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Neurodevelopmental Phenotyping to Inform Treatment in Neurogenetic Syndromes

Guest Editor:

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Message from the Guest Editor

A critical goal for translational research in neurogenetic syndromes is to identify developmental, behavioral, and pharmaceutical treatments that address neurodevelopmental challenges associated with specific conditions (e.g., Down syndrome, fragile X syndrome, Angelman syndrome, Rett syndrome). Innovations in this area have the potential to offer novel and effective biomedical and behavioral intervention for individuals with specific neurogenetic conditions; however, there are gaps in our knowledge that hinder the readiness of treatments, namely the need for (1) validated outcome measures that quantify the effects of new treatments on cognition and behavior in specific conditions; and (2) more advanced characterizations of neurodevelopmental heterogeneity specific conditions within to inform treatment development. In this Special Issue, all related articles and reviews are welcome













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Editor-in-Chief

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Message from the Editor-in-Chief

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