Azeez Butali

List of Publications by Year
in descending order

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61 papers	1,362 citations	430874 18 h-index	395702 33 g-index
64 all docs	64 docs citations	64 times ranked	1410 citing authors

#	Article	IF	Citations
1	Variant analyses of candidate genes in orofacial clefts in multiâ€ethnic populations. Oral Diseases, 2022, 28, 1921-1935.	3.0	3
2	Genetic and epigenetic studies in nonâ€syndromic oral clefts. Oral Diseases, 2022, 28, 1339-1350.	3.0	16
3	Genomeâ€wide association study of multiethnic nonsyndromic orofacial cleft families identifies novel loci specific to family and phenotypic subtypes. Genetic Epidemiology, 2022, , .	1.3	4
4	Whole-genome sequencing reveals de-novo mutations associated with nonsyndromic cleft lip/palate. Scientific Reports, 2022, 12, .	3.3	11
5	Correlation Between Height and Impacted Third Molars and Genetics Role in Third Molar Impaction. Journal of Maxillofacial and Oral Surgery, 2021, 20, 149-153.	1.4	3
6	Coâ€occurrence of orofacial clefts and clubfoot phenotypes in a subâ€Saharan African cohort: Wholeâ€exome sequencing implicates multiple syndromes and genes. Molecular Genetics & Genomic Medicine, 2021, 9, e1655.	1.2	3
7	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.	3.7	16
8	Replication of GWAS significant loci in a sub-Saharan African Cohort with early childhood caries: a pilot study. BMC Oral Health, 2021, 21, 274.	2.3	3
9	Variant Analyses of Candidate Genes in Orofacial Clefts in Multiâ€Ethnic Populations. FASEB Journal, 2021, 35, .	0.5	0
10	Missense Variants Within GJB2 Gene Locus and the Risk of Hearing Defects in Nonsyndromic Cleft Lip and Palate. Journal of Craniofacial Surgery, 2021, Publish Ahead of Print, e676-e678.	0.7	1
11	The Influence of Sex and Ancestry on Three-Dimensional Palate Shape. Journal of Craniofacial Surgery, 2021, 32, 2883-2887.	0.7	1
12	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.	0.9	1
13	MTHFR promoter methylation might mitigate the effect of smoking at the level of LINE â€1 in cleft lip tissues: A preliminary study. Birth Defects Research, 2021, 113, 1463-1469.	1.5	0
14	Assessing the Practice of Birth Defect Registration at Addis Ababa Health Facilities. Ethiopian Journal of Health Sciences, 2021, 31, 683-687.	0.4	0
15	The Role of Environmental Factors in the Etiology of Nonsyndromic Orofacial Clefts. Journal of Craniofacial Surgery, 2020, 31, 113-116.	0.7	8
16	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.	3.8	19
17	Six2 regulates Pax9 expression, palatogenesis and craniofacial bone formation. Developmental Biology, 2020, 458, 246-256.	2.0	13
18	Nonâ€random distribution of deleterious mutations in the DNA and proteinâ€binding domains of IRF6 are associated with Van Der Woude syndrome. Molecular Genetics & Genomic Medicine, 2020, 8, e1355.	1.2	13

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19	SPECC1L regulates palate development downstream of IRF6. Human Molecular Genetics, 2020, 29, 845-858.	2.9	18
20	Breastfeeding Practices Among Mothers of Children With Orofacial Clefts in an African Cohort. Cleft Palate-Craniofacial Journal, 2020, 57, 1018-1023.	0.9	5
21	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.	0.7	5
22	Management of orofacial cleft in Nigeria - A retrospective study. Annals of Maxillofacial Surgery, 2020, 10, 434.	0.7	8
23	Descriptive epidemiology of salivary gland neoplasms in Nigeria: An AOPRC multicenter tertiary hospital study. Oral Diseases, 2019, 25, 142-149.	3.0	5
24	Missense Pathogenic variants in KIF4A Affect Dental Morphogenesis Resulting in X-linked Taurodontism, Microdontia and Dens-Invaginatus. Frontiers in Genetics, 2019, 10, 800.	2.3	7
25	A systematic genetic analysis and visualization of phenotypic heterogeneity among orofacial cleft GWAS signals. Genetic Epidemiology, 2019, 43, 704-716.	1.3	36
26	A multi-centre evaluation of 566 cases of ameloblastoma in Nigeria by the African Oral Pathology Research Consortium. Oral Cancer, 2019, 3, 9-15.	0.3	1
27	Automated phone call and text reminders for childhood immunisations (PRIMM): a randomised controlled trial in Nigeria. BMJ Global Health, 2019, 4, e001232.	4.7	23
28	Genomic analyses in African populations identify novel risk loci for cleft palate. Human Molecular Genetics, 2019, 28, 1038-1051.	2.9	61
29	Molecular Screening of VAX1 Gene Polymorphisms Uncovered the Genetic Heterogeneity of Nonsyndromic Orofacial Cleft Among Saudi Arabian Patients. Genetic Testing and Molecular Biomarkers, 2019, 23, 45-50.	0.7	6
30	A Preexperimental Study to Assess the Impact of an Interdisciplinary Educational Intervention on Nurses' Knowledge of Perinatal and Infant Oral Health Care. Journal of International Society of Preventive and Community Dentistry, 2019, 9, 619-629.	1.0	3
31	Novel GREM1 Variations in Sub-Saharan African Patients With Cleft Lip and/or Cleft Palate. Cleft Palate-Craniofacial Journal, 2018, 55, 736-742.	0.9	9
32	Geriatric dentistry education and context in a selection of countries in 5 continents. Special Care in Dentistry, 2018, 38, 123-132.	0.8	37
33	Oral Healthâ€“Related Quality of Life of Children Born With Orofacial Clefts in Ethiopia and Their Parents. Cleft Palate-Craniofacial Journal, 2018, 55, 1153-1157.	0.9	11
34	A comparative study of immediate wound healing complications following cleft lip repair using either absorbable or non-absorbable skin sutures. Journal of the Korean Association of Oral and Maxillofacial Surgeons, 2018, 44, 159.	0.8	9
35	Genomeâ€wide interaction studies identify sexâ€specific risk alleles for nonsyndromic orofacial clefts. Genetic Epidemiology, 2018, 42, 664-672.	1.3	15
36	Identification of paternal uniparental disomy on chromosome 22 and a de novo deletion on chromosome 18 in individuals with orofacial clefts. Molecular Genetics & Genomic Medicine, 2018, 6, 924-932.	1.2	4

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37	Unilateral cleft lip repair: a comparison of treatment outcome with two surgical techniques using quantitative (anthropometry) assessment. Journal of the Korean Association of Oral and Maxillofacial Surgeons, 2018, 44, 3.	0.8	21
38	Six2 regulates palate development by inhibiting palatal bone formation during development. FASEB Journal, 2018, 32, 15.1.	0.5	0
39	The dentist-scientist career pathway in Africa: opportunities and obstacles. Korean Journal of Medical Education, 2018, 30, 189-198.	1.3	6
40	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.	3.8	139
41	Genetics and genomics etiology of nonsyndromic orofacialÂclefts. Molecular Genetics & Genomic Medicine, 2017, 5, 3-7.	1.2	13
42	The prevalence, penetrance, and expressivity of etiologic <scp>IRF</scp>6 variants in orofacial clefts patients from subâ€Saharan Africa. Molecular Genetics & Genomic Medicine, 2017, 5, 164-171.	1.2	9
43	Exome sequencing provides additional evidence for the involvement of ARHGAP29 in Mendelian orofacial clefting and extends the phenotypic spectrum to isolated cleft palate. Birth Defects Research, 2017, 109, 27-37.	1.5	49
44	Identification of 16q21 as a modifier of nonsyndromic orofacial cleft phenotypes. Genetic Epidemiology, 2017, 41, 887-897.	1.3	24
45	Omics-based molecular techniques in oral pathology centred cancer: prospect and challenges in Africa. Cancer Cell International, 2017, 17, 61.	4.1	7
46	A comparative study of quality of life of families with children born with cleft lip and/or palate before and after surgical treatment. Journal of the Korean Association of Oral and Maxillofacial Surgeons, 2017, 43, 247.	0.8	25
47	A multi-centre evaluation of oral cancer in Southern and Western Nigeria: an African oral pathology research consortium initiative. Pan African Medical Journal, 2017, 28, 64.	0.8	9
48	Characteristics and risk factors of preterm births in a tertiary center in Lagos, Nigeria. Pan African Medical Journal, 2016, 24, 1.	0.8	73
49	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.	2.9	163
50	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.	6.2	146
51	Congenital Palatal Fistula Associated with Submucous Cleft Palate. Plastic and Reconstructive Surgery - Global Open, 2016, 4, e613.	0.6	5
52	Irf6 directly regulates Klf17 in zebrafish periderm and Klf4 in murine oral epithelium, and dominant-negative KLF4 variants are present in patients with cleft lip and palate. Human Molecular Genetics, 2016, 25, 766-776.	2.9	48
53	Cleft lip and palate: Parental experiences of stigma, discrimination, and social/structural inequalities. Annals of Maxillofacial Surgery, 2016, 6, 195.	0.7	39
54	Multidisciplinary approach to genomics research in Africa: the AfriCRAN model. Pan African Medical Journal, 2015, 21, 229.	0.8	6

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55	Foreign Bodies Simulating a Congenital Palatal Fistula and Vascular Anomaly. Case Reports in Dentistry, 2015, 2015, 1-4.	0.5	1
56	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.	1.2	31
57	Novel IRF6 mutations in families with Van Der Woude syndrome and popliteal pterygium syndrome from subâ€Saharan Africa. Molecular Genetics & Genomic Medicine, 2014, 2, 254-260.	1.2	24
58	Rare functional variants in genomeâ€“wide association identified candidate genes for nonsyndromic clefts in the African population. American Journal of Medical Genetics, Part A, 2014, 164, 2567-2571.	1.2	35
59	Replication of Genome Wide Association Identified Candidate Genes Confirm the Role of Common and Rare Variants in <scp>PAX</scp>7 and <scp>VAX</scp>1 in the Etiology of Nonsyndromic CL(P). American Journal of Medical Genetics, Part A, 2013, 161, 965-972.	1.2	60
60	Folic acid supplementation use and the MTHFR C677T polymorphism in orofacial clefts etiology: An individual participant data pooledâ€analysis. Birth Defects Research Part A: Clinical and Molecular Teratology, 2013, 97, 509-514.	1.6	46
61	Amniotic band syndrome associated with extremely severe atypical clefts of the orofacial region. European Journal of Plastic Surgery, 0, , 1.	0.6	3