



Tracking Back Proteotoxicity in Neurodegenerative Diseases

Guest Editors:

Prof. Dr. Lucio Tremolizzo

1. School of Medicine and Surgery, Milan Center for Neuroscience (NeuroMI), University of Milano-Bicocca, Monza (MB), Italy
2. Department of Neurology, San Gerardo Hospital, Monza (MB), Italy

Dr. Gessica Sala

Milan Center for Neuroscience (NeuroMI), University of Milano-Bicocca, Milan, Italy

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Message from the Guest Editors

Many neurodegenerative diseases belong to the large category of proteinopathies, conditions characterized by the presence of proteinaceous inclusions within and/or outside the degenerating neurons. The identification of such aggregates supports the view that misfolded proteins represent a basic requirement for the neurodegenerative process and provides input to verify the existence of possible dysfunctions of the biological systems influencing protein homeostasis.

Besides the impairment of intracellular protein catabolic systems, epigenetic mechanisms are often involved, implying a putative environmental contribution to altering the physiological processes involved in the synthesis of disease-specific proteins. This may result in the overproduction of post-translationally modified proteins more prone to aggregation.

The aim of this Special Issue consists in enlarging the focus of our previous Special Issue “Proteotoxicity and Neurodegenerative Diseases”, tracing back even more the steps leading to protein aggregates, including also chromatin and DNA modification and transcription factor interactions, besides RNA metabolism alterations.





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Prof. Dr. José L. Quiles

Department of Physiology,
Institute of Nutrition and Food
Technology “Jose Mataix”,
Biomedical Research Center,
University of Granada, Avda.
Conocimiento s/n, 18100 Armilla,
Granada, Spain

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MDPI, Grosspeteranlage 5
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