



Gene Editing for Disease Modeling and Therapeutics

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submissions:

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Message from the Guest Editor

Dear Colleagues,

This issue will accept manuscripts that use genome-edited, human-induced pluripotent stem cells (hiPSCs) as models for monogenic/complex diseases and as cells relevant for therapeutics.

The gene editing methodology can be used for fixing mutations, creating mutations, and knocking out repressors or enhancers, etc., and as such can serve to generate disease models and their controls for both the studying of diseases and for their therapeutic potential (including issues of off target).

Original research papers, reviews, and methodology papers on these topics and related topics are encouraged. They can deal with any issue relevant to genome editing.

Dr. Rivka Ofir
Guest Editor





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Editor-in-Chief

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Message from the Editor-in-Chief

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