F Vansenne

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/10102966/publications.pdf

Version: 2024-02-01

18	349	933447 10	888059 17
papers	citations	h-index	g-index

18	18	18	785
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Opposite Modulation of RAC1 by Mutations in TRIO Is Associated with Distinct, Domain-Specific Neurodevelopmental Disorders. American Journal of Human Genetics, 2020, 106, 338-355.	6.2	58
2	The psychological impact of testing for thrombophilia: a systematic review. Journal of Thrombosis and Haemostasis, 2008, 6, 1099-1104.	3.8	57
3	Disruption of RFX family transcription factors causes autism, attention-deficit/hyperactivity disorder, intellectual disability, and dysregulated behavior. Genetics in Medicine, 2021, 23, 1028-1040.	2.4	34
4	A genotype-first approach identifies an intellectual disability-overweight syndrome caused by PHIP haploinsufficiency. European Journal of Human Genetics, 2018, 26, 54-63.	2.8	32
5	The effects of screening on health behaviour: a summary of the results of randomized controlled trials. Journal of Public Health, 2011, 33, 71-79.	1.8	31
6	Two Novel Mutations in the SLC25A4 Gene in a Patient with Mitochondrial Myopathy. JIMD Reports, 2015, 22, 39-45.	1.5	28
7	De novo variants in CDK13 associated with syndromic ID/DD: Molecular and clinical delineation of 15 individuals and a further review. Clinical Genetics, 2018, 93, 1000-1007.	2.0	20
8	Genotypic diversity and phenotypic spectrum of infantile liver failure syndrome type 1 due to variants in LARS1. Genetics in Medicine, 2020, 22, 1863-1873.	2.4	19
9	Delineating the molecular and phenotypic spectrum of the SETD1B-related syndrome. Genetics in Medicine, 2021, 23, 2122-2137.	2.4	16
10	The effect of cervical X-irradiation on activity index of thyrocytes and plasma TSH: A pre-clinical model for radiation-induced thyroid damage. Journal of Endocrinological Investigation, 2005, 28, 261-269.	3.3	13
11	Economic analysis of chromosome testing in couples with recurrent miscarriage to prevent handicapped offspring. Human Reproduction, 2013, 28, 1737-1742.	0.9	8
12	Session 07: Psychology & Counselling 1. Human Reproduction, 2010, 25, i14-i17.	0.9	7
13	Fever-Induced Paroxysmal Weakness and Encephalopathy (FIPWE)â€"Part of a Phenotypic Continuum in Patients With ATP1A3 Mutations?. Pediatric Neurology, 2018, 81, 57-58.	2.1	7
14	De novo variants in CAMTA1 cause a syndrome variably associated with spasticity, ataxia, and intellectual disability. European Journal of Human Genetics, 2020, 28, 763-769.	2.8	7
15	Endocrine Intervention During Irradiation Does Not Prevent Damage To the Thyroid Gland. Thyroid, 2006, 16, 387-395.	4.5	5
16	Knowledge and perceived risks in couples undergoing genetic testing after recurrent miscarriage or for poor semen quality. Reproductive BioMedicine Online, 2011, 23, 525-533.	2.4	4
17	How to detect late-onset inborn errors of metabolism in patients with movement disorders â€“ A modern diagnostic approach. Parkinsonism and Related Disorders, 2021, 85, 124-132.	2.2	3
18	SESSION 05: EARLY PREGNANCY. Human Reproduction, 2012, 27, ii9-ii11.	0.9	0