

Stefanie Brock

List of Publications by Year in descending order

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Version: 2024-02-01

10
papers

118
citations

1478505

6
h-index

1588992

8
g-index

10
all docs

10
docs citations

10
times ranked

243
citing authors

#	ARTICLE	IF	CITATIONS
1	Overlapping cortical malformations in patients with pathogenic variants in <i>GRIN1</i> and <i>GRIN2B</i> . <i>Journal of Medical Genetics</i> , 2023, 60, 183-192.	3.2	1
2	Malformations of cerebral development and clues from the peripheral nervous system: A systematic literature review. <i>European Journal of Paediatric Neurology</i> , 2022, 37, 155-164.	1.6	1
3	Defining the phenotypical spectrum associated with variants in <i>TUBB2A</i> . <i>Journal of Medical Genetics</i> , 2021, 58, 33-40.	3.2	11
4	Neuropathology of genetically defined malformations of cortical development – A systematic literature review. <i>Neuropathology and Applied Neurobiology</i> , 2021, 47, 585-602.	3.2	9
5	Overcoming the Challenges of High Quality RNA Extraction from Core Needle Biopsy. <i>Biomolecules</i> , 2021, 11, 621.	4.0	7
6	A rare cause of postmenopausal hyperandrogenism. <i>BMJ Case Reports</i> , 2021, 14, e237505.	0.5	0
7	Genetic heterogeneity of polymicrogyria: study of 123 patients using deep sequencing. <i>Brain Communications</i> , 2021, 3, fcaa221.	3.3	22
8	Recurrent NEDD4L Variant in Periventricular Nodular Heterotopia, Polymicrogyria and Syndactyly. <i>Frontiers in Genetics</i> , 2020, 11, 26.	2.3	12
9	TMX2 Is a Crucial Regulator of Cellular Redox State, and Its Dysfunction Causes Severe Brain Developmental Abnormalities. <i>American Journal of Human Genetics</i> , 2019, 105, 1126-1147.	6.2	25
10	Tubulinopathies continued: refining the phenotypic spectrum associated with variants in TUBG1. <i>European Journal of Human Genetics</i> , 2018, 26, 1132-1142.	2.8	30