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**Mutation in transcription factor POU4F3 associated with inherited progressive hearing loss in humans**

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299	Mutations in a novel cochlear gene cause DFNA9, a human nonsyndromic deafness with vestibular dysfunction. <b>1998</b> , 20, 299-303		286
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296	Single gene influences on radiologically-detectable malformations of the inner ear. <b>1998</b> , 31, 391-408; quiz 409-10		12
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