without cleft palate identifies risk variants near MAFB and ABCA4. <i>Nature Genetics</i> , 2010, 42, 525-529.	9.4	518
3	Interaction of lifestyle, behaviour or systemic diseases with dental caries and periodontal diseases: consensus report of group 2 of the joint <scp>EFP</scp>/<scp>ORCA</scp> workshop on the boundaries between caries and periodontal diseases. <i>Journal of Clinical Periodontology</i> , 2017, 44, S39-S51.	2.3	306
4	Complete sequencing shows a role for MSX1 in non-syndromic cleft lip and palate. <i>Journal of Medical Genetics</i> , 2003, 40, 399-407.	1.5	254
5	Impaired FGF signaling contributes to cleft lip and palate. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 4512-4517.	3.3	246
6	Medical Sequencing of Candidate Genes for Nonsyndromic Cleft Lip and Palate. <i>PLoS Genetics</i> , 2005, 1, e64.	1.5	212
7	Unraveling Human Cleft Lip and Palate Research. <i>Journal of Dental Research</i> , 2008, 87, 119-125.	2.5	209
8	Mutations in BMP4 Are Associated with Subepithelial, Microform, and Overt Cleft Lip. <i>American Journal of Human Genetics</i> , 2009, 84, 406-411.	2.6	176
9	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. <i>Human Molecular Genetics</i> , 2016, 25, dddw104.	1.4	163
10	<i>MSX1, PAX9</i>, and <i>TGFA</i> Contribute to Tooth Agenesis in Humans. <i>Journal of Dental Research</i> , 2004, 83, 723-727.	2.5	158
11	Terminology of Erosive Tooth Wear: Consensus Report of a Workshop Organized by the ORCA and the Cariology Research Group of the IADR. <i>Caries Research</i> , 2020, 54, 2-6.	0.9	155
12	Genes expressed in dental enamel development are associated with molar-incisor hypomineralization. <i>Archives of Oral Biology</i> , 2013, 58, 1434-1442.	0.8	152
13	Enamel Formation Genes Are Associated with High Caries Experience in Turkish Children. <i>Caries Research</i> , 2008, 42, 394-400.	0.9	148
14	A Genome-wide Association Study of Nonsyndromic Cleft Palate Identifies an Etiologic Missense Variant in GRHL3. <i>American Journal of Human Genetics</i> , 2016, 98, 744-754.	2.6	146
15	Defining Subphenotypes for Oral Clefts Based on Dental Development. <i>Journal of Dental Research</i> , 2007, 86, 986-991.	2.5	145
16	Possible Association of <i>Amelogenin</i> to High Caries Experience in a Guatemalan-Mayan Population. <i>Caries Research</i> , 2008, 42, 8-13.	0.9	140
17	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. <i>Human Genetics</i> , 2017, 136, 275-286.	1.8	139
18	Caries: Review of Human Genetics Research. <i>Caries Research</i> , 2014, 48, 491-506.	0.9	127

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19	Explaining Gender Differences in Caries: A Multifactorial Approach to a Multifactorial Disease. <i>International Journal of Dentistry</i> , 2010, 2010, 1-5.	0.5	125
20	Genome-wide Scan Finds Suggestive Caries Loci. <i>Journal of Dental Research</i> , 2008, 87, 435-439.	2.5	123
21	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. <i>Human Heredity</i> , 2009, 68, 151-170.	0.4	113
22	Genome-wide Association Scan for Childhood Caries Implicates Novel Genes. <i>Journal of Dental Research</i> , 2011, 90, 1457-1462.	2.5	108
23	MSX1 and TGFB3 Contribute to Clefting in South America. <i>Journal of Dental Research</i> , 2003, 82, 289-292.	2.5	105
24	Oral Clefts and Syndromic Forms of Tooth Agenesis as Models for Genetics of Isolated Tooth Agenesis. <i>Journal of Dental Research</i> , 2003, 82, 162-165.	2.5	101
25	Dental Anomalies as Part of the Cleft Spectrum. <i>Cleft Palate-Craniofacial Journal</i> , 2008, 45, 414-419.	0.5	97
26	Axis inhibition protein 2 (AXIN2) polymorphisms and tooth agenesis. <i>Archives of Oral Biology</i> , 2009, 54, 45-49.	0.8	95
27	Candidate gene/loci studies in cleft lip/palate and dental anomalies finds novel susceptibility genes for clefts. <i>Genetics in Medicine</i> , 2008, 10, 668-674.	1.1	91
28	Interferon regulatory factor 6 (IRF6) and fibroblast growth factor receptor 1 (FGFR1) contribute to human tooth agenesis. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 538-545.	0.7	89
29	The Antimicrobial Peptide <i>DEFB1</i> Is Associated with Caries. <i>Journal of Dental Research</i> , 2010, 89, 631-636.	2.5	89
30	On the Etiology of Molar-Incisor Hypomineralization. <i>Caries Research</i> , 2016, 50, 166-169.	0.9	86
31	Genetic variation in Myosin 1H contributes to mandibular prognathism. <i>American Journal of Orthodontics and Dentofacial Orthopedics</i> , 2012, 141, 51-59.	0.8	85
32	Genetic Susceptibility to Periapical Disease: Conditional Contribution of MMP2 and MMP3 Genes to the Development of Periapical Lesions and Healing Response. <i>Journal of Endodontics</i> , 2012, 38, 604-607.	1.4	84
33	Enamel Formation Genes Influence Enamel Microhardness Before and After Cariogenic Challenge. <i>PLoS ONE</i> , 2012, 7, e45022.	1.1	82
34	Prevalence of Dental Anomalies in Nonsyndromic Individuals with Cleft Lip and Palate: A Systematic Review and Meta-analysis. <i>Cleft Palate-Craniofacial Journal</i> , 2012, 49, 194-200.	0.5	79
35	Studies with <i>Wnt</i> genes and nonsyndromic cleft lip and palate. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2010, 88, 995-1000.	1.6	78
36	AXIS inhibition protein 2, orofacial clefts and a family history of cancer. <i>Journal of the American Dental Association</i> , 2009, 140, 80-84.	0.7	77

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37	Evidence of genetic variations associated with rotator cuff disease. <i>Journal of Shoulder and Elbow Surgery</i> , 2014, 23, 227-235.	1.2	77
38	Studies of dental anomalies in a large group of school children. <i>Archives of Oral Biology</i> , 2008, 53, 941-946.	0.8	75
39	Association between the Transforming Growth Factor Alpha Gene and Nonsyndromic Oral Clefts: A HuGE Review. <i>American Journal of Epidemiology</i> , 2006, 163, 790-810.	1.6	74
40	<i>AXIN2</i> and <i>CDH1</i> polymorphisms, tooth agenesis, and oral clefts. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2009, 85, 169-173.	1.6	73
41	Genetic variation in MMP20 contributes to higher caries experience. <i>Journal of Dentistry</i> , 2012, 40, 381-386.	1.7	72
42	Early Childhood Caries Is Associated with Genetic Variants in Enamel Formation and Immune Response Genes. <i>Caries Research</i> , 2015, 49, 70-77.	0.9	72
43	PVRL1 variants contribute to non-syndromic cleft lip and palate in multiple populations. <i>American Journal of Medical Genetics, Part A</i> , 2006, 140A, 2562-2570.	0.7	70
44	Role of genetic factors in the pathogenesis of aggressive periodontitis. <i>Periodontology 2000</i> , 2014, 65, 92-106.	6.3	70
45	Genome-wide association Scan of dental caries in the permanent dentition. <i>BMC Oral Health</i> , 2012, 12, 57.	0.8	69
46	Host genetics role in the pathogenesis of periodontal disease and caries. <i>Journal of Clinical Periodontology</i> , 2017, 44, S52-S78.	2.3	68
47	What Is the Heritability of Periodontitis? A Systematic Review. <i>Journal of Dental Research</i> , 2019, 98, 632-641.	2.5	63
48	<i>MMP13</i> Polymorphism Decreases Risk for Dental Caries. <i>Caries Research</i> , 2012, 46, 401-407.	0.9	60
49	A genome wide linkage scan for cleft lip and palate and dental anomalies. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 1406-1413.	0.7	55
50	Elderly at Greater Risk for Root Caries: A Look at the Multifactorial Risks with Emphasis on Genetics Susceptibility. <i>International Journal of Dentistry</i> , 2011, 2011, 1-6.	0.5	53
51	Side of Dental Anomalies and Taurodontism as Potential Clinical Markers for Cleft Subphenotypes. <i>Cleft Palate-Craniofacial Journal</i> , 2011, 48, 103-108.	0.5	52
52	Tooth Agensis Association with Self-reported Family History of Cancer. <i>Journal of Dental Research</i> , 2013, 92, 149-155.	2.5	52
53	<i>TLR4</i> as a risk factor for periodontal disease: a reappraisal. <i>Journal of Clinical Periodontology</i> , 2009, 36, 279-286.	2.3	51
54	Genes Regulating Immune Response and Amelogenesis Interact in Increasing the Susceptibility to Molar-Incisor Hypomineralization. <i>Caries Research</i> , 2019, 53, 217-227.	0.9	50

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55	The PDGF-C regulatory region SNP rs28999109 decreases promoter transcriptional activity and is associated with CL/P. <i>European Journal of Human Genetics</i> , 2009, 17, 774-784.	1.4	48
56	Position and course of the mandibular canal in skulls. <i>Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology</i> , 2012, 113, 453-458.	0.2	48
57	Maternal age and oral clefts: A reappraisal. <i>Oral Surgery Oral Medicine Oral Pathology Oral Radiology and Endodontics</i> , 2002, 94, 530-5.	1.6	46
58	Insights from Studies with Oral Cleft Genes Suggest Associations between WNT-pathway Genes and Risk of Oral Cancer. <i>Journal of Dental Research</i> , 2011, 90, 740-746.	2.5	46
59	Interaction between IRF6 and TGFA Genes Contribute to the Risk of Nonsyndromic Cleft Lip/Palate. <i>PLoS ONE</i> , 2012, 7, e45441.	1.1	46
60	Genetic polymorphisms underlying the skeletal Class III phenotype. <i>American Journal of Orthodontics and Dentofacial Orthopedics</i> , 2017, 151, 700-707.	0.8	45
61	Birth order and oral clefts: A meta analysis. <i>Teratology</i> , 2002, 66, 209-216.	1.8	44
62	Genetic and Environmental Factors in Human Cleft Lip and Palate. <i>Frontiers of Oral Biology</i> , 2012, 16, 19-31.	1.5	44
63	The use of chronic gingivitis as reference status increases the power and odds of periodontitis genetic studies – a proposal based in the exposure concept and clearer resistance and susceptibility phenotypes definition. <i>Journal of Clinical Periodontology</i> , 2012, 39, 323-332.	2.3	42
64	Genome wide association scan for chronic periodontitis implicates novel locus. <i>BMC Oral Health</i> , 2014, 14, 84.	0.8	42
65	A genome-wide linkage scan for cleft lip and cleft palate identifies a novel locus on 8p11-23. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 846-852.	0.7	41
66	Rethinking isolated cleft palate: Evidence of occult lip defects in a subset of cases. <i>American Journal of Medical Genetics, Part A</i> , 2008, 146A, 1670-1675.	0.7	40
67	MMP3 and TIMP1 variants contribute to chronic periodontitis and may be implicated in disease progression. <i>Journal of Clinical Periodontology</i> , 2012, 39, 707-716.	2.3	40
68	The WNT10A gene in ectodermal dysplasias and selective tooth agenesis. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 2455-2460.	0.7	40
69	Spectrum of Dental Phenotypes in Nonsyndromic Orofacial Clefting. <i>Journal of Dental Research</i> , 2015, 94, 905-912.	2.5	40
70	Genetic evidence for the role of loci at 19q13 in cleft lip and palate. <i>Journal of Medical Genetics</i> , 2006, 43, e26-e26.	1.5	38
71	The many faces of the genetics contribution to temporomandibular joint disorder. <i>Orthodontics and Craniofacial Research</i> , 2008, 11, 125-135.	1.2	38
72	Effects of enamel matrix genes on dental caries are moderated by fluoride exposures. <i>Human Genetics</i> , 2015, 134, 159-167.	1.8	38

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73	<i>WDR72</i> Mutations Associated with Amelogenesis Imperfecta and Acidosis. <i>Journal of Dental Research</i> , 2019, 98, 541-548.	2.5	38
74	Interferon regulatory factor 6 (IRF6) is associated with oral-facial cleft in individuals that originate in South America. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 2075-2078.	0.7	37
75	Cleft Lip and Palate in Family Members of Cancer Survivors. <i>Cancer Investigation</i> , 2010, 28, 958-962.	0.6	37
76	Analysis of Multiple Cytokine Polymorphisms in Individuals with Untreated Deep Carious Lesions Reveals IL1B (rs1143643) as a Susceptibility Factor for Periapical Lesion Development. <i>Journal of Endodontics</i> , 2015, 41, 197-200.	1.4	36
77	The identification of peptides by nanoLC-MS/MS from human surface tooth enamel following a simple acid etch extraction. <i>RSC Advances</i> , 2016, 6, 61673-61679.	1.7	36
78	Association studies of low-frequency coding variants in nonsyndromic cleft lip with or without cleft palate. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 1531-1538.	0.7	36
79	Redefining the Phenotype of Dental Caries. <i>Caries Research</i> , 2018, 52, 263-271.	0.9	36
80	A systematic genetic analysis and visualization of phenotypic heterogeneity among orofacial cleft GWAS signals. <i>Genetic Epidemiology</i> , 2019, 43, 704-716.	0.6	36
81	Genetic variants in ACTN3 and MYO1H are associated with sagittal and vertical craniofacial skeletal patterns. <i>Archives of Oral Biology</i> , 2019, 97, 85-90.	0.8	36
82	Caries is Associated with Asthma and Epilepsy. <i>European Journal of Dentistry</i> , 2009, 03, 297-303.	0.8	35
83	Risk of cancer in relatives of children born with isolated cleft lip and palate. <i>American Journal of Medical Genetics, Part A</i> , 2012, 158A, 1503-1504.	0.7	35
84	Role of estrogen related receptor beta (ESRRB) in DFN35B hearing impairment and dental decay. <i>BMC Medical Genetics</i> , 2014, 15, 81.	2.1	35
85	ACTN3 R577X genotypes associate with Class II and deepbite malocclusions. <i>American Journal of Orthodontics and Dentofacial Orthopedics</i> , 2014, 146, 603-611.	0.8	35
86	Dental anomalies in different growth and skeletal malocclusion patterns. <i>Angle Orthodontist</i> , 2018, 88, 195-201.	1.1	35
87	Genetic origins in a South American clefting population. <i>Clinical Genetics</i> , 2002, 62, 458-463.	1.0	34
88	Polymorphisms in BMP4 and FGFR1 genes are associated with fracture non-union. <i>Journal of Orthopaedic Research</i> , 2013, 31, 1971-1979.	1.2	34
89	Genetic influences on dental enamel that impact caries differ between the primary and permanent dentitions. <i>European Journal of Oral Sciences</i> , 2015, 123, 327-334.	0.7	33
90	Oral manifestations in coronavirus disease 2019 (COVID-19). <i>Oral Diseases</i> , 2021, 27, 770-770.	1.5	33

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91	Whorl patterns on the lower lip are associated with nonsyndromic cleft lip with or without cleft palate. <i>American Journal of Medical Genetics, Part A</i> , 2009, 149A, 2673-2679.	0.7	32
92	Association of <i>MMP3</i> and <i>TIMP2</i> promoter polymorphisms with nonsyndromic oral clefts. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2012, 94, 540-548.	1.6	32
93	Fine-Mapping of 5q12.1-13.3 Unveils New Genetic Contributors to Caries. <i>Caries Research</i> , 2013, 47, 273-283.	0.9	31
94	Enamel Formation Genes Associated with Dental Erosive Wear. <i>Caries Research</i> , 2015, 49, 236-242.	0.9	31
95	Association of <i>AXIN2</i> with Non-syndromic Oral Clefts in Multiple Populations. <i>Journal of Dental Research</i> , 2012, 91, 473-478.	2.5	29
96	On the Variable Clinical Presentation of Molar-Incisor Hypomineralization. <i>Caries Research</i> , 2019, 53, 482-488.	0.9	29
97	Genetic Association of <i>MPPED2</i> and <i>ACTN2</i> with Dental Caries. <i>Journal of Dental Research</i> , 2014, 93, 626-632.	2.5	28
98	<i>CRISPLD2</i> Variants Including a C471T Silent Mutation May Contribute to Nonsyndromic Cleft Lip with or without Cleft Palate. <i>Cleft Palate-Craniofacial Journal</i> , 2011, 48, 363-370.	0.5	27
99	Genetic variation in <i>Ameloblastin</i> is associated with caries in asthmatic children. <i>European Archives of Paediatric Dentistry: Official Journal of the European Academy of Paediatric Dentistry</i> , 2014, 15, 211-216.	0.7	27
100	COVID-19-Related Challenges in Dental Education: Experiences From Brazil, the USA, and Australia. <i>Pesquisa Brasileira Em Odontopediatria E Clinica Integrada</i> , 2020, 20, .	0.7	27
101	Defining Predictors of Cleft Lip and Palate Risk. <i>Journal of Dental Research</i> , 2012, 91, 556-561.	2.5	26
102	Role of <i>TRAV</i> locus in low caries experience. <i>Human Genetics</i> , 2013, 132, 1015-1025.	1.8	26
103	Genetic variation in the promoter region of beta-defensin 1 (<i>DEFB 1</i>) is associated with high caries experience in children born with cleft lip and palate. <i>Acta Odontologica Scandinavica</i> , 2014, 72, 235-240.	0.9	26
104	Heat Shock 70 Protein Genes and Genetic Susceptibility to Apical Periodontitis. <i>Journal of Endodontics</i> , 2016, 42, 1467-1471.	1.4	26
105	Aggressive periodontitis is likely influenced by a few small effect genes. <i>Journal of Clinical Periodontology</i> , 2009, 36, 468-473.	2.3	25
106	Follow-up association studies of chromosome region 9q and nonsyndromic cleft lip/palate. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 1701-1710.	0.7	25
107	Saving More Teeth—A Case for Personalized Care. <i>Journal of Personalized Medicine</i> , 2015, 5, 30-35.	1.1	25
108	Weaker Dental Enamel Explains Dental Decay. <i>PLoS ONE</i> , 2015, 10, e0124236.	1.1	25

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109	MMP20 rs1784418 Protects Certain Populations against Caries. Caries Research, 2017, 51, 46-51.	0.9	25
110	Profiling microorganisms in whole saliva of children with and without dental caries. Clinical and Experimental Dental Research, 2019, 5, 438-446.	0.8	25
111	Gene-environment interaction in molar-incisor hypomineralization. PLoS ONE, 2021, 16, e0241898.	1.1	25
112	Polymorphisms in Nonamelogenin Enamel Matrix Genes Are Associated with Dental Fluorosis. Caries Research, 2018, 52, 1-6.	0.9	25
113	Genetic variation may explain why females are less susceptible to dental erosion. European Journal of Oral Sciences, 2016, 124, 426-432.	0.7	24
114	Identification of 16q21 as a modifier of nonsyndromic orofacial cleft phenotypes. Genetic Epidemiology, 2017, 41, 887-897.	0.6	24
115	Novel Cleft Susceptibility Genes in Chromosome 6q. Journal of Dental Research, 2010, 89, 927-932.	2.5	23
116	FAM5C Contributes to Aggressive Periodontitis. PLoS ONE, 2010, 5, e10053.	1.1	23
117	Caries experience and overall health status. Oral Health & Preventive Dentistry, 2014, 12, 163-70.	0.3	23
118	Birth order and neural tube defects: a reappraisal. Journal of the Neurological Sciences, 2004, 217, 65-72.	0.3	22
119	Candidate Genes for Oral-Facial Clefts in Guatemalan Families. Annals of Plastic Surgery, 2006, 56, 518-521.	0.5	22
120	Evidence of linkage disequilibrium between polymorphisms at the IRF6 locus and isolate tooth agenesis, in a Turkish population. Archives of Oral Biology, 2008, 53, 780-784.	0.8	22
121	ENPP1 and ESR1 genotypes influence temporomandibular disorders development and surgical treatment response in dentofacial deformities. Journal of Cranio-Maxillo-Facial Surgery, 2016, 44, 1226-1237.	0.7	22
122	Genes Involved in the Enamel Development Are Associated with Calcium and Phosphorus Level in Saliva. Caries Research, 2017, 51, 225-230.	0.9	22
123	Aquaporin 5 Interacts with Fluoride and Possibly Protects against Caries. PLoS ONE, 2015, 10, e0143068.	1.1	22
124	Fluoride uptake and release by composites and glass ionomers in a high caries challenge situation. American Journal of Dentistry, 1999, 12, 14-8.	0.1	22
125	Reduced folate carrier 1 (RFC1) is associated with cleft of the lip only. Brazilian Journal of Medical and Biological Research, 2008, 41, 689-693.	0.7	21
126	BMP4 and FGF3 haplotypes increase the risk of tendinopathy in volleyball athletes. Journal of Science and Medicine in Sport, 2015, 18, 150-155.	0.6	21

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127	MMP1 and MMP20 contribute to tooth agenesis in humans. Archives of Oral Biology, 2011, 56, 506-511.	0.8	20
128	Molecular motor MYO1C, acetyltransferase KAT6B and osteogenetic transcription factor RUNX2 expression in human masseter muscle contributes to development of malocclusion. Archives of Oral Biology, 2014, 59, 601-607.	0.8	20
129	A Pragmatic Study Shows Failure of Dental Composite Fillings Is Genetically Determined: A Contribution to the Discussion on Dental Amalgams. Frontiers in Medicine, 2017, 4, 186.	1.2	20
130	Studies of genes in the <i>FGF</i> signaling pathway and oral clefts with or without dental anomalies. American Journal of Medical Genetics, Part A, 2008, 146A, 1614-1617.	0.7	19
131	Family history of cleft lip and palate in subjects diagnosed with leukemia. American Journal of Medical Genetics, Part A, 2012, 158A, 678-679.	0.7	19
132	Functional Significance of <i>MMP3</i> and <i>TIMP2</i> Polymorphisms in Cleft Lip/Palate. Journal of Dental Research, 2014, 93, 651-656.	2.5	19
133	Rethinking isolated cleft lip and palate as a syndrome. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, 2018, 125, 307-312.	0.2	19
134	Two-fold excess of fluoride in the drinking water has no obvious health effects other than dental fluorosis. Journal of Trace Elements in Medicine and Biology, 2018, 50, 216-222.	1.5	19
135	Dental Anomalies in Children Born with Clefts: A Case-Control Study. Cleft Palate-Craniofacial Journal, 2012, 49, 64-68.	0.5	18
136	Candidate gene studies in hypodontia suggest role for FGF3. European Archives of Paediatric Dentistry: Official Journal of the European Academy of Paediatric Dentistry, 2013, 14, 405-410.	0.7	18
137	Fine mapping of locus Xq25.1-27-2 for a low caries experience phenotype. Archives of Oral Biology, 2014, 59, 479-486.	0.8	18
138	Colorectal Cancer-Associated Genes Are Associated with Tooth Agenesis and May Have a Role in Tooth Development. Scientific Reports, 2018, 8, 2979.	1.6	18
139	The association of genetic polymorphisms in serotonin transporter and catecholâ€methyltransferase on temporomandibular disorders and anxiety in adolescents. Journal of Oral Rehabilitation, 2019, 46, 597-604.	1.3	18
140	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
141	Estrogen receptor gene is associated with dental fluorosis in Brazilian children. Clinical Oral Investigations, 2019, 23, 3565-3570.	1.4	18
142	On the genetics contribution to molar incisor hypomineralization. International Journal of Paediatric Dentistry, 2019, 29, 2-3.	1.0	18
143	Caries is Associated with Asthma and Epilepsy. European Journal of Dentistry, 2009, 3, 297-303.	0.8	18
144	Genetic mapping of high caries experience on human chromosome 13. BMC Medical Genetics, 2013, 14, 116.	2.1	17

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145	Different contribution of BRINP3 gene in chronic periodontitis and peri-implantitis: a cross-sectional study. BMC Oral Health, 2015, 15, 33.	0.8	17
146	Association between TNF α - 308 G/A polymorphism and oral lichen planus (OLP): a meta-analysis. Journal of Applied Oral Science, 2018, 26, e20170184.	0.7	16
147	Oestrogen receptor alpha, growth hormone receptor, and developmental defect of enamel. International Journal of Paediatric Dentistry, 2019, 29, 29-35.	1.0	16
148	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
149	Genome-Wide Association Study (GWAS) of dental caries in diverse populations. BMC Oral Health, 2021, 21, 377.	0.8	16
150	ENPP1 and ESR1 genotypes associated with subclassifications of craniofacial asymmetry and severity of temporomandibular disorders. American Journal of Orthodontics and Dentofacial Orthopedics, 2017, 152, 631-645.	0.8	15
151	Association Between Polymorphisms in the Genes of Estrogen Receptors and the Presence of Temporomandibular Disorders and Chronic Arthralgia. Journal of Oral and Maxillofacial Surgery, 2018, 76, 314.e1-314.e9.	0.5	15
152	Genome-wide interaction studies identify sex-specific risk alleles for nonsyndromic orofacial clefts. Genetic Epidemiology, 2018, 42, 664-672.	0.6	15
153	MMP13 Contributes to Dental Caries Associated with Developmental Defects of Enamel. Caries Research, 2019, 53, 441-446.	0.9	15
154	Measuring the Microscopic Structures of Human Dental Enamel Can Predict Caries Experience. Journal of Personalized Medicine, 2020, 10, 5.	1.1	15
155	Transforming growth factor α gene (<i>TGFA</i>), human tooth agenesis, and evidence of segmental uniparental isodisomy. European Journal of Oral Sciences, 2009, 117, 20-26.	0.7	14
156	In Vitro Acid-Mediated Initial Dental Enamel Loss Is Associated with Genetic Variants Previously Linked to Caries Experience. Frontiers in Physiology, 2017, 8, 104.	1.3	14
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