

# Fe Amalia Garcia-Santiago

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8327024/publications.pdf>

Version: 2024-02-01

10  
papers

261  
citations

1683354

5  
h-index

1588620

8  
g-index

10  
all docs

10  
docs citations

10  
times ranked

634  
citing authors

#	ARTICLE	IF	CITATIONS
1	New microdeletion and microduplication syndromes: a comprehensive review. <i>Genetics and Molecular Biology</i> , 2014, 37, 210-219.	0.6	84
2	Clinical utility of chromosomal microarray analysis in invasive prenatal diagnosis. <i>Human Genetics</i> , 2012, 131, 513-523.	1.8	82
3	A New Overgrowth Syndrome is due to Mutations in <i>RNF125</i> . <i>Human Mutation</i> , 2014, 35, 1436-1441.	1.1	33
4	Customized high resolution CGH array for clinical diagnosis reveals additional genomic imbalances in previous well-defined pathological samples. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 1950-1960.	0.7	32
5	Analysis of invdupdel(8p) rearrangement: Clinical, cytogenetic and molecular characterization. <i>American Journal of Medical Genetics, Part A</i> , 2015, 167, 1018-1025.	0.7	22
6	Familial imbalance in 16p13.11 leads to a dosage compensation rearrangement in an unaffected carrier. <i>BMC Medical Genetics</i> , 2014, 15, 116.	2.1	3
7	Unusual four-generation chromosome 22 rearrangement: When "normality" masks abnormality. <i>American Journal of Medical Genetics, Part A</i> , 2009, 149A, 1561-1564.	0.7	2
8	Prenatal ultrasound findings in Koolen-de Vries fetuses: Central nervous system anomalies are frequent markers of this syndrome. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2021, 9, e1649.	0.6	2
9	Rapidly Progressing to ESRD in an Individual with Coexisting ADPKD and Masked Klinefelter and Citelman Syndromes. <i>Genes</i> , 2022, 13, 394.	1.0	1
10	Prenatal Diagnosis of Acromelic Frontonasal Dysostosis. <i>Molecular Syndromology</i> , 2021, 12, 1-5.	0.3	0