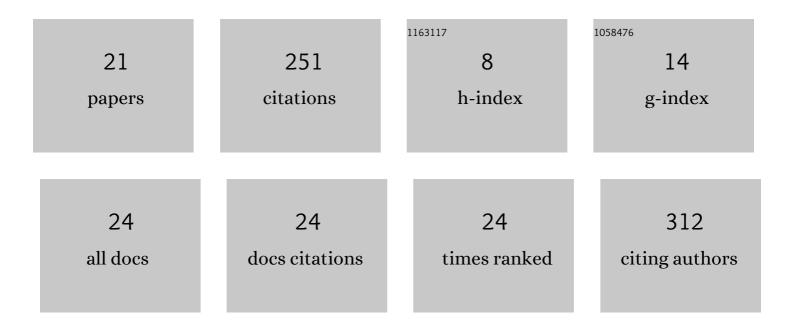
Miriam Kessi

List of Publications by Year in descending order

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MIDIAM KESSI

| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Genetic etiologies of the electrical status epilepticus during slow wave sleep: systematic review. BMC Genetics, 2018, 19, 40. | 2.7 | 39 |
| 2 | Calcium channelopathies and intellectual disability: a systematic review. Orphanet Journal of Rare Diseases, 2021, 16, 219. | 2.7 | 33 |
| 3 | Myelin Oligodendrocyte Glycoprotein (MOG) Antibody Diseases in Children in Central South China: Clinical Features, Treatments, Influencing Factors, and Outcomes. Frontiers in Neurology, 2019, 10, 868. | 2.4 | 29 |
| 4 | Intellectual Disability and Potassium Channelopathies: A Systematic Review. Frontiers in Genetics, 2020, 11, 614. | 2.3 | 28 |
| 5 | Rare Copy Number Variations and Predictors in Children With Intellectual Disability and Epilepsy. Frontiers in Neurology, 2018, 9, 947. | 2.4 | 23 |
| 6 | The Contribution of HCN Channelopathies in Different Epileptic Syndromes, Mechanisms, Modulators, and Potential Treatment Targets: A Systematic Review. Frontiers in Molecular Neuroscience, 2022, 15, . | 2.9 | 15 |
| 7 | Efficacy of different treatment modalities for acute and chronic phases of the febrile infection-related epilepsy syndrome: A systematic review. Seizure: the Journal of the British Epilepsy Association, 2020, 79, 61-68. | 2.0 | 13 |
| 8 | The Study of Genetic Susceptibility and Mitochondrial Dysfunction in Mesial Temporal Lobe Epilepsy. Molecular Neurobiology, 2020, 57, 3920-3930. | 4.0 | 11 |
| 9 | Familial SYN1 variants related neurodevelopmental disorders in Asian pediatric patients. BMC Medical Genomics, 2021, 14, 182. | 1.5 | 9 |
| 10 | Treatment for the Benign Childhood Epilepsy With Centrotemporal Spikes: A Monocentric Study. Frontiers in Neurology, 2021, 12, 670958. | 2.4 | 8 |
| 11 | De Novo KCNQ2 Mutation in One Case of Epilepsy of Infancy With Migrating Focal Seizures That Evolved to Infantile Spasms. Child Neurology Open, 2018, 5, 2329048X1876773. | 1.1 | 7 |
| 12 | Immunotherapies for Anti-N-M-methyl-D-aspartate Receptor Encephalitis: Multicenter Retrospective Pediatric Cohort Study in China. Frontiers in Pediatrics, 2021, 9, 691599. | 1.9 | 7 |
| 13 | Etiologic Classification of 541 Infantile Spasms Cases: A Cohort Study. Frontiers in Pediatrics, 2022, 10, 774828. | 1.9 | 7 |
| 14 | A Case With 4 de Novo Copy Number Variations With Clinical Features That Overlap 1q43q44 Microdeletion and 3q29 Microduplication Syndromes. Child Neurology Open, 2018, 5, 2329048X1879820. | 1.1 | 4 |
| 15 | Correlation Analyses of Clinical Manifestations and Variant Effects in KCNB1-Related Neurodevelopmental Disorder. Frontiers in Pediatrics, 2021, 9, 755344. | 1.9 | 4 |
| 16 | Urine Organic Acids as Metabolic Indicators for Global Developmental Delay/Intellectual Disability in Chinese Children. Frontiers in Molecular Biosciences, 2021, 8, 792319. | 3.5 | 4 |
| 17 | The Recommendations for the Management of Chinese Children With Epilepsy During the COVID-19 Outbreak. Frontiers in Pediatrics, 2020, 8, 495. | 1.9 | 3 |
| 18 | Observed and self-reported hand hygiene compliance and associated factors among healthcare workers at a county referral hospital in Kenya. Scientific African, 2021, 14, e00984. | 1.5 | 2 |

| # | Article | IF | CITATIONS |
|----|--|-----|-----------|
| 19 | Restoration of Sarco/Endoplasmic Reticulum Ca2+-ATPase Activity Functions as a Pivotal Therapeutic Target of Anti-Glutamate-Induced Excitotoxicity to Attenuate Endoplasmic Reticulum Ca2+ Depletion. Frontiers in Pharmacology, 2022, 13, 877175. | 3.5 | 2 |
| 20 | The patterns of response of 11 regimens for infantile spasms. Scientific Reports, 2020, 10, 11509. | 3.3 | 1 |
| 21 | A survey on pediatric anti-N-methyl-D-aspartate-receptor encephalitis treatment strategies in China. Chinese Medical Journal, 2021, 134, 1498-1499. | 2.3 | 1 |