Special Issue

Advances in Single Molecule, Real-Time (SMRT) Sequencing

Message from the Guest Editors

PacBio's single molecule, real-time (SMRT) sequencing technology offers important advantages over the shortread DNA sequencing technologies that currently dominate the market. This includes exceptionally long read lengths (20 kb or more), unparalleled consensus accuracy, and the ability to sequence native, nonamplified DNA molecules. From microbes to vertebrates, long reads are now used to create highly accurate de novo genome assemblies, characterize complex structural variations, permit full-length RNA isoform sequencing, and directly phase variants. The high accuracy further enables low frequency mutation detection and clonal evolution determination. Besides reducing biases, sequencing native DNA also permits the direct measurement of DNA base modifications. Therefore, SMRT sequencing has become an attractive technology in many fields, such as agriculture, basic science, and medical research. This Special Issue is a collection of articles showcasing the latest developments and the breadth of applications enabled by SMRT sequencing technology.

Guest Editors

Dr. Adam Ameur

Department of Immunology, Genetics and Pathology, Uppsala University, Science for Life Laboratory, 75108 Uppsala, Sweden

Dr. Matthew S. Hestand

- 1. Division of Human Genetics, Cincinnati Children's Hospital Medical Center, 3333 Burnet Ave, Cincinnati, OH 45229, USA
- 2. Department of Pediatrics, University of Cincinnati, 2600 Clifton Ave, Cincinnati, OH 45220, USA

Deadline for manuscript submissions

closed (31 January 2019)

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/16577

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))

