AndrÃ© M Travessa

List of Publications by Year
in descending order

8 papers	109 citations	1684188 5 h-index	1588992 8 g-index
8 all docs	8 docs citations	8 times ranked	227 citing authors

#	Article	IF	Citations
1	Fifteen Years of Clinical Trials inÂHuntingtonâ€™s Disease: A Very Low ClinicalÂDrug Development Success Rate. Journal of Huntington's Disease, 2017, 6, 157-163.	1.9	50
2	Genomic sequencing highlights the diverse molecular causes of Perrault syndrome: a peroxisomal disorder (PEX6), metabolic disorders (CLPP, GGPS1), and mtDNA maintenance/translation disorders (LARS2, TFAM). Human Genetics, 2020, 139, 1325-1343.	3.8	21
3	High prevalence of variants in skeletal dysplasia associated genes in individuals with short stature and minor skeletal anomalies. European Journal of Endocrinology, 2021, 185, 691-705.	3.7	13
4	Prenatal diagnosis of holoprosencephaly associated with Smithâ€“Lemliâ€“Opitz syndrome (SLOS) in a 46,XX fetus. Taiwanese Journal of Obstetrics and Gynecology, 2017, 56, 541-544.	1.3	7
5	Clinical and Molecular Description of 16 Families With Heterozygous IHH Variants. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 2654-2666.	3.6	6
6	Reporting and methodological quality of clinical trials on exercise therapy for Parkinson's disease. Parkinsonism and Related Disorders, 2019, 69, 150-156.	2.2	5
7	Upper limb phocomelia: A prenatal case of thrombocytopenia-absent radius (TAR) syndrome illustrating the importance of chromosomal microarray in limb reduction defects. Taiwanese Journal of Obstetrics and Gynecology, 2020, 59, 318-322.	1.3	5
8	Spondyloepiphyseal dysplasia type Stanescu: Expanding the clinical and molecular spectrum of a very rare type II collagenopathy. American Journal of Medical Genetics, Part A, 2020, 182, 2715-2721.	1.2	2