

# Fernando Mayo-Merino

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/341858/publications.pdf>

Version: 2024-02-01

4  
papers

635  
citations

1937685

4  
h-index

2272923

4  
g-index

4  
all docs

4  
docs citations

4  
times ranked

1237  
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutations in the seed region of human miR-96 are responsible for nonsyndromic progressive hearing loss. <i>Nature Genetics</i> , 2009, 41, 609-613.	21.4	483
2	DFNA8/12 caused by TECTA mutations is the most identified subtype of nonsyndromic autosomal dominant hearing loss. <i>Human Mutation</i> , 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. <i>Human Molecular Genetics</i> , 2009, 18, 3075-3089.	2.9	64
4	Insights into the pathophysiology of DFNA10 hearing loss associated with novel EYA4 variants. <i>Scientific Reports</i> , 2020, 10, 6213.	3.3	15