Daniela Rusconi

List of Publications by Year
in descending order

12	277	1040056 9	1199594 12
papers	citations	h-index	g-index

12	12	12	734
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Erythroid differentiation and maturation from peripheral CD34+ cells in liquid culture: Cellular and molecular characterization. Blood Cells, Molecules, and Diseases, 2008, 40, 148-155.	1.4	45
2	Gene dosage as a relevant mechanism contributing to the determination of ovarian function in Turner syndrome. Human Reproduction, 2014, 29, 368-379.	0.9	39
3	Characterization of 14 novel deletions underlying Rubinsteinâ€“Taybi syndrome: an update of the CREBBP deletion repertoire. Human Genetics, 2015, 134, 613-626.	3.8	38
4	From Whole Gene Deletion to Point Mutations ofEP300-Positive Rubinstein-Taybi Patients: New Insights into the Mutational Spectrum and Peculiar Clinical Hallmarks. Human Mutation, 2016, 37, 175-183.	2.5	36
5	Oligoclonal bands in the cerebrospinal fluid of amyotrophic lateral sclerosis patients with disease-associated mutations. Journal of Neurology, 2013, 260, 85-92.	3.6	24
6	Juxtaposition of heterochromatic and euchromatic regions by chromosomal translocation mediates a heterochromatic long-range position effect associated with a severe neurological phenotype. Molecular Cytogenetics, 2012, 5, 16.	0.9	22
7	Familial intragenic duplication of ANKRD11 underlying three patients of KBG syndrome. Molecular Cytogenetics, 2015, 8, 20.	0.9	18
8	Combined characterization of a pituitary adenoma and a subcutaneous lipoma in a MEN1 patient with a whole gene deletion. Cancer Genetics, 2011, 204, 309-315.	0.4	16
9	Genomic imbalances in patients with a clinical presentation in the spectrum of Cornelia de Lange syndrome. BMC Medical Genetics, 2013, 14, 41.	2.1	15
10	SNPs and real-time quantitative PCR method for constitutional allelic copy number determination, the VPREB1 marker case. BMC Medical Genetics, 2011, 12, 61.	2.1	8
11	Design and validation of a pericentromeric BAC clone set aimed at improving diagnosis and phenotype prediction of supernumerary marker chromosomes. Molecular Cytogenetics, 2013, 6, 45.	0.9	8
12	A novel mosaic NSD1 intragenic deletion in a patient with an atypical phenotype. American Journal of Medical Genetics, Part A, 2013, 161, 611-618.	1.2	8