# Micol Busi

List of Publications by Year
in descending order



| 9 | 129 | 1307594 7 | 1474206 9 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 9 | 9 | 9 | 224 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Cochlear Implant Outcomes and Genetic Mutations in Children with Ear and Brain Anomalies. BioMed Research International, 2015, 2015, 1-19. | 1.9 | 20 |
| 2 | Syndromic hearing loss: An update. Hearing, Balance and Communication, 2013, 11, 146-159. | 0.4 | 9 |
| 3 | 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 |
| 4 | Novel mutations in the SLC26A4 gene. International Journal of Pediatric Otorhinolaryngology, 2012, 76, 1249-1254. | 1.0 | 9 |
| 5 | Association of the 4â€‰g/5â€‰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 |
| 6 | 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 |
| 7 | 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 |
| 8 | 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 |
| 9 | 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 |