

Edi LÃ³cia Sartorato

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

259
citations

933447

10
h-index

940533

16
g-index

21
all docs

21
docs citations

21
times ranked

322
citing authors

#	ARTICLE	IF	CITATIONS
1	Prevalence of the GJB2 mutations and the del(GJB6-D13S1830) mutation in Brazilian patients with deafness. <i>Hearing Research</i> , 2004, 196, 87-93.	2.0	33
2	Searching for Digenic Inheritance in Deaf Brazilian Individuals Using the Multiplex Ligation-Dependent Probe Amplification Technique. <i>Genetic Testing and Molecular Biomarkers</i> , 2011, 15, 849-853.	0.7	31
3	Determination of the frequency of the 35delG allele in Brazilian neonates. <i>Clinical Genetics</i> , 2000, 58, 339-340.	2.0	29
4	Molecular genetics study of deafness in Brazil: 8-year experience. <i>American Journal of Medical Genetics, Part A</i> , 2007, 143A, 1574-1579.	1.2	24
5	Screening of genetic alterations related to non-syndromic hearing loss using MassARRAY iPLEX® technology. <i>BMC Medical Genetics</i> , 2015, 16, 85.	2.1	23
6	Optimization of simultaneous screening of the main mutations involved in non-syndromic deafness using the TaqMan® OpenArray, Genotyping Platform. <i>BMC Medical Genetics</i> , 2013, 14, 112.	2.1	21
7	Molecular analysis of SLC26A4 gene in patients with nonsyndromic hearing loss and EVA: Identification of two novel mutations in Brazilian patients. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2013, 77, 410-413.	1.0	18
8	Analysis of mitochondrial alterations in Brazilian patients with sensorineural hearing loss using MALDI-TOF mass spectrometry. <i>BMC Medical Genetics</i> , 2016, 17, 41.	2.1	14
9	Screening for the <i>GJB2</i> c.-3170 G>A (IVS 1+1 G>A) Mutation in Brazilian Deaf Individuals Using Multiplex Ligation-Dependent Probe Amplification. <i>Genetic Testing and Molecular Biomarkers</i> , 2009, 13, 701-704.	0.7	12
10	Molecular study of patients with auditory neuropathy. <i>Molecular Medicine Reports</i> , 2016, 14, 481-490.	2.4	11
11	Single Nucleotide Polymorphisms of the GJB2 and GJB6 Genes Are Associated with Autosomal Recessive Nonsyndromic Hearing Loss. <i>BioMed Research International</i> , 2015, 2015, 1-8.	1.9	9
12	A investigaÃ§Ã£o genÃ©tica na surdez hereditÃ¡ria nÃ£o-sindrÃ´mica. <i>Revista Brasileira De Otorrinolaringologia</i> , 2004, 70, 182-186.	0.2	9
13	Molecular investigation in children candidates and submitted to cochlear implantation. <i>Brazilian Journal of Otorhinolaryngology</i> , 2006, 72, 333-336.	1.0	6
14	Correlation between audiometric data and the 35delG mutation in ten patients. <i>Brazilian Journal of Otorhinolaryngology</i> , 2007, 73, 777-783.	1.0	5
15	Multiplex MALDI-TOF MS detection of mitochondrial variants in Brazilian patients with hereditary optic neuropathy. <i>Molecular Vision</i> , 2016, 22, 1024-35.	1.1	5
16	CorrelaÃ§Ã£o entre dados audiomÃ©tricos e mutaÃ§Ã£o 35delG em dez pacientes. <i>Revista Brasileira De Otorrinolaringologia</i> , 2007, 73, 777-783.	0.2	2
17	Interaction between audiology and genetics in the study of a family: the complexity of molecular diagnosis and genetic counseling. <i>Brazilian Journal of Otorhinolaryngology</i> , 2008, 74, 698-702.	1.0	2
18	Differentiating Leber Hereditary Optic Neuropathy from Normal-Tension Glaucoma. <i>Neuro-Ophthalmology</i> , 2017, 41, 99-102.	1.0	2

#	ARTICLE	IF	CITATIONS
19	Estudo molecular em crianÃ§as candidatas e submetidas ao implante coclear. Revista Brasileira De Otorrinolaringologia, 2006, 72, 333-336.	0.2	2
20	An auditory health program for neonates in ICU and/or intermediate care settings. Brazilian Journal of Otorhinolaryngology, 2013, 79, 709-715.	1.0	1
21	InteraÃ§Ã£o entre audiologia e genÃ©tica no estudo de uma famÃlia: a complexidade do diagnÃstico molecular e do aconselhamento genÃ©tico. Revista Brasileira De Otorrinolaringologia, 2008, 74, 698-702.	0.2	0