# Natalie Loundon

List of Publications by Year
in descending order



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|---|---|---|---|
| 46 papers | 1,495 citations | 236925 25 h-index | 315739 38 g-index |
| 46 all docs | 46 docs citations | 46 times ranked | 1831 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | GJB2 and GJB6 Mutations. JAMA Otolaryngology, 2005, 131, 481. | 1.2 | 93 |
| 2 | Medical and Surgical Complications in Pediatric Cochlear Implantation. JAMA Otolaryngology, 2010, 136, 12. | 1.2 | 88 |
| 3 | International Pediatric Otolaryngology Group (IPOG) consensus recommendations: Hearing loss in the pediatric patient. International Journal of Pediatric Otorhinolaryngology, 2016, 90, 251-258. | 1.0 | 88 |
| 4 | European Bilateral Pediatric Cochlear Implant Forum Consensus Statement. Otology and Neurotology, 2012, 33, 561-565. | 1.3 | 79 |
| 5 | Reliability of the Language ENvironment Analysis system (LENAâ„¢) in European French. Behavior Research Methods, 2016, 48, 1109-1124. | 4.0 | 75 |
| 6 | Temperature-sensitive auditory neuropathy associated with an otoferlin mutation: Deafening fever!. Biochemical and Biophysical Research Communications, 2010, 394, 737-742. | 2.1 | 74 |
| 7 | Evaluation of Cytomegalovirus (CMV) DNA Quantification in Dried Blood Spots: Retrospective Study of CMV Congenital Infection. Journal of Clinical Microbiology, 2007, 45, 3804-3806. | 3.9 | 59 |
| 8 | Usher Syndrome and Cochlear Implantation. Otology and Neurotology, 2003, 24, 216-221. | 1.3 | 55 |
| 9 | Cochlear implantation and vestibular function in children. International Journal of Pediatric Otorhinolaryngology, 2015, 79, 101-104. | 1.0 | 53 |
| 10 | Clinical evidence of the nonpathogenic nature of the M34T variant in the connexin 26 gene. European Journal of Human Genetics, 2004, 12, 279-284. | 2.8 | 48 |
| 11 | Screening of SLC26A4, FOXI1 and KCNJ10 genes in unilateral hearing impairment with ipsilateral enlarged vestibular aqueduct. International Journal of Pediatric Otorhinolaryngology, 2010, 74, 1049-1053. | 1.0 | 46 |
| 12 | An Application of NGS for Molecular Investigations in Perrault Syndrome: Study of 14 Families and Review of the Literature. Human Mutation, 2016, 37, 1354-1362. | 2.5 | 46 |
| 13 | Unilateral Sensorineural Hearing Loss: Medical Context and Etiology. Audiology and Neuro-Otology, 2017, 22, 83-88. | 1.3 | 43 |
| 14 | Alu-mediated deletion of SOX10 regulatory elements in Waardenburg syndrome type 4. European Journal of Human Genetics, 2012, 20, 990-994. | 2.8 | 37 |
| 15 | Cochlear Implant Failure and Revision Surgery in Pediatric Population. Annals of Otology, Rhinology and Laryngology, 2015, 124, 227-231. | 1.1 | 37 |
| 16 | Velopharyngoplasty for Noncleft Velopharyngeal Insufficiency. JAMA Otolaryngology, 2009, 135, 652. | 1.2 | 34 |
| 17 | EDNRB mutations cause Waardenburg syndrome type II in the heterozygous state. Human Mutation, 2017, 38, 581-593. | 2.5 | 33 |
| 18 | New Closed Skin Bone-Anchored Implant. Otology and Neurotology, 2013, 34, 275-281. | 1.3 | 32 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Mycophenolate mofetil as a treatment of steroid dependent Cogan's syndrome in childhood. International Journal of Pediatric Otorhinolaryngology, 2009, 73, 1477-1479. | 1.0 | 31 |
| 20 | Congenital Cytomegalovirus Is the Second Most Frequent Cause of Bilateral Hearing Loss in Young French Children. Journal of Pediatrics, 2013, 162, 593-599. | 1.8 | 31 |
| 21 | Mutations in TUBB4B Cause a Distinctive Sensorineural Disease. American Journal of Human Genetics, 2017, 101, 1006-1012. | 6.2 | 30 |
| 22 | A preliminary study of computer assisted evaluation of congenital tracheal stenosis: A new tool for surgical decision-making. International Journal of Pediatric Otorhinolaryngology, 2012, 76, 1552-1557. | 1.0 | 29 |
| 23 | Hearing rehabilitation with the closed skin bone-anchored implant Sophono Alpha1: Results of a prospective study in 15 children with ear atresia. International Journal of Pediatric Otorhinolaryngology, 2015, 79, 382-387. | 1.0 | 29 |
| 24 | TheGJB2 mutation R75Q can cause nonsyndromic hearing loss DFNA3 or hereditary palmoplantar keratoderma with deafness. American Journal of Medical Genetics, Part A, 2005, 137A, 225-227. | 1.2 | 27 |
| 25 | Pediatric Cochlear Implantation in Residual Hearing Candidates. Annals of Otology, Rhinology and Laryngology, 2015, 124, 443-451. | 1.1 | 27 |
| 26 | Cochlear implant and inner ear malformation. International Journal of Pediatric Otorhinolaryngology, 2008, 72, 541-547. | 1.0 | 26 |
| 27 | Audiophonological results after cochlear implantation in 40 congenitally deaf patients: preliminary results. International Journal of Pediatric Otorhinolaryngology, 2000, 56, 9-21. | 1.0 | 25 |
| 28 | Stapedectomy in Children. JAMA Otolaryngology, 2010, 136, 1005. | 1.2 | 25 |
| 29 | Pain After Cochlear Implantation: An Unusual Complication?. Otology and Neurotology, 2017, 38, 956-961. | 1.3 | 21 |
| 30 | Head and neck infantile myofibromatosis â€" a report of three cases. International Journal of Pediatric Otorhinolaryngology, 1999, 51, 181-186. | 1.0 | 20 |
| 31 | Benefits from upgrade to the CP810â„¢ sound processor for NucleusÂ® 24 cochlear implant recipients. European Archives of Oto-Rhino-Laryngology, 2014, 271, 49-57. | 1.6 | 19 |
| 32 | Usher syndrome type 1: Early detection of electroretinographic changes. European Journal of Paediatric Neurology, 2009, 13, 505-507. | 1.6 | 17 |
| 33 | Results of VSB implantation at the short process of the incus in children with ear atresia. International Journal of Pediatric Otorhinolaryngology, 2017, 93, 83-87. | 1.0 | 17 |
| 34 | Intra-cochlear electrode tip fold-over. Cochlear Implants International, 2018, 19, 225-229. | 1.2 | 16 |
| 35 | Hearing Outcomes in Functional Surgery for Middle Ear Malformations. Otology and Neurotology, 2013, 34, 1417-1420. | 1.3 | 13 |
| 36 | Cochlear Implantation and Congenital Deafness. Otology and Neurotology, 2012, 33, 539-544. | 1.3 | 11 |

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|---|---|---|---|
| 37 | Evolution of the Bacteriologic Features of Persistent Acute Otitis Media Compared With Acute Otitis Media. JAMA Otolaryngology, 1999, 125, 1134. | 1.2 | 10 |
| 38 | Discovery of a Large Deletion of KAL1 in 2 Deaf Brothers. Otology and Neurotology, 2013, 34, 1590-1594. | 1.3 | 9 |
| 39 | Efficiency of Melatonin as Compared to Pentobarbital for Audiometry Brainstem Response in Children With Associated Disorders. American Journal of Audiology, 2016, 25, 206-210. | 1.2 | 9 |
| 40 | A case of mild CHARGE syndrome associated with a splice site mutation in CHD7. European Journal of Medical Genetics, 2016, 59, 195-197. | 1.3 | 7 |
| 41 | High prevalence of congenital deafness on Reunion Island is due to a founder variant of LHFPL5. Clinical Genetics, 2019, 95, 177-181. | 2.0 | 7 |
| 42 | Noonan Syndrome: An Underestimated Cause of Severe to Profound Sensorineural Hearing Impairment. Which Clues to Suspect the Diagnosis?. Otology and Neurotology, 2017, 38, 1081-1084. | 1.3 | 6 |
| 43 | The Oticon Medical Neuro Zti cochlear implant and the Neuro 2 sound processor: multicentric evaluation of outcomes in adults and children. International Journal of Audiology, 2020, 59, 153-160. | 1.7 | 5 |
| 44 | The role of computed tomography and magnetic resonance imaging for preoperative pediatric cochlear implantation work-up in academic institutions. Cochlear Implants International, 2021, 22, 96-102. | 1.2 | 5 |
| 45 | Arterial spin labeling brain MRI study to evaluate the impact of deafness on cerebral perfusion in 79 children before cochlear implantation. NeuroImage: Clinical, 2021, 29, 102510. | 2.7 | 3 |
| 46 | Congenital Cytomegalovirus Is the Second Most Frequent Cause of Bilateral Hearing Loss in Young French Children. Obstetrical and Gynecological Survey, 2013, 68, 501-503. | 0.4 | 0 |