

Thomas Böldicke *Editor*

# Protein Targeting Compounds

Prediction, Selection and Activity of  
Specific Inhibitors

 Springer

# Protein Targeting Compounds Prediction Inhibitors

**Thomas Böldicke**



## **Protein Targeting Compounds Prediction Inhibitors:**

**Protein Targeting Compounds** Thomas Böldicke, 2016-05-28 This book presents an overview of the most relevant protein knockdown techniques Readers will find a description of the generation and use of recombinant human antibodies ER intrabodies and peptides as well as a description of the working mechanism and potential of the inhibitory action of each of these molecules The book will also describe the selection and activity of a number of phosphatases aptamers and allosteric modulators focusing on A G protein coupled receptors The book starts with a chapter on the in silico prediction of target inhibitor interactions Key aspects of this book are Selection strategies for monoclonal and recombinant human antibodies Selection strategies for ER intrabodies peptides and aptamers Examples of the most efficient inhibitors and their applications in protein biochemistry and cell biology Antibodies for cancer therapy and inhibitors of angiogenesis The book will be of great interest to scientists and students working in the life sciences on all areas related to protein biology The variety of methods to modulate the activities of specific proteins which are introduced here will be of great benefit to the reader who is interested in general protein research or to readers who have very specific protein related questions

**Protein Targeting with Small Molecules** Hiroyuki Osada, 2009-07-31 Discover the link between the latest chemical biology approaches and novel drug therapies Protein Targeting with Small Molecules Chemical Biology Techniques and Applications takes readers beyond the use of chemical biology in basic research providing a highly relevant look at techniques that can address the challenges of biology and drug design and development This indispensable bench companion features up to date coverage of advances in chemistry and assesses their impact on developing new therapeutics making it ideal for chemical biologists and medicinal chemists who are developing small molecule drugs to target proteins and treat diseases In addition the book examines the full range of complex biological systems and their interrelationship with chemistry from the interaction of biological response modifiers with proteins to the chemical biology of cell surface oligosaccharides Distinguished by an overview of chemical biology that is reinforced and clarified by detailed examples and descriptions of techniques Protein Targeting with Small Molecules Chemical Biology Techniques and Applications Introduces key technologies and methods of chemical biology designed to detect the interactions of small molecules and proteins Facilitates the discovery of small molecules that bind to proteins and describes the molecules application in the investigation of biological processes Presents timely coverage of the development of fluorescent probes for small molecules as well as the generation of small molecule ligands and inhibitors Reviews important techniques such as chemical genomics target profiling immobilization technology detection methods chemical inhibition and structure based targeting Offers a compelling synopsis of data that underscores the recent progress made in the area of targeting proteins by small molecules

*Inhibitors of Protein-Protein Interactions* Ali Tavassoli, 2020-11-25 Protein protein interactions PPI are at the heart of the majority of cellular processes and are frequently dysregulated or usurped in disease Given this central role the inhibition of PPIs has been of significant interest as

a means of treating a wide variety of diseases. However, there are inherent challenges in developing molecules capable of disrupting the relatively featureless and large interfacial areas involved. Despite this, there have been a number of successes in this field in recent years using both traditional drug discovery approaches and innovative interdisciplinary strategies using novel chemical scaffolds. This book comprehensively covers the various aspects of PPI inhibition encompassing small molecules, peptidomimetics, cyclic peptides, stapled peptides, and macrocycles. Illustrated throughout with successful case studies, this book provides a holistic cutting edge view of the subject area and is ideal for chemical biologists and medicinal chemists interested in developing PPI inhibitors.

*Multi-targeted Tyrosine Kinase Inhibitors in the Treatment of Cancer and Neurodegenerative Disorders* Halilibrahim Ciftci, Belgin Sever, Luciano Saso, Rumiana Tzoneva, Valentina Onnis, 2024-08-07

Tyrosine kinases, which catalyze the transfer of a phosphate from ATP to a hydroxyl group of a tyrosine residue, play a plethora of roles in the regulation of diverse functions in normal cells, including cell growth, motility, differentiation, and metabolism. Moreover, they are also associated with oncogenesis. Tyrosine kinases are mainly classified as receptor tyrosine kinases and non-receptor tyrosine kinases, including crucial members. Epidermal growth factor receptor (EGFR) belongs to the ERBB family of receptor tyrosine kinases, along with three other closely related receptors, namely HER2, HER3, and HER4. EGFR and HER2 lead to autophosphorylation of the intracellular domain through tyrosine kinase activity, and subsequent stimulation of downstream cascade that may result in proliferation, suppression of apoptosis, metastasis, and angiogenesis. On the other hand, c-Abl (Abl1) is a non-receptor tyrosine kinase, which is also essential in the regulation of several antiapoptotic and proliferative signal transduction pathways. They have mainly been identified as important targets for several types of cancer, such as EGFR for non-small cell lung cancer, glioma, and colorectal cancer; HER2 for breast and colorectal cancers; and Abl for chronic myeloid leukemia. One of the major platforms that they have participated in is neurodegenerative disorders, such as Parkinson's disease, Alzheimer's disease, and amyotrophic lateral sclerosis. Aberrant activity of tyrosine kinases, in particular EGFR and Abl, have been reported to induce apoptosis and cell cycle arrest in response to a wide range of stimuli, resulting in neurodegeneration and neuroinflammation. The main goal of this Research Topic is to identify new and potent, effective tyrosine kinase inhibitors to be effective in cancer and neurodegenerative disorders.

**Spectral Techniques In Proteomics** Daniel S. Sem, 2007-03-30 Facilitating the innovation, development, and application of new spectroscopic methods in proteomics. *Spectral Techniques in Proteomics* provides a broad overview of the spectroscopic toolbox that can be used either with proteome or sub-proteome mixtures or with individual purified proteins studied in parallel. It gives a modest overview of

**Ovarian Cancer Targeted Medication: PARP Inhibitors, Anti-Angiogenic Drugs, Immunotherapy, and More, volume II** Zhaoqian Liu, Jing Wang, Naiyuan Wu, Guang Lei, Zhi-Bin Wang, 2025-02-21 Ovarian cancer is the malignancy with the highest rate of death from tumors of the female reproductive system, and the main treatment options are tumor reduction and post-surgical platinum-based chemotherapy. Unfortunately, some patients will

develop platinum resistance after multiple recurrences Given that ovarian cancer is a heterogeneous disease with complex molecular and genetic alterations the identification of specific molecular targets will be of great benefit in gaining a deeper understanding of the mechanisms of ovarian cancer development and progression As research into targeted therapies continues the treatment of ovarian cancer is gradually shifting from conventional chemotherapy to targeted therapies Targeted drugs such as PARP inhibitors and anti angiogenic drugs have become important modalities for the maintenance treatment of ovarian cancer At the same time targeted therapy also suffers from ineffective use high rates of adverse reactions and high prices With the development of next generation sequencing technology targeted therapeutic agents developed for specific molecules in ovarian cancer may provide a wider range of treatment options for ovarian cancer patients and offer new strategies for individualized treatment Immunotherapy as well as ferroptosis pathway modulation has also provided new ideas for targeted therapy in ovarian cancer

*Chemoinformatics Approaches to Structure- and Ligand-Based Drug Design* Adriano D. Andricopulo, Leonardo L. G. Ferreira, 2019-02-05 Chemoinformatics is paramount to current drug discovery Structure and ligand based drug design strategies have been used to uncover hidden patterns in large amounts of data and to disclose the molecular aspects underlying ligand receptor interactions This Research Topic aims to share with a broad audience the most recent trends in the use of chemoinformatics in drug design To that end experts in all areas of drug discovery have made their knowledge available through a series of articles that report state of the art approaches Readers are provided with outstanding contributions focusing on a wide variety of topics which will be of great value to those interested in the many different and exciting facets of drug design

*Advances in Virus Research* , 2024-06-17 Advances in Virus Research Volume 119 the latest release in this esteemed series presents the latest research and developments in viruses Provides the authority and expertise of leading contributors from an international board of authors Presents the latest release in Advances in Virus Research series

**Structural Biology in Drug Discovery** Jean-Paul Renaud, 2020-01-27 With the most comprehensive and up to date overview of structure based drug discovery covering both experimental and computational approaches Structural Biology in Drug Discovery Methods Techniques and Practices describes principles methods applications and emerging paradigms of structural biology as a tool for more efficient drug development Coverage includes successful examples academic and industry insights novel concepts and advances in a rapidly evolving field The combined chapters by authors writing from the frontlines of structural biology and drug discovery give readers a valuable reference and resource that Presents the benefits limitations and potentiality of major techniques in the field such as X ray crystallography NMR neutron crystallography cryo EM mass spectrometry and other biophysical techniques and computational structural biology Includes detailed chapters on druggability allostery complementary use of thermodynamic and kinetic information and powerful approaches such as structural chemogenomics and fragment based drug design Emphasizes the need for the in depth biophysical characterization of protein targets as well as of therapeutic

proteins and for a thorough quality assessment of experimental structures Illustrates advances in the field of established therapeutic targets like kinases serine proteinases GPCRs and epigenetic proteins and of more challenging ones like protein protein interactions and intrinsically disordered proteins      **Ovarian Cancer Targeted Medication: PARP Inhibitors, Anti-Angiogenic Drugs, Immunotherapy, and More** Zhaoqian Liu,Dan Liu,Jing Wang,Guang Lei,Naiyiyuan Wu,2023-06-29

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