

Phenotype-Driven Virtual Panel Is an Effective Method Neurological Disease

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Citation Report

| # | ARTICLE | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Genomic testing and counseling: The contribution of next-generation sequencing to epilepsy genetics. <i>Annals of Human Genetics</i> , 2020, 84, 431-436. | 0.3 | 5 |
| 2 | Clinical and Genetic Study on a Chinese Patient with Infantile Onset Epileptic Encephalopathy carrying a PPP3CA Null Variant: a case report. <i>BMC Pediatrics</i> , 2020, 20, 315. | 0.7 | 4 |
| 3 | Molecular analysis and clinical diversity of distal hereditary motor neuropathy. <i>European Journal of Neurology</i> , 2020, 27, 1319-1326. | 1.7 | 28 |
| 4 | Molecular diagnosis of muscular diseases in outpatient clinics. <i>Neurology: Genetics</i> , 2020, 6, e408. | 0.9 | 15 |
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| 9 | More autosomal dominant SPG18 cases than recessive? The first ADâ€¦SPG18 pedigree in Chinese and literature review. <i>Brain and Behavior</i> , 2021, 11, e32395. | 1.0 | 5 |
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| 12 | The pathogenetic basis for a disease continuum in early- and late-onset ataxia-dystonia supports a unified genetic diagnostic approach. <i>European Journal of Paediatric Neurology</i> , 2023, 43, 44-51. | 0.7 | 2 |
| 13 | Solving inherited white matter disorder etiologies in the neurology clinic: Challenges and lessons learned using next-generation sequencing. <i>Frontiers in Neurology</i> , 0, 14, . | 1.1 | 2 |