Elio Marciano

List of Publications by Year
in descending order

43 papers	1,490 citations	394421 19 h-index	315739 38 g-index
43 all docs	43 docs citations	43 times ranked	2062 citing authors

#	Article	IF	Citations
1	Performance and characteristics of the Newborn Hearing Screening Program in Campania region (Italy) between 2013 and 2019. European Archives of Oto-Rhino-Laryngology, 2022, 279, 1221-1231.	1.6	5
2	NeonaTal Assisted TelerehAbilitation (T.A.T.A. Web App) for Hearing-Impaired Children: A Family-Centered Care Model for Early Intervention in Congenital Hearing Loss. Audiology Research, 2022, 12, 182-190.	1.8	5
3	Novel USH1G homozygous variant underlying USH2-like phenotype of Usher syndrome. European Journal of Ophthalmology, 2021, 31, NP18-NP22.	1.3	4
4	Targeted Audiological Surveillance Program in Campania, Italy. Indian Pediatrics, 2021, 58, 441-444.	0.4	4
5	Integrated Bimodal Fitting for Unilateral CI Users with Residual Contralateral Hearing. Audiology Research, 2021, 11, 200-206.	1.8	2
6	The Natural History of Hearing Disorders in Asymptomatic Congenital Cytomegalovirus Infection. Frontiers in Pediatrics, 2020, 8, 217.	1.9	9
7	Prevalence of macular abnormalities assessed by optical coherence tomography in patients with Usher syndrome. Ophthalmic Genetics, 2018, 39, 17-21.	1.2	10
8	Voice Disorder Detection via an m-Health System: Design and Results of a Clinical Study to Evaluate Vox4Health. BioMed Research International, 2018, 2018, 1-19.	1.9	23
9	Dialogic reading in the rehabilitation of Children with Hearing Loss andÂthe â€œBorn to Readâ€•Project: A pilot study. Scandinavian Journal of Psychology, 2018, 59, 518-523.	1.5	3
10	Genetic characterization of Italian patients with Bardet-Biedl syndrome and correlation to ocular, renal and audio-vestibular phenotype: identification of eleven novel pathogenic sequence variants. BMC Medical Genetics, 2017, 18, 10.	2.1	59
11	Novel compound heterozygous mutations in BCS1L gene causing Bjornstad syndrome in two siblings. , 2017, 173, 1348-1352.		7
12	SLC26A4 genotypes associated with enlarged vestibular aqueduct malformation in south Italian children with sensorineural hearing loss. Clinical Chemistry and Laboratory Medicine, 2016, 54, e259-63.	2.3	2
13	Phenotypic and genetic characterization of a family carrying two Xq21.1-21.3 interstitial deletions associated with syndromic hearing loss. Molecular Cytogenetics, 2015, 8, 18.	0.9	19
14	Exclusion of TNFRSF11B as Candidate Gene for Otosclerosis in Campania Population. Indian Journal of Otolaryngology and Head and Neck Surgery, 2014, 66, 297-301.	0.9	3
15	Very good performance with bimodal stimulation in a like-hybrid modality in a patient with profound bilateral sensorineural hearing loss with low-frequencies preservation. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2014, 35, 70-72.	1.3	0
16	Abnormal thalamic function in patients with vestibular migraine. Neurology, 2014, 82, 2120-2126.	1.1	114
17	Feasibility and effectiveness of a population-based newborn hearing screening in an economically deprived region of Italy. International Journal of Pediatric Otorhinolaryngology, 2013, 77, 329-333.	1.0	19
18	Hearing impairment in Parkinson's disease: Expanding the nonmotor phenotype. Movement Disorders, 2012, 27, 1530-1535.	3.9	93

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19	The development of language in babies and the role of the family. Journal of Maternal-Fetal and Neonatal Medicine, 2011, 24, 120-121.	1.5	2
20	Mutational analysis forGJB2, GJB6, andGJB3genes in Campania within a universal neonatal hearing screening programme. International Journal of Audiology, 2011, 50, 866-870.	1.7	8
21	Vestibular impairment and adaptive postural imbalance in parkinsonian patients with lateral trunk flexion. Movement Disorders, 2011, 26, 1458-1463.	3.9	75
22	GJB2 Gene Mutations in Syndromic Skin Diseases with Sensorineural Hearing Loss.. Current Genomics, 2011, 12, 475-485.	1.6	42
23	R75Q dominant mutation in GJB2 gene silenced by the in cis recessive mutation c.35delG. American Journal of Medical Genetics, Part A, 2010, 152A, 2658-2660.	1.2	6
24	Vestibular Pathways Involvement in Children With Migraine: A Neuroâ€Otological Study. Headache, 2010, 50, 71-76.	3.9	39
25	Screening forGJB2andGJB6gene mutations in patients from Campania region with sensorineural hearing loss. International Journal of Audiology, 2010, 49, 326-331.	1.7	8
26	Functional dissection of auditory cortex with magnetic resonance imaging. Audiological Medicine, 2010, 8, 88-99.	0.4	0
27	Age-related hearing loss in four Italian genetic isolates: An epidemiological study. International Journal of Audiology, 2009, 48, 465-472.	1.7	17
28	New evidence for the correlation of the p.G130V mutation in the GJB2 gene and syndromic hearing loss with palmoplantar keratoderma. American Journal of Medical Genetics, Part A, 2009, 149A, 685-688.	1.2	22
29	Spatio-temporal pattern of vestibular information processing after brief caloric stimulation. European Journal of Radiology, 2009, 70, 312-316.	2.6	69
30	Independent component model of the default-mode brain function: combining individual-level and population-level analyses in resting-state fMRI. Magnetic Resonance Imaging, 2008, 26, 905-913.	1.8	126
31	Neuro-otological features of Benign Paroxysmal Vertigo and Benign Paroxysmal Positioning Vertigo in children: A follow-up study. Brain and Development, 2006, 28, 80-84.	1.1	46
32	Identification of a Novel Mutation in the Myosin VIIA Motor Domain in a Family with Autosomal Dominant Hearing Loss (DFNA11). Audiology and Neuro-Otology, 2006, 11, 157-164.	1.3	25
33	Trigeminal Stimulation Elicits a Peripheral Vestibular Imbalance in Migraine Patients. Headache, 2005, 45, 325-331.	3.9	101
34	Audiometric evaluation of carriers of the connexin 26 mutation 35delG. European Archives of Oto-Rhino-Laryngology, 2005, 262, 921-924.	1.6	14
35	Functional characterization of a novel Cx26 (T55N) mutation associated to non-syndromic hearing loss. Biochemical and Biophysical Research Communications, 2005, 337, 799-805.	2.1	27
36	Paraoxonase and Superoxide Dismutase Gene Polymorphisms and Noise-Induced Hearing Loss. Clinical Chemistry, 2004, 50, 2012-2018.	3.2	100

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37	Down-regulation of otospiralin mRNA in response to acoustic stress in guinea pig. Hearing Research, 2004, 198, 36-40.	2.0	5
38	FMRI of the auditory system: understanding the neural basis of auditory gestalt. Magnetic Resonance Imaging, 2003, 21, 1213-1224.	1.8	33
39	Otosclerosis: exclusion of linkage to the OTSC1 and OTSC2 loci in four Italian families: Otoesclerosis: exclusiÃ³n de enlaces entre los loci OTSC1 y OTSC2 en cuatro familias italianas. International Journal of Audiology, 2003, 42, 475-480.	1.7	5
40	Effect over time of allopurinol on noise-induced hearing loss in guinea pigs: Efecto en el tiempo del alopurinol sobre la hipoacusia inducida por ruido en cobayos. International Journal of Audiology, 2003, 42, 227-234.	1.7	16
41	Postural restrictions in labyrintholithiasis. European Archives of Oto-Rhino-Laryngology, 2002, 259, 262-265.	1.6	49
42	Functional Fields in Human Auditory Cortex Revealed by Time-Resolved fMRI without Interference of EPI Noise. NeuroImage, 2001, 13, 328-338.	4.2	51
43	Quality of Life Determinants and Hearing Function in an Elderly Population: Osservatorio Geriatrico Campano Study Group. Gerontology, 1999, 45, 323-328.	2.8	223