# Special Issue

# Molecular Genetics of Neurodevelopmental Disorders

### Message from the Guest Editors

Neurodevelopmental disorders affect a significant number of children during vulnerable periods of their lives and can have a substantial lifelong impact on them and their families. In particular, although the genetic component is large and many genetic variants have been identified, the mechanism through which they predispose towards the autism phenotype and how treatments can mitigate the development process, remain unclear. The genetic component is highly complex, not only in terms of the number of genes known to be involved, but also in terms of multiple pathways, low penetrance/high prevalence variants versus the opposite, de novo versus inherited variants, polygenetic interaction, and microRNA gene regulation. We encourage the submission of manuscripts describing any of the above-listed factors that provide insight into the etiological or pathophysiological processes underlying neurodevelopmental disorders. We also encourage submissions improving the description of genetic variants or conditions that help elucidate underlying biological mechanisms.

#### **Guest Editors**

Prof. Dr. Richard Eugene Frye

Autism Discovery and Treatment Foundation and Rossignol Medical Center, Phoenix, AZ, USA

Dr. Richard G. Boles

The Center for Neurological and Neurodevelopmental Health, Vorhees, NJ, USA

Prof. Dr. Stephen G. Kahler

UK Healthcare, University of Kentucky, Lexington, KY, USA

### Deadline for manuscript submissions

closed (15 December 2024)

G C A T T A C G G C A T

# Genes

an Open Access Journal by MDPI

Impact Factor 2.8
CiteScore 5.5
Indexed in PubMed



mdpi.com/si/177435

Genes Editorial Office MDPI, Grosspeteranlage 5 4052 Basel, Switzerland Tel: +41 61 683 77 34

mdpi.com/journal/genes

genes@mdpi.com



# G C A T T A C G G C A T

# Genes

an Open Access Journal by MDPI

Impact Factor 2.8 CiteScore 5.5 Indexed in PubMed



## **About the Journal**

### Message from the Editor-in-Chief

Genes is central to our understanding of biology, and modern advances such as genomics and genome editing have maintained genetics as a vibrant, diverse and fast-moving 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. Why not consider Genes for your next genetics paper?

#### Editor-in-Chief

### Prof. Dr. Selvarangan Ponnazhagan

Department of Pathology, The University of Alabama at Birmingham, 1825 University Blvd, SHEL 814, Birmingham, AL 35294-2182, USA

#### **Author Benefits**

## Open Access:

free for readers, with article processing charges (APC) paid by authors or their institutions.

### **High Visibility:**

indexed within Scopus, SCIE (Web of Science), PubMed, MEDLINE, PMC, Embase, PubAg, and other databases.

#### **Journal Rank:**

JCR - Q2 (Genetics and Heredity) / CiteScore - Q2 (Genetics (clinical))

