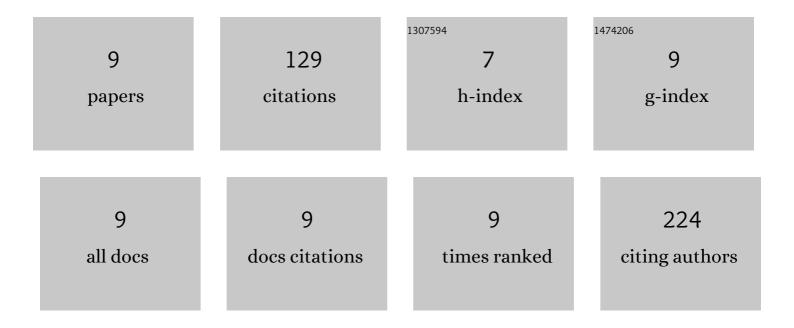
Micol Busi

List of Publications by Year in descending order

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MICOL RUSI

#	Article	IF	CITATIONS
1	The universal newborn hearing screening program at the University Hospital of Ferrara: Focus on costs and software solutions. International Journal of Pediatric Otorhinolaryngology, 2008, 72, 807-816.	1.0	23
2	LAMM syndrome with middle ear dysplasia associated with compound heterozygosity for FGF3 mutations. American Journal of Medical Genetics, Part A, 2011, 155, 1096-1101.	1.2	23
3	Cochlear Implant Outcomes and Genetic Mutations in Children with Ear and Brain Anomalies. BioMed Research International, 2015, 2015, 1-19.	1.9	20
4	Association of the 4 g/5 g polymorphism of plasminogen activator inhibitor-1 gene with sudden sensorineural hearing loss. A case control study. BMC Ear, Nose and Throat Disorders, 2012, 12, 5.	2.6	16
5	Association between idiopathic hearing loss and mitochondrial DNA mutations: A study on 169 hearing-impaired subjects. International Journal of Molecular Medicine, 2013, 32, 785-794.	4.0	16
6	Novel mutations in the SLC26A4 gene. International Journal of Pediatric Otorhinolaryngology, 2012, 76, 1249-1254.	1.0	9
7	Syndromic hearing loss: An update. Hearing, Balance and Communication, 2013, 11, 146-159.	0.4	9
8	Hearing threshold assessment in young children with electrocochleograpy (EcochG) and auditory brainstem responses (ABR): Experience at the University Hospital of Ferrara. Auris Nasus Larynx, 2010, 37, 553-557.	1.2	7
9	Audiological profiles and gjb2, gjb6 mutations: A retrospective study on genetic and clinical data from 2003 to 2008. Audiological Medicine, 2009, 7, 93-105.	0.4	6