Special Issue

CRISPR/Cas Systems and Genome Editing—3rd Edition

Message from the Guest Editor

The harnessing of CRISPR/Cas (Clustered Regularly Interspaced Short Palindromic Repeats/CRISPRassociated genes) systems for the detection, chemical modification, and sequence editing of nucleic acids has dramatically changed many fields of fundamental science, biotechnology, and biomedicine. Natural CRISPR/Cas systems are well suited to disrupting coding and non-coding sequences in the genome, are much less effective at distinguishing and editing sequences with single-nucleotide differences, and are prone to recognizing similar sequences in the genome. Therefore, further exploration of the CRISPR/Cas field continues both to find new genome and epigenome editors, and to engineer known editors to overcome their shortcomings to make their applications more efficient, predictable, and safe. The aim of this Special Issue is to gather knowledge on the current state of the rapidly growing field of CRIPSR/Cas systems and other genome editing tools in fundamental science, biotechnology, and biomedicine.

Guest Editor

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