Special Issue

Pediatric Epilepsy: Insights from Genetics to Mechanisms

Message from the Guest Editors

Pediatric epilepsy (PE) comprises a heterogeneous group of disorders characterized by recurrent, unprovoked seizures with onset in infancy or childhood, often resistant to standard anti-epileptic drugs. Over the past two decades, advances in genomics and nextgeneration sequencing have significantly expanded our understanding of the molecular basis of PE. Complementary multi-omics approaches have further illuminated the complex genetic architecture of PE, uncovering key molecular pathways involved in its pathogenesis. Despite these advances, significant challenges remain in translating genetic findings into biological insight and clinical impact. Nowadays, emerging technologies such as single-cell and spatial transcriptomics are revealing the cellular heterogeneity and spatial organization of brain development in both health and disease. This area of research is critically important—not only for advancing our understanding of the biological basis of PE, but also for driving the development of effective therapeutic strategies.

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