Peter Anderson

List of Publications by Year
in descending order

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60 papers	1,265 citations	361045 20 h-index	377514 34 g-index
60 all docs	60 docs citations	60 times ranked	1703 citing authors

#	Article	IF	Citations
1	Proboscis Lateralis With Basal Encephalocele: A Report of Clinical Management and Reconstructive Approach. Cleft Palate-Craniofacial Journal, 2023, 60, 1331-1336.	0.5	1
2	Childhood Experiences and Perspectives of Individuals With Orofacial Clefts: A Qualitative Systematic Review. Cleft Palate-Craniofacial Journal, 2022, , 105566562210845.	0.5	1
3	Ex Vivo Culture of Human Cranial Suture Cells. Methods in Molecular Biology, 2022, 2403, 215-222.	0.4	0
4	Editorial: Genetic, Environmental and Synergistic Gene-Environment Contributions to Craniofacial Defects. Frontiers in Cell and Developmental Biology, 2022, 10, 887051.	1.8	0
5	Cognitive, behavioral and psychological functioning of children and adults with conservatively managed metopic synostosis. Child Neuropsychology, 2021, 27, 190-208.	0.8	4
6	Delineating the roles of Grhl2 in craniofacial development through tissueâ€specific conditional deletion and epistasis approaches in mouse. Developmental Dynamics, 2021, 250, 1191-1209.	0.8	2
7	Incidence of persistent metopic suture in Australia: findings from 1034 three-dimensional computed tomography scans. Child's Nervous System, 2021, 37, 3871-3879.	0.6	2
8	Secondary Surgery in Metopic Craniosynostosis. Journal of Craniofacial Surgery, 2021, Publish Ahead of Print, 539-542.	0.3	1
9	Neo-Sagittal Suture Formation After Cranial Vault Remodeling in Sagittal Craniosynostosis. Journal of Craniofacial Surgery, 2021, 32, 282-284.	0.3	3
10	The Normal Fetal Cephalic Index in the Second and Third Trimesters of Pregnancy. Ultrasound Quarterly, 2020, 36, 255-262.	0.3	4
11	Interrogating the Grainyhead-like 2 (Grhl2) genomic locus identifies an enhancer element that regulates palatogenesis in mouse. Developmental Biology, 2020, 459, 194-203.	0.9	7
12	HOPX regulates bone marrow-derived mesenchymal stromal cell fate determination via suppression of adipogenic gene pathways. Scientific Reports, 2020, 10, 11345.	1.6	11
13	Pharmacological targeting of KDM6A and KDM6B, as a novel therapeutic strategy for treating craniosynostosis in Saethre-Chotzen syndrome. Stem Cell Research and Therapy, 2020, 11, 529.	2.4	12
14	Sonographic indicators of isolated fetal sagittal craniosynostosis during pregnancy. Journal of Medical Imaging and Radiation Oncology, 2020, 64, 626-633.	0.9	2
15	Vessel-derived angiocrine IGF-1 promotes Meckel's cartilage proliferation to drive jaw growth during embryogenesis. Development (Cambridge), 2020, 147, .	1.2	10
16	Panoramic radiography is of limited value in the evaluation of maxillary sinus disease. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, 2019, 127, 237-246.	0.2	21
17	Cognitive, behavioral and psychological functioning in children with metopic synostosis: a meta-analysis examining the impact of surgical status. Child Neuropsychology, 2019, 25, 263-277.	0.8	21
18	Tyrosine kinase receptor c-ros-oncogene 1 inhibition alleviates aberrant bone formation of TWIST-1 haploinsufficient calvarial cells from Saethre-Chotzen syndrome patients. Journal of Cellular Physiology, 2018, 233, 7320-7332.	2.0	8

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19	YY1 Haploinsufficiency Causes an Intellectual Disability Syndrome Featuring Transcriptional and Chromatin Dysfunction. American Journal of Human Genetics, 2017, 100, 907-925.	2.6	125
20	The prevalence of dental anomalies in an Australian population. Australian Dental Journal, 2017, 62, 161-164.	0.6	20
21	Tyrosine kinase receptor c-ros-oncogene 1 mediates TWIST-1 regulation of human mesenchymal stem cell lineage commitment. Bone, 2017, 94, 98-107.	1.4	13
22	The use of obstetric ultrasound in the antenatal diagnosis of craniosynostosis: We need to do better. Australasian Journal of Ultrasound in Medicine, 2016, 19, 91-98.	0.3	4
23	Feeding and reflux in children after mandibular distraction osteogenesis for micrognathia: A systematic review. International Journal of Pediatric Otorhinolaryngology, 2016, 85, 128-135.	0.4	29
24	The ophthalmic sequelae of Pfeiffer syndrome and the long-term visual outcomes after craniofacial surgery. Journal of AAPOS, 2016, 20, 315-319.	0.2	14
25	Mice lacking the conserved transcription factor Grainyhead-like 3 (Grhl3) display increased apposition of the frontal and parietal bones during embryonic development. BMC Developmental Biology, 2016, 16, 37.	2.1	17
26	Flightless I is a key regulator of the fibroproliferative process in hypertrophic scarring and a target for a novel antiscarring therapy. British Journal of Dermatology, 2016, 174, 786-794.	1.4	18
27	Does the Rate of Distraction or Type of Distractor Affect the Outcome of Mandibular Distraction in Children With Micrognathia?. Journal of Oral and Maxillofacial Surgery, 2016, 74, 1441-1453.	0.5	20
28	Neural crest cell-derived VEGF promotes embryonic jaw extension. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 6086-6091.	3.3	54
29	Application of threeâ€dimensional computed tomography in craniofacial clinical practice and research. Australian Dental Journal, 2014, 59, 174-185.	0.6	57
30	A complicated case of plagiocephaly followed by delayed craniosynostosis. Child's Nervous System, 2013, 29, 1395-1396.	0.6	3
31	Muencke Syndrome With Cleft Lip and Palate. Journal of Craniofacial Surgery, 2013, 24, 1484-1485.	0.3	6
32	Twist-1 Induces Ezh2 Recruitment Regulating Histone Methylation along the Ink4A/Arf Locus in Mesenchymal Stem Cells. Molecular and Cellular Biology, 2012, 32, 1433-1441.	1.1	106
33	Heterozygous Mutations of FREM1 Are Associated with an Increased Risk of Isolated Metopic Craniosynostosis in Humans and Mice. PLoS Genetics, 2011, 7, e1002278.	1.5	80
34	A New Syndrome With Craniosynostosis and Cleft Lip and Palate. Journal of Craniofacial Surgery, 2011, 22, 122-124.	0.3	0
35	Intracranial volume measurement of sagittal craniosynostosis. Journal of Clinical Neuroscience, 2007, 14, 455-458.	0.8	29
36	Unravelling the molecular control of calvarial suture fusion in children with craniosynostosis. BMC Genomics, 2007, 8, 458.	1.2	84

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37	Scanning Electron Microscope and Micro-CT Evaluation of Cranial Sutures in Health and Disease. Journal of Craniofacial Surgery, 2006, 17, 909-919.	0.3	13
38	Do Craniosynostosis Syndrome Phenotypes with Both FGFR2 and TWIST Mutations have a Worse Clinical Outcome?. Journal of Craniofacial Surgery, 2006, 17, 166-172.	0.3	2
39	Spinal Anomalies in Goldenhar Syndrome. Cleft Palate-Craniofacial Journal, 2005, 42, 477-480.	0.5	43
40	Late Results After Unicoronal Craniosynostosis Correction. Journal of Craniofacial Surgery, 2005, 16, 37-44.	0.3	28
41	Simultaneous multiple vector distraction for craniosynostosis syndromes. Journal of Plastic, Reconstructive and Aesthetic Surgery, 2005, 58, 626-631.	1.1	33
42	Craniosynostosis and childbirth. European Journal of Plastic Surgery, 2005, 28, 94-98.	0.3	9
43	Familial Parryâ€“Romberg disease. International Journal of Pediatric Otorhinolaryngology, 2005, 69, 705-708.	0.4	35
44	Apocrine Hydrocysadenoma of the Ear. Otolaryngology - Head and Neck Surgery, 2005, 133, 981-982.	1.1	13
45	Hyperostosis as a late sequel of parasymphyseal mandibular fractures in 2 children. Journal of Cranio-Maxillo-Facial Surgery, 2005, 33, 188-190.	0.7	4
46	Management of facial dysmorphogenesis in nemaline myopathy: a case report. World Journal of Orthodontics, 2005, 6, 156-60.	0.2	4
47	Analysis of Intracranial Volume in Apert Syndrome Genotypes. Pediatric Neurosurgery, 2004, 40, 161-164.	0.4	39
48	Intracranial Volume Measurement of Metopic Craniosynostosis. Journal of Craniofacial Surgery, 2004, 15, 1014-1016.	0.3	35
49	Modified costochondral graft osteotomy in hemifacial microsomia. Journal of Plastic, Reconstructive and Aesthetic Surgery, 2003, 56, 414-415.	1.1	4
50	Teratomas of the head and neck region. Journal of Cranio-Maxillo-Facial Surgery, 2003, 31, 369-377.	0.7	35
51	Assessment of various parameters in the estimation of differential renal function using technetium-99m mercaptoacetyltriglycine. European Journal of Nuclear Medicine and Molecular Imaging, 1999, 26, 155-162.	3.3	31
52	The Feet in Apert's Syndrome. Journal of Pediatric Orthopaedics, 1999, 19, 504-507.	0.6	8
53	Re: Metacarpophalangeal analysis in Crouzon syndrome. , 1998, 80, 439-439.		5
54	Assessment of drainage in PUJ dilatation. Nuclear Medicine Communications, 1997, 18, 823-826.	0.5	27

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55	Anomalous venous drainage in a case of non-syndromic craniosynostosis. Child's Nervous System, 1997, 13, 97-100.	0.6	38
56	Hand anomalies in Crouzon syndrome. Skeletal Radiology, 1997, 26, 113-115.	1.2	24
57	The feet in Crouzon syndrome. Journal of Craniofacial Genetics and Developmental Biology, 1997, 17, 43-7.	0.1	4
58	Finger Duplication in Apert's Syndrome. Journal of Hand Surgery, 1996, 21, 649-651.	0.9	4
59	The hands in Saethre-Chotzen syndrome. Journal of Craniofacial Genetics and Developmental Biology, 1996, 16, 228-33.	0.1	4
60	Effect of renal maturation on the clearance of technetium-99m mercaptoacetyltriglycine. European Journal of Nuclear Medicine and Molecular Imaging, 1994, 21, 1333-1337.	2.2	34