



Prions and Prion-Like Mechanisms in Disease and Biological Function

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Deadline for manuscript submissions:

closed (28 February 2023)

Message from the Guest Editors

Research into prions has vastly expanded our knowledge and understanding of an infectious pathogen. Prions are replication-competent assemblies of a host-encoded protein. Recent data suggest that prion transmission can be dissociated from toxicity, raising the question of why some prion and prion-like amyloid aggregates are toxic while others are not, and how self-replication, infectivity, and toxicity are linked at the structural and mechanistic level. These key questions of fundamental importance will be highlighted in this Special Issue. They are by their nature multidisciplinary, and we thus strongly welcome approaches that merge structural, cellular, and molecular biology, biochemistry, biophysics, imaging and computational techniques on topics including, but not limited to the following:

Structural properties of prions and amyloid

Prion-like mechanisms in neurodegenerative disease, in systemic diseases, and in biological functions

Structure–toxicity and structure–infectivity relationships

Structural and dynamic basis of prion strains

Functional role of disease-associated mutations





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