Special Issue

Adeno-Associated Virus Biology and AAV Vector-Mediated Gene Therapy

Message from the Guest Editors

Following their successful use as vectors in gene therapy clinical trials, adeno-associated viruses (AAV) have increased in popularity and have become emblematic viruses of the Parvovirinae subfamily. However, grey areas remain in the wild-type AAV life cycle. Whereas AAV was disclosed to the scientific community in 1964, novel coding sequences and proteins were only recently discovered, such as the MAAP (membrane-associated accessory protein) identified by Church's group. Additional knowledge on fundamental biology has a potential impact on AAV vector production, purification and efficiency. In this regard, the assembly-activating protein (AAP) described in 2010 shed new light on the capsid assembly mechanism, leading to a rethinking of Cap ORF molecular design for the generation of AAV variants. This Special Issue of *Microorganisms* invites submissions of manuscripts that address AAV biology and vectorology. Scientific findings must be relevant to recombinant AAV-mediated gene therapy. Original research articles and reviews are welcome. Research areas may include (but are not limited to) the following: genomics, epigenetics, vector design and quality control studies.

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Deadline for manuscript submissions

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