Special Issue

Fetal Genetic Disorders: Diagnosis and Therapy

Message from the Guest Editor

In recent years, technological advances have led to excellent results in prenatal genomic sequencing, enabling the identification of a significant number of fetal genetic abnormalities that until recently were labeled only after birth. Although improved practitioner expertise has led to a greater understanding of fetal malformations from an ultrasound-focused point of view, adequate classification in the context of genetic abnormalities still remains a challenge. In this Special Issue, we aim to attract cutting-edge studies on the role of genetic research in the diagnosis of genetic syndromes associated with fetal malformations.

Guest Editor

Dr. Gianluca Campobasso

Department of Obstetrics and Gynecology, Vito Fazzi Hospital, 73100 Lecce, Italy

Deadline for manuscript submissions

20 July 2026

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/262578

Genes

Editorial Office MDPI, Grosspeteranlage 5 4052 Basel, Switzerland Tel: +41 61 683 77 34 genes@mdpi.com

mdpi.com/journal/genes



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))

