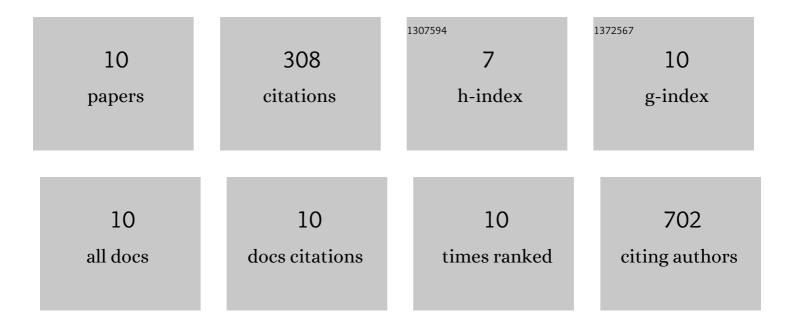
## Fleur Vansenne

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11263709/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Phenotype, cancer risk, and surveillance in Beckwith–Wiedemann syndrome depending on molecular genetic subgroups. American Journal of Medical Genetics, Part A, 2016, 170, 2248-2260.	1.2	163
2	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
3	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
4	<i>ZMYND11</i> â€related syndromic intellectual disability: 16 patients delineating and expanding the phenotypic spectrum. Human Mutation, 2020, 41, 1042-1050.	2.5	20
5	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
6	Delineating the molecular and phenotypic spectrum of the SETD1B-related syndrome. Genetics in Medicine, 2021, 23, 2122-2137.	2.4	16
7	Expanding the <i>ADCY5</i> phenotype toward spastic paraparesis. Neurology: Genetics, 2018, 4, e214.	1.9	11
8	Fever-Induced Paroxysmal Weakness and Encephalopathy (FIPWE)—Part of a Phenotypic Continuum in Patients With ATP1A3 Mutations?. Pediatric Neurology, 2018, 81, 57-58.	2.1	7
9	Trisomy 4 mosaicism: Delineation of the phenotype. American Journal of Medical Genetics, Part A, 2016, 170, 1040-1045.	1.2	3
10	How to detect late-onset inborn errors of metabolism in patients with movement disorders – A modern diagnostic approach. Parkinsonism and Related Disorders, 2021, 85, 124-132.	2.2	3