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Selected Papers from the International Workshop on Fragile X and Other Neurodevelopmental Disorders

Guest Editors:

Dr. Pietro Chiurazzi

Prof. Dr. Giovanni Neri

Dr. Maria Giuseppina Miano

Deadline for manuscript submissions:
closed (28 February 2020)

Message from the Guest Editors

The 19th International Workshop Fragile X and other Neurodevelopmental Disorders will be held on September 18-21, 2019 in Sorrento, Italy.

This Workshop has two main aims: 1) To foster interdisciplinary and multidisciplinary research on the genetic bases, molecular pathogenesis, clinical presentation of, and innovative therapies for, intellectual disability, including fragile X syndrome, other X-linked conditions, autism, and other neurodevelopmental disorders; 2) to provide an informal setting where trainees and young investigators across a range of scientific disciplines can build a network among both established and junior investigators. All accepted papers from the Workshop will be entitled a 15% discount.

The current Special Issue invites submissions of unpublished original work describing recent advances on all aspects related to the following topics:

Fragile X and FMR1-related diseases; Other XLID and autosomal ID conditions; Syndromes caused by multiple “single gene” defects; Mechanisms of disease using human models; Autism, the unending nightmare of the geneticist; Epigenetics signatures and diagnostic biomarkers; Therapeutic perspectives.

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Message from the Editor-in-Chief

Genes are central to our understanding of biology, and modern advances such as genomics and genome editing have maintained genetics as a vibrant, diverse and fastmoving field. There is a need for good quality, open access journals in this area, and the *Genes* team aims to provide expert manuscript handling, serious peer review, and rapid publication across the whole discipline of genetics. Starting in 2010, the journal is now well established and recognised.

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