Special Issue

New Insights in Huntington's Disease

Message from the Guest Editor

Huntington's Disease (HD) is a rare, progressive and invariably fatal neurological disease. The genetic mutation that leads to HD was discovered in 1993, but a cure remains elusive. We have made great strides in our understanding of the disease regarding pathology, brain networking, radiological findings, possible biomarkers and clinical manifestations. The possible treatments for the symptoms of HD have progressed. There are now two FDA-approved drugs in the US with more medications under investigation. The development of therapies for HD continues to focus on symptomatic management. This Special Issue will highlight advances in the genetics, epidemiology, diagnosis and treatment of HD, including pathology, imaging, current therapies and potential future therapies. Input from physicians and allied health fields will be considered in the ethical and clinical care advances across all stages of Huntington's Disease.

Guest Editor

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You are invited to contribute a research article or a comprehensive review for consideration and publication in *Brain Sciences* (ISSN 2076-3425). *Brain Sciences* is an open access, peer-reviewed scientific journal that publishes original articles, critical reviews, research notes, and short communications on neuroscience. The scientific community and the general public can access the content free of charge as soon as it is published.

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