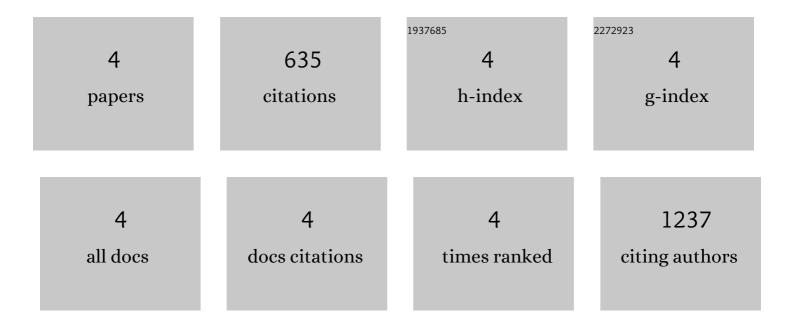
## Fernando Mayo-Merino

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/341858/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Insights into the pathophysiology of DFNA10 hearing loss associated with novel EYA4 variants. Scientific Reports, 2020, 10, 6213.	3.3	15
2	DFNA8/12 caused by TECTA mutations is the most identified subtype of nonsyndromic autosomal dominant hearing loss. Human Mutation, 2011, 32, 825-834.	2.5	73
3	In vivo and in vitro effects of two novel gamma-actin (ACTG1) mutations that cause DFNA20/26 hearing impairment. Human Molecular Genetics, 2009, 18, 3075-3089.	2.9	64
4	Mutations in the seed region of human miR-96 are responsible for nonsyndromic progressive hearing loss. Nature Genetics, 2009, 41, 609-613.	21.4	483