## **ALESSANDRA SIRONI**

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/4309906/publications.pdf

Version: 2024-02-01

8 papers

120 citations

6 h-index

1478280

1588896 8 g-index

8 all docs 8 docs citations

times ranked

8

269 citing authors

#	Article	IF	CITATIONS
1	Smith-Magenis Syndromeâ€"Clinical Review, Biological Background and Related Disorders. Genes, 2022, 13, 335.	1.0	18
2	Expanding the Molecular Spectrum of ANKRD11 Gene Defects in 33 Patients with a Clinical Presentation of KBG Syndrome. International Journal of Molecular Sciences, 2022, 23, 5912.	1.8	6
3	High-resolution array-CGH analysis on 46,XX patients affected by early onset primary ovarian insufficiency discloses new genes involved in ovarian function. Human Reproduction, 2019, 34, 574-583.	0.4	32
4	9q34.3 microduplications lead to neurodevelopmental disorders through EHMT1 overexpression. Neurogenetics, 2019, 20, 145-154.	0.7	12
5	Molecular Etiology Disclosed by Array CGH in Patients With Silver–Russell Syndrome or Similar Phenotypes. Frontiers in Genetics, 2019, 10, 955.	1.1	11
6	13q mosaic deletion including RB1 associated to mild phenotype and no cancer outcome – case report and review of the literature. Molecular Cytogenetics, 2018, 11, 53.	0.4	2
7	Rothmund-Thomson Syndrome: Insights from New Patients on the Genetic Variability Underpinning Clinical Presentation and Cancer Outcome. International Journal of Molecular Sciences, 2018, 19, 1103.	1.8	20
8	iPSC-derived neurons of CREBBP - and EP300 -mutated Rubinstein-Taybi syndrome patients show morphological alterations and hypoexcitability. Stem Cell Research, 2018, 30, 130-140.	0.3	19