

# Federica Isidori

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/447086/publications.pdf>

Version: 2024-02-01

9  
papers

99  
citations

1937685  
4  
h-index

1474206  
9  
g-index

9  
all docs

9  
docs citations

9  
times ranked

254  
citing authors

#	ARTICLE	IF	CITATIONS
1	A New Homozygous CACNB2 Mutation has Functional Relevance and Supports a Role for Calcium Channels in Autism Spectrum Disorder. <i>Journal of Autism and Developmental Disorders</i> , 2021, 51, 377-381.	2.7	5
2	Biallelic variants in <i>LIG3</i> cause a novel mitochondrial neurogastrointestinal encephalomyopathy. <i>Brain</i> , 2021, 144, 1451-1466.	7.6	28
3	Targeted Sequencing of Sorted Esophageal Adenocarcinoma Cells Unveils Known and Novel Mutations in the Separated Subpopulations. <i>Clinical and Translational Gastroenterology</i> , 2020, 11, e00202.	2.5	3
4	RASAL1 and ROS1 Gene Variants in Hereditary Breast Cancer. <i>Cancers</i> , 2020, 12, 2539.	3.7	2
5	<i>HDAC8</i> ; Loss of Function and <i>SHOX</i> ; Haploinsufficiency: Two Independent Genetic Defects Responsible for a Complex Phenotype. <i>Cytogenetic and Genome Research</i> , 2019, 157, 135-140.	1.1	3
6	Mutant <i>MYO1F</i> alters the mitochondrial network and induces tumor proliferation in thyroid cancer. <i>International Journal of Cancer</i> , 2018, 143, 1706-1719.	5.1	35
7	Genomic profiles of primary and metastatic esophageal adenocarcinoma identified via digital sorting of pure cell populations: results from a case report. <i>BMC Cancer</i> , 2018, 18, 889.	2.6	3
8	A <i>de novo</i> <i>PUF60</i> mutation in a child with a syndromic form of coloboma and persistent fetal vasculature. <i>Ophthalmic Genetics</i> , 2017, 38, 590-592.	1.2	16
9	Search for genetic factors in bicuspid aortic valve disease: <i>ACTA2</i> mutations do not play a major role. <i>Interactive Cardiovascular and Thoracic Surgery</i> , 2017, 25, 813-817.	1.1	4