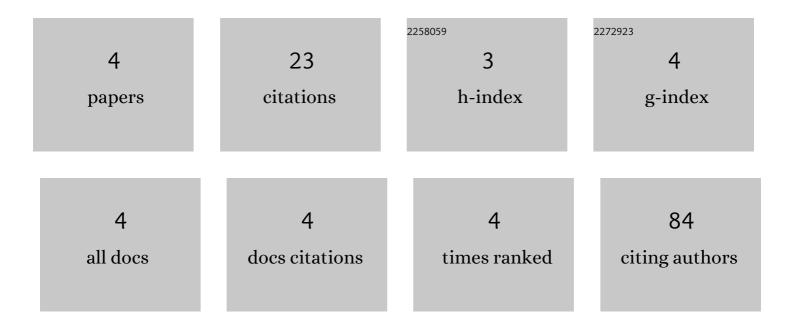
Virginie Benoit

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/758740/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	15q24.1 BP4-BP1 microdeletion unmasking paternally inherited functional polymorphisms combined with distal 15q24.2q24.3 duplication in a patient with epilepsy, psychomotor delay, overweight, ventricular arrhythmia. European Journal of Medical Genetics, 2018, 61, 459-464.	1.3	11
2	Prenatal Diagnosis of a 2.5 Mb De Novo 17q24.1q24.2 Deletion Encompassing <i> KPNA2</i> and <i> PSMD12</i> Genes in a Fetus with Craniofacial Dysmorphism, Equinovarus Feet, and Syndactyly. Case Reports in Genetics, 2017, 2017, 1-5.	0.2	7
3	First prenatal case of proximal 19p13.12 microdeletion syndrome: New insights and new delineation of the syndrome. European Journal of Medical Genetics, 2018, 61, 322-328.	1.3	3
4	Two new cases of interstitial 7q35q36.1 deletion including <i>CNTNAP2</i> and <i>KMT2C</i> . Molecular Genetics & Genomic Medicine, 2021, 9, e1645.	1.2	2