## **Special Issue**

# Update on the Treatment of Fragile X Syndrome

## Message from the Guest Editors

Fragile X syndrome (FXS) is a well-defined genetic cause of inherited intellectual disability (ID). It is also the bestunderstood single-gene factor associated with autism spectrum disorder (ASD). Deficits in the fragile X gene's key protein seem to be the critical unifying factor linked, at a synaptic level, to dysfunction in brain pathways and to at least some aspects of behavioral symptoms in idiopathic ASD. Thus, further advances in the understanding of FXS may help to inform studies on ID and other ASD in highly heterogenous idiopathic ASD. In last decade, major progress in elucidating the underlying causes of FXS has generated new targeted approaches to manage them. Indeed, among all neurodevelopmental disabilities, FXS has been at the forefront of efforts to test preclinical evidence for these interventions in clinical studies. In this issue, we focus on the objective and/or directly observable quantitative measures of FXS pathophysiology of meaningful relevance not only to the treatment challenges but also to understanding the developmental trajectory in FXS, including novel topics with implications on ASD and ID.

#### **Guest Editors**

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## Deadline for manuscript submissions

closed (15 June 2020)



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Impact Factor 2.8
CiteScore 5.6
Indexed in PubMed



mdpi.com/si/29245

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You are invited to contribute a research article or a comprehensive review for consideration and publication in *Brain Sciences* (ISSN 2076-3425). *Brain Sciences* is an open access, peer-reviewed scientific journal that publishes original articles, critical reviews, research notes, and short communications on neuroscience. The scientific community and the general public can access the content free of charge as soon as it is published.

## Editor-in-Chief

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