

Siren Berland

List of Publications by Year in descending order

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

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8
papers

133
citations

1684188

5
h-index

1588992

8
g-index

9
all docs

9
docs citations

9
times ranked

395
citing authors

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