Special Issue

Calpain Family in Health and Diseases: The Road Ahead

Message from the Guest Editor

Calpains, Ca2+-dependent intracellular proteases, are comprised of 15 homologues in mammals, and are classified into conventional and unconventional isozymes. Conventional isozymes are composed of calpain-1 and -2, which are expressed in almost all eukaryotes. Growing evidence suggests that these conventional isozymes modify intracellular signalling molecules, thereby altering cellular processes including inflammatory cascades. Accordingly, defective calpainmediated proteolysis may be involved in the pathogenesis of human diseases, such as cardiometabolic disease, neurodegenerative disorders, and cancer progression. In contrast to conventional isozymes, unconventional calpains are expressed in a tissue-specific manner. Investigations have identified pathogenic roles of unconventional calpains in a variety of diseases, including cancer and retinal degeneration, targeting calpain-3 can induce limb girdle muscular dystrophy type 2A. The current Special Issue highlights recent advances in molecular-based analyses of conventional and unconventional calpains to elucidate the pathophysiological aspects of these molecules and possible clinical applications.

Guest Editor

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Deadline for manuscript submissions

closed (20 January 2025)



International Journal of Molecular Sciences

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