

## Special Issue

# Fragile X Syndrome: From Molecular Mechanisms to Therapeutic Approaches

### Message from the Guest Editors

This Special Issue will cover all aspects of the molecular and cellular mechanisms leading to neuropathology in fragile X syndrome. Since the discovery of the *FMR1* gene more than 30 years ago and the establishment of different research models from knockout mice to induced pluripotent stem cells derived from patients, the understanding of FXS has advanced remarkably. However, critical questions about the role of FMRP in neural tissue remain unanswered: what are the most important targets of FMRP that react to its disappearance and are responsible to downstream detrimental effects? What aspects of neuronal plasticity are most affected by lack of FMRP and how are they manifested during embryonic development and adult life? What type of neuronal and synaptic deficits caused by absence of FMRP are reversible and what strategies could be used to promote reversibility? We invite authors to submit details of their research focused on these and other open questions as original research articles or literature reviews.

### Guest Editors

Prof. Dr. Menahem Segal

Department of Neurobiology, The Weizmann Institute, Herzl St 234, Rehovot 7610001, Israel

Dr. Telias Michael

Department of Molecular and Cell Biology, University of California, Berkeley, CA, USA

### Deadline for manuscript submissions

closed (15 November 2022)



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Editorial Office  
MDPI, Grosspeteranlage 5  
4052 Basel, Switzerland  
Tel: +41 61 683 77 34  
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### Editor-in-Chief

Prof. Dr. Maurizio Battino

Department of Odontostomatologic and Specialized Clinical Sciences,  
Sez-Biochimica, Faculty of Medicine, Università Politecnica delle  
Marche, Via Ranieri 65, 60100 Ancona, Italy

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