Alonso, Lg ; Alonso, L

List of Publications by Year
in descending order

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23 papers	1,238 citations	687363 13 h-index	677142 22 g-index
24 all docs	24 docs citations	24 times ranked	1375 citing authors

#	Article	IF	Citations
1	Analysis by Light, Scanning, and Transmission Microscopy of the Intima Synovial of the Temporomandibular Joint of Human Fetuses during the Development. Anatomy Research International, 2014, 2014, 1-6.	1.1	8
2	Ring chromosome 10: report on two patients and review of the literature. Journal of Applied Genetics, 2013, 54, 35-41.	1.9	10
3	Morphometric Analysis of the Internal Auditory Canal by Computed Tomography Imaging. Iranian Journal of Radiology, 2012, 9, 71-78.	0.2	28
4	Toriello Carey syndrome: genetic, clinical, and oral considerations: a case report. Special Care in Dentistry, 2011, 31, 68-72.	0.8	4
5	Immunohistochemical expression of types I and III collagen antibodies in the temporomandibular joint disc of human foetuses. European Journal of Histochemistry, 2011, 55, e24.	1.5	16
6	High frequency of submicroscopic chromosomal imbalances in patients with syndromic craniosynostosis detected by a combined approach of microsatellite segregation analysis, multiplex ligation-dependent probe amplification and array-based comparative genome hybridisation. Journal of Medical Genetics, 2008, 45, 447-450.	3.2	36
7	Anatomic and Dynamic Aspects of Stomatognathic Structures in Osteogenesis Imperfecta: A Case Report. Cranio - Journal of Craniomandibular Practice, 2007, 25, 144-149.	1.4	3
8	Radiological Findings and Dynamic Aspects of Stomatognathic Structures in Treacher Collins Syndrome: Clinical Case Report. Cleft Palate-Craniofacial Journal, 2007, 44, 678-682.	0.9	8
9	Morphological analysis of the vestibular aqueduct by computerized tomography images. European Journal of Radiology, 2007, 61, 79-83.	2.6	12
10	RAB23 Mutations in Carpenter Syndrome Imply an Unexpected Role for Hedgehog Signaling in Cranial-Suture Development and Obesity. American Journal of Human Genetics, 2007, 80, 1162-1170.	6.2	229
11	Mutational Screening of FGFR1, CER1, and CDON in a Large Cohort of Trigonocephalic Patients. Cleft Palate-Craniofacial Journal, 2006, 43, 148-151.	0.9	15
12	Temporomandibular joint disc position and configuration in children with functional unilateral posterior crossbite: A magnetic resonance imaging evaluation. American Journal of Orthodontics and Dentofacial Orthopedics, 2006, 129, 785-793.	1.7	22
13	Further evidence of association between mutations inFGFR2 and syndromic craniosynostosis with sacrococcygeal eversion. Birth Defects Research Part A: Clinical and Molecular Teratology, 2006, 76, 629-633.	1.6	15
14	Molecular screening for microdeletions at 9p22â€p24 and 11q23â€q24 in a large cohort of patients with trigonocephaly. Clinical Genetics, 2005, 67, 503-510.	2.0	48
15	Functional unilateral posterior crossbite effects on mastication movements using axiography. Angle Orthodontist, 2005, 75, 362-7.	2.4	12
16	Tetraploid/diploid mosaicism: case report and review of the literature. Annales De GÃ©nÃ©tique, 2002, 45, 177-180.	0.4	7
17	Clinical and cytogenetic characterisation of a patient with Down syndrome resulting from a 21q22.1->qter duplication. Journal of Medical Genetics, 2001, 38, 73-76.	3.2	7
18	High mutation detection rate inTCOF1 among Treacher Collins syndrome patients reveals clustering of mutations and 16 novel pathogenic changes. Human Mutation, 2000, 16, 315-322.	2.5	112

#	Article	IF	Citations
19	CACP, encoding a secreted proteoglycan, is mutated in camptodactyly-arthropathy-coxa vara-pericarditis syndrome. Nature Genetics, 1999, 23, 319-322.	21.4	286
20	Clinical spectrum of fibroblast growth factor receptor mutations. Human Mutation, 1999, 14, 115-125.	2.5	284
21	Clinical spectrum of fibroblast growth factor receptor mutations. Human Mutation, 1999, 14, 115.	2.5	35
22	Description of a new mutation and characterization ofFGFR1, FGFR2, andFGFR3 mutations among Brazilian patients with syndromic craniosynostoses. American Journal of Medical Genetics Part A, 1998, 78, 237-241.	2.4	41
23	Description of a new mutation and characterization of FGFR1, FGFR2, and FGFR3 mutations among Brazilian patients with syndromic craniosynostoses. American Journal of Medical Genetics Part A, 1998, 78, 237-241.	2.4	0