## Siren Berland

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6247019/publications.pdf

Version: 2024-02-01

		1684188	1588992	
8	133	5	8	
papers	citations	h-index	g-index	
9	9	9	395	
all docs	docs citations	times ranked	citing authors	

#	Article	IF	CITATIONS
1	CTCF variants in 39 individuals with a variable neurodevelopmental disorder broaden the mutational and clinical spectrum. Genetics in Medicine, 2019, 21, 2723-2733.	2.4	48
2	Population prevalence and inheritance pattern of recurrent CNVs associated with neurodevelopmental disorders in 12,252 newborns and their parents. European Journal of Human Genetics, 2021, 29, 205-215.	2.8	40
3	A clinical scoring system for congenital contractural arachnodactyly. Genetics in Medicine, 2020, 22, 124-131.	2.4	17
4	Further delineation of the clinical spectrum of White $\hat{a} \in \text{``Sutton syndrome: } 12$ new individuals and a review of the literature. European Journal of Human Genetics, 2022, 30, 95-100.	2.8	8
5	Janus-faced EPHB4-associated disorders: novel pathogenic variants and unreported intrafamilial overlapping phenotypes. Genetics in Medicine, 2021, 23, 1315-1324.	2.4	6
6	Deep exploration of a <i>CDKN1C</i> mutation causing a mixture of Beckwith-Wiedemann and IMAGe syndromes revealed a novel transcript associated with developmental delay. Journal of Medical Genetics, 2022, 59, 155-164.	3.2	6
7	Double paternal uniparental isodisomy 7 and 15 presenting with Beckwith–Wiedemann spectrum features. Journal of Physical Education and Sports Management, 2021, 7, a006113.	1.2	5
8	The blended phenotype of a germline <i>RIT1</i> and a mosaic <i>PIK3CA</i> variant. Journal of Physical Education and Sports Management, 2021, 7, a006121.	1.2	3