Carla Laria

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/7907302/publications.pdf

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17	239	1163117	996975
papers	citations	8	15
		h-index	g-index

17	17	17	337
all docs	docs citations	times ranked	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	Tinnitus and equilibrium disorders in COVID-19 patients: preliminary results. European Archives of Oto-Rhino-Laryngology, 2021, 278, 3725-3730.	1.6	82
4	Sudden olfactory loss as an early marker of COVID-19: a nationwide Italian survey. European Archives of Oto-Rhino-Laryngology, 2021, 278, 247-255.	1.6	15
5	Targeted Audiological Surveillance Program in Campania, Italy. Indian Pediatrics, 2021, 58, 441-444.	0.4	4
6	Tinnitus and Neuropsychological Dysfunction in the Elderly: A Systematic Review on Possible Links. Journal of Clinical Medicine, 2021, 10, 1881.	2.4	6
7	Integrated Bimodal Fitting for Unilateral CI Users with Residual Contralateral Hearing. Audiology Research, 2021, 11, 200-206.	1.8	2
8	The role of endogenous Antisecretory Factor (AF) in the treatment of MÃ©niÃ¨re's Disease: A two-year follow-up study. Preliminary results. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2020, 41, 102673.	1.3	12
9	Bimodal strategy for excellent audiological rehabilitation in a subject with a novel nonsense mutation of the SLC26A4 gene: A case report. International Journal of Pediatric Otorhinolaryngology, 2020, 134, 110018.	1.0	1
10	X-Linked Sensorineural Hearing Loss: A Literature Review. Current Genomics, 2018, 19, 327-338.	1.6	35
11	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
12	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
13	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
14	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
15	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
16	Screening forGJB2andGJB6gene mutations in patients from Campania region with sensorineural hearing loss. International Journal of Audiology, 2010, 49, 326-331.	1.7	8
17	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