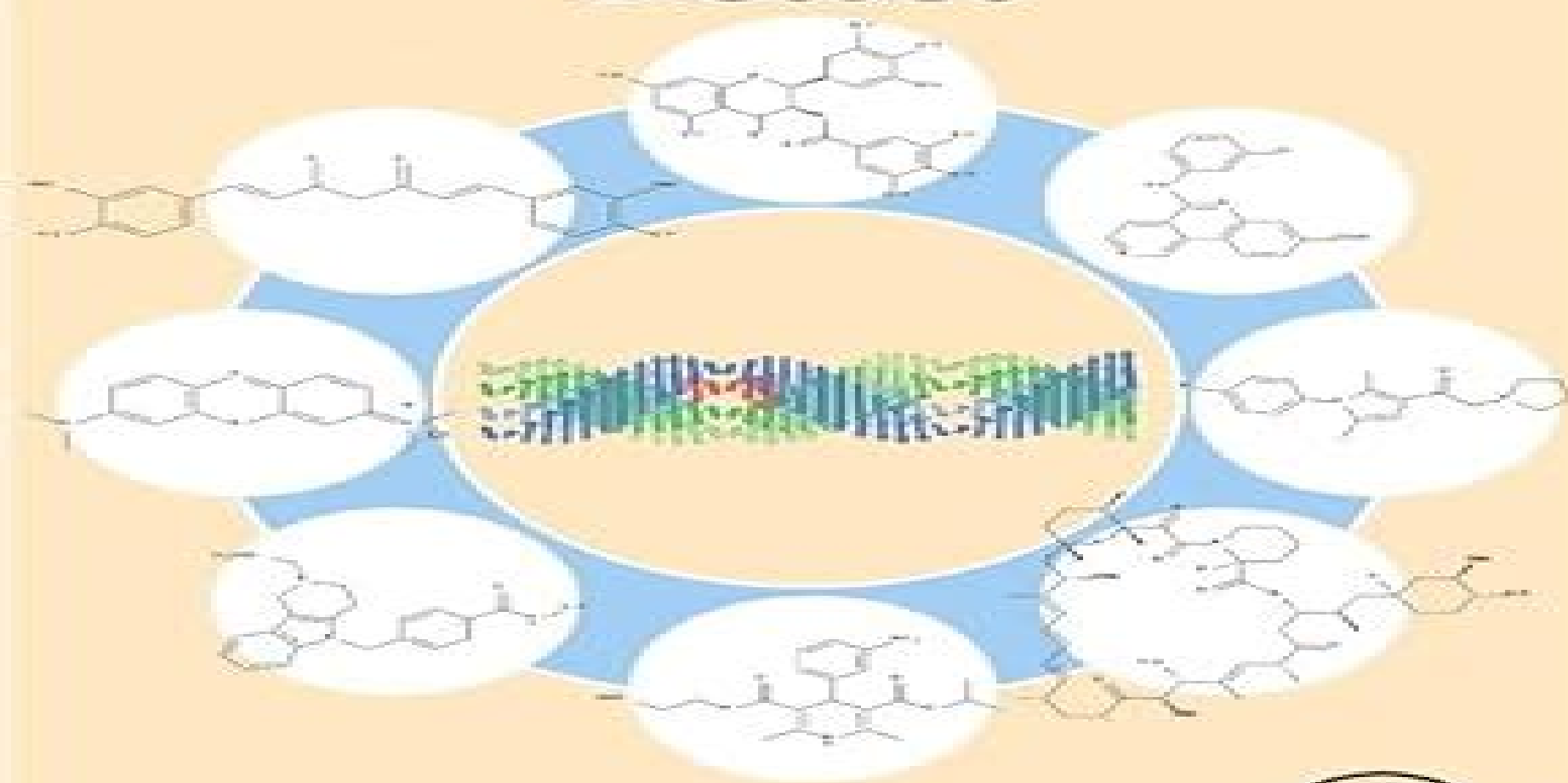
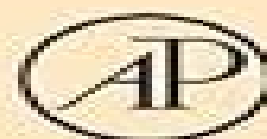


Chemical Modulators of Protein Misfolding and Neurodegenerative Disease



Pierfausto Seneci



**Chemical Modulators Of Protein Misfolding And
Neurodegenerative Disease Hardcover 2015 By
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**Cintia Roodveldt,Tiago F.
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Chemical Modulators Of Protein Misfolding And Neurodegenerative Disease Hardcover 2015 By Pierfausto Seneci:

Chemical Modulators of Protein Misfolding and Neurodegenerative Disease Pierfausto Seneci, 2015-01-14 This book is a neurochemistry based companion for Protein Misfolding and Neurodegenerative Diseases Molecular Targets an Elsevier title by the same author publishing in December 2014 While the first book focuses on biology and molecular targets this companion book describes how these targets are regulated by small molecules and disease modifying compounds The book begins with a brief introduction to how key proteins become dysfunctional and each subsequent chapter describes major disease mechanisms in Alzheimer s and other tauopathies Properties and development status of these molecular targets and disease mechanisms are thoroughly described as are small molecule effectors of autophagy and dis aggregating agents Written to provide comprehensive coverage of neurodegenerative disease modifying compounds Provides discipline specific chapters that cover medicinal chemistry and clinical applications Provides an overview of more than 200 chemical classes and lead compounds acting on selected molecular targets that are of relevance to any neurodegenerative disorder Coverage of misfolding diseases chaperone proteins ubiquitination and autophagy oncology makes this book suitable for structural neurochemists chemists biologists non CNS scientists and scientists interested in drug discovery

Molecular Targets in Protein Misfolding and Neurodegenerative Disease Pierfausto Seneci, 2014-10-07 Aimed at drug discoverers i e any scientist who is interested in neurodegenerative diseases in general and in finding disease modifying treatments in particular the first edition of Molecular Targets in Protein Misfolding and Neurodegenerative Disease will contain both a detailed discipline specific coverage paragraphs on medicinal chemistry on clinical and preclinical characterization of compounds in development on target identification and validation on genetic factors influencing a pathology etc and a drug discovery oriented overall evaluation of each target validation druggability existing leads etc Together these will satisfy the needs of various audiences including in vitro biologists pharmacologists medicinal chemists etc Written to provide a comprehensive coverage of disease modifying mechanisms and compounds against neurodegenerative diseases Provides a drug discovery application oriented perspective evaluating targets and candidates for their overall therapeutic potential Provides discipline specific chapters medicinal chemistry target validation preclinical and clinical development Provides an overview on a number of molecular mechanisms e g phosphorylation chaperon refolding ubiquitination autophagy microtubule transportation protease cleavage etc with relevance for any disease area Contains a more thorough description of the therapeutic relevance of 10 specific molecular targets

Protein Misfolding in Neurodegenerative Diseases Robert D. E. Sewell, 2007-12-03 Current research suggests that neurodegenerative diseases such as Alzheimer s Parkinson s Huntington s and Creutzfeldt Jacob may be linked to disorders in protein shape referred to as protein misfolding Continued study in this area could lead to promising advances in future treatment of these diseases This groundbreaking text describes

the latest findings regarding protein misfolding in the context of it being a marker and perhaps a cause in neurodegenerative diseases Comprehensive coverage includes the diverse biochemical targets markers for each disease the currently limited success of drug therapies and the cutting edge research that could lead to more promising treatments *Protein folding and misfolding: neurodegenerative diseases* Judit Ovádi,Ferenc Orosz,2008-12-21 Offering all the latest in the study of neurodegenerative diseases this book reviews the molecular events initiated by unfolded or misfolded proteins leading to conformational human diseases especially those found in Parkinson s and Alzheimer s diseases Molecular Chaperones and Neurodegeneration Cintia Roodveldt,Tiago F. Outeiro,Janice E. Braun,2017-12-06 Molecular chaperones or heat shock proteins HSPs play essential roles in safeguarding structural stability and preventing misfolding and aggregation of proteins and maintaining the proteome functionality in the cell For over two decades until the present time new functions have been discovered and several molecular mechanisms have been elucidated for many chaperones while the field is being continuously challenged by new open questions Probably as a consequence of the increasing research on the molecular bases of neurodegenerative diseases and the realisation that many such disorders are linked to protein misfolding processes unleashing the roles and mechanisms of chaperones in the context of neurodegeneration has become a prime scientific goal This e book contains a diversity of reviews perspective and original research articles highlighting the importance and potential of this emerging subject **Protein Quality Control in Neurodegenerative Diseases** Richard I. Morimoto,Yves Christen,2012-12-13 The health of the proteome depends upon protein quality control to regulate the proper synthesis folding translocation and clearance of proteins The cell is challenged constantly by environmental and physiological stress aging and the chronic expressions of disease associated misfolded proteins Substantial evidence supports the hypothesis that the expression of damaged proteins initiates a cascade of molecular events that leads to Alzheimer s disease Parkinson s disease amyotrophic lateral sclerosis Huntington s disease and other diseases of protein conformation *Template-directed Protein Misfolding in Neurodegenerative Disease* William Clay Guest,2012

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Introduction

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