Sven Kreiborg

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/72688/publications.pdf

Version: 2024-02-01

41 papers	1,530 citations	516710 16 h-index	330143 37 g-index
41 all docs	41 docs citations	41 times ranked	1497 citing authors

#	Article	IF	Citations
1	Facial Asymmetry in Nonsyndromic and Muenke Syndromeâ€“Associated Unicoronal Synostosis: A 3-Dimensional Study Based on Facial Surfaces Extracted From CT Scans. Cleft Palate-Craniofacial Journal, 2021, 58, 687-696.	0.9	1
2	Dental Subphenotypes in Infants With Orofacial Cleftsâ€”A Longitudinal Population-Based Retrospective Radiographic Study of the Primary and Secondary Dentitions. Cleft Palate-Craniofacial Journal, 2021, 58, 1526-1535.	0.9	2
3	Comparison of 3â€D mandibular surfaces generated by MRI and CT. Orthodontics and Craniofacial Research, 2021, , .	2.8	2
4	Diffusion-weighted magnetic resonance imaging of the oral and maxillofacial region: optimal fat suppression method. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, 2021, 131, 738-745.	0.4	1
5	Three-dimensional assessment of facial morphology in children and adolescents with juvenile idiopathic arthritis and moderate to severe TMJ involvement using 3D surface scans. Clinical Oral Investigations, 2020, 24, 799-807.	3.0	10
6	Phenotypic variability in Muenke syndromeâ€”observations from five Danish families. Clinical Dysmorphology, 2020, 29, 1-9.	0.3	6
7	Description of a family with Xâ€linked oculoâ€auriculoâ€vertebral spectrum associated with polyalanine tract expansion in <scp>ZIC3</scp>. Clinical Genetics, 2020, 98, 384-389.	2.0	11
8	Deep orofacial phenotyping of population-based infants with isolated cleft lip and isolated cleft palate. Scientific Reports, 2020, 10, 21666.	3.3	3
9	Delayed maturation and reduced crown width of the permanent first mandibular molar in all subgroups of cleft lip and palate. Orthodontics and Craniofacial Research, 2020, 23, 432-438.	2.8	2
10	Longterm Outcomes of Temporomandibular Joints in Juvenile Idiopathic Arthritis: 17 Years of Followup of a Nordic Juvenile Idiopathic Arthritis Cohort. Journal of Rheumatology, 2020, 47, 730-738.	2.0	34
11	Novel de novo mutation in ZBTB20 in primrose syndrome in boy with short stature. Clinical Dysmorphology, 2019, 28, 41-45.	0.3	7
12	Spatially Detailed 3D Quantification of Improved Facial Symmetry After Surgery in Children With Unicoronal Synostosis. Cleft Palate-Craniofacial Journal, 2019, 56, 918-928.	0.9	8
13	A complex phenotype in a family with a pathogenic SOX3 missense variant. European Journal of Medical Genetics, 2018, 61, 168-172.	1.3	12
14	Accuracy and precision of manual segmentation of the maxillary sinus in MR imagesâ€”a method study. British Journal of Radiology, 2018, 91, 20170663.	2.2	14
15	Neonatal hyperinsulinemic hypoglycemia in a patient with 9p deletion syndrome. European Journal of Medical Genetics, 2018, 61, 473-477.	1.3	4
16	Dentinogenesis imperfecta type <scp>II</scp>â€genotype and phenotype analyses in three Danish families. Molecular Genetics & Genomic Medicine, 2018, 6, 339-349.	1.2	11
17	A study of familial Char syndrome involving the TFAP2B gene with a focus on facial shape characteristics. Clinical Dysmorphology, 2018, 27, 71-77.	0.3	5
18	Tooth formation and eruption â€“ lessons learnt from cleidocranial dysplasia. European Journal of Oral Sciences, 2018, 126, 72-80.	1.5	42

#	Article	IF	Citations
19	Parental age in relation to the severity of cleft lip and/or palate. Orthodontics and Craniofacial Research, 2018, 21, 236-241.	2.8	13
20	Stereoscopy in Dental Education: An Investigation. Journal of Dental Education, 2017, 81, 450-457.	1.2	3
21	Automatic measurement of orbital volume in unilateral coronal synostosis. , 2016, , .		1
22	A new implementation of digital X-ray radiogrammetry and reference curves of four indices of cortical bone for healthy European adults. Archives of Osteoporosis, 2016, 11, 17.	2.4	8
23	Phenotypic subregions within the split-hand/foot malformation 1 locus. Human Genetics, 2016, 135, 345-357.	3.8	15
24	Facial morphology in children and adolescents with juvenile idiopathic arthritis and moderate to severe temporomandibular joint involvement. American Journal of Orthodontics and Dentofacial Orthopedics, 2016, 149, 182-191.	1.7	29
25	3D digital surface imaging for quantification of facial development and asymmetry in juvenile idiopathic arthritis. Seminars in Orthodontics, 2015, 21, 121-124.	1.4	2
26	3D analysis of facial asymmetry in subjects with juvenile idiopathic arthritis. Rheumatology, 2011, 50, 586-592.	1.9	26
27	Automated quantification and analysis of facial asymmetry in children with arthritis in the temporomandibular joint. , 2011, , .		2
28	Ocular Manifestations of Apert and Crouzon Syndromes. Journal of Craniofacial Surgery, 2010, 21, 1354-1357.	0.7	76
29	Automated quantification and analysis of mandibular asymmetry. , 2010, , .		2
30	Orofacial pain, jaw function, and temporomandibular disorders in women with a history of juvenile chronic arthritis or persistent juvenile chronic arthritis. Oral Surgery Oral Medicine Oral Pathology Oral Radiology and Endodontics, 2001, 92, 406-414.	1.4	53
31	Mutations in the gene encoding c-Abl-binding protein SH3BP2 cause cherubism. Nature Genetics, 2001, 28, 125-126.	21.4	341
32	Suture formation, premature sutural fusion, and suture default zones in Apert syndrome. American Journal of Medical Genetics Part A, 1996, 62, 339-344.	2.4	47
33	FIRST-TRIMESTER PRENATAL DIAGNOSIS OF CROUZON SYNDROME. , 1996, 16, 155-158.		25
34	Hands and feet in the Apert syndrome. American Journal of Medical Genetics Part A, 1995, 57, 82-96.	2.4	88
35	Cutaneous manifestations of Apert syndrome. American Journal of Medical Genetics Part A, 1995, 58, 94-96.	2.4	66
36	Visceral anomalies in the Apert syndrome. American Journal of Medical Genetics Part A, 1993, 45, 758-760.	2.4	128

#	Article	IF	Citations
37	Growth pattern in the Apert syndrome. American Journal of Medical Genetics Part A, 1993, 47, 617-623.	2.4	44
38	Cranial base and face in mandibulofacial dysostosis. American Journal of Medical Genetics Part A, 1993, 47, 753-760.	2.4	30
39	Skeletal abnormalities in the Apert syndrome. American Journal of Medical Genetics Part A, 1993, 47, 624-632.	2.4	81
40	Upper and lower airway compromise in the apert syndrome. American Journal of Medical Genetics Part A, 1992, 44, 90-93.	2.4	89
41	Development of the dentition in cleidocranial dysplasia. Journal of Oral Pathology and Medicine, 1990, 19, 89-93.	2.7	186