Ilaria Catusi

List of Publications by Year
in descending order

13	88	1478505 6	1474206 9
papers	citations	h-index	g-index

13	13	13	157
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Characterization of Chromosomal Breakpoints in 12 Cases with 8p Rearrangements Defines a Continuum of Fragility of the Region. International Journal of Molecular Sciences, 2022, 23, 3347.	4.1	6
2	12q21 Interstitial Deletions: Seven New Syndromic Cases Detected by Array-CGH and Review of the Literature. Genes, 2022, 13, 780.	2.4	0
3	8p23.2-pter Microdeletions: Seven New Cases Narrowing the Candidate Region and Review of the Literature. Genes, 2021, 12, 652.	2.4	11
4	Histone Deacetylase Inhibitors Ameliorate Morphological Defects and Hypoexcitability of iPSC-Neurons from Rubinstein-Taybi Patients. International Journal of Molecular Sciences, 2021, 22, 5777.	4.1	7
5	Ten new cases of Balanced Reciprocal Translocation Mosaicism (BRTM): Reproductive implications, frequency and mechanism. European Journal of Medical Genetics, 2020, 63, 103639.	1.3	6
6	Testing single/combined clinical categories on 5110 Italian patients with developmental phenotypes to improve arrayâ€based detection rate. Molecular Genetics & Genomic Medicine, 2020, 8, e1056.	1.2	6
7	Recombinant Chromosome 7 Driven by Maternal Chromosome 7 Pericentric Inversion in a Girl with Features of Silver-Russell Syndrome. International Journal of Molecular Sciences, 2020, 21, 8487.	4.1	1
8	Instability of Short Arm of Acrocentric Chromosomes: Lesson from Non-Acrocentric Satellited Chromosomes. Report of 24 Unrelated Cases. International Journal of Molecular Sciences, 2020, 21, 3431.	4.1	3
9	Molecular cytogenetics characterization of seven small supernumerary marker chromosomes derived from chromosome 19: Genotype-phenotype correlation and review of the literature. European Journal of Medical Genetics, 2018, 61, 173-180.	1.3	5
10	13q mosaic deletion including RB1 associated to mild phenotype and no cancer outcome â€“ case report and review of the literature. Molecular Cytogenetics, 2018, 11, 53.	0.9	2
11	iPSC-derived neurons of CREBBP - and EP300 -mutated Rubinstein-Taybi syndrome patients show morphological alterations and hypoexcitability. Stem Cell Research, 2018, 30, 130-140.	0.7	19
12	7p22.1 microduplication syndrome: Clinical and molecular characterization of an adult case and review of the literature. European Journal of Medical Genetics, 2015, 58, 578-583.	1.3	9
13	Cdc14 Inhibition by the Spindle Assembly Checkpoint Prevents Unscheduled Centrosome Separation in Budding Yeast. Molecular Biology of the Cell, 2009, 20, 2626-2637.	2.1	13