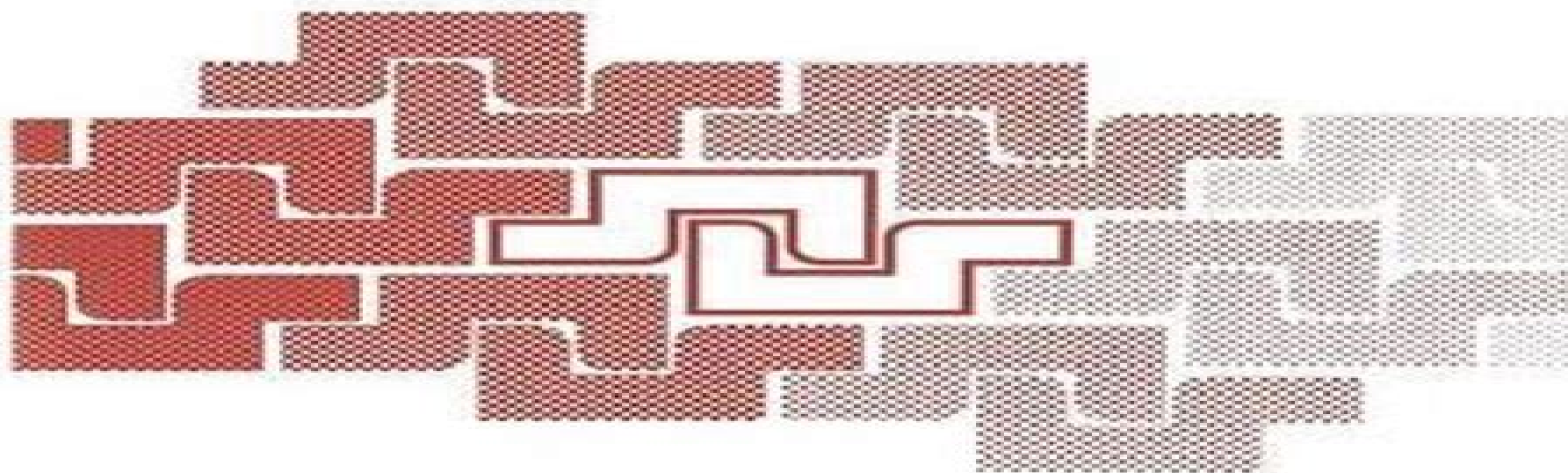




Targets for the Design of Antiviral Agents

Edited by
E. De Clercq and
R. T. Walker

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Targets For The Design Of Antiviral Agents

Erik De Clercq



Targets For The Design Of Antiviral Agents:

Targets for the Design of Antiviral Agents Erik De Clercq, 2013-06-29 This publication contains the Review Lectures presented at a joint NATO Advanced Study Institute and FEES Advanced Study Course held at Les Arcs Bourg Saint Maurice France from the 19th June 2nd July 1983 The Course entitled Targets for the Design of Antiviral Agents was in some ways a sequel to the NATO FEES Course held at SOGESTA near Urbino Italy from the 7th 18th May 1979 and published as volume A26 in this series During the subsequent four years we have witnessed the first of the new generation of antiviral compounds which are more efficacious and less toxic than the classical antiviral drugs reach the clinic and we felt that it was the right time to assess the future prospects of this very important and exciting field The vast majority of the drugs developed recently have proved active against various members of the herpesvirus family and elsewhere in this publication we learn that the cure for only rather few viral diseases such as the common cold influenza and herpes promises the return on investment required by the pharmaceutical industry However the aim of this Course was for eminent virologists to identify possible targets among the various virus classes against which the chemists could then design suitable therapeutic agents Recent advances with antiherpesvirus drugs have shown that a far greater selectivity and therapeutic index can be obtained than was previously thought to be possible Targets for the Design of Antiviral Agents Erik De Clercq, 2014-01-15

Biomedical Index to PHS-supported Research, 1990 **Current Catalog** National Library of Medicine (U.S.), 1985 First multi year cumulation covers six years 1965 70 **Aspartic Acid Proteases as Therapeutic Targets** Arun K. Ghosh, 2011-08-04 In this ground breaking practical reference the family of aspartic acid proteases is described from a drug developer's perspective The first part provides a general introduction to the family of aspartic acid proteases their physiological functions molecular structure and inhibition Parts two to five present various case studies of successful protease inhibitor drug design and development as well as current and potential uses of such inhibitors in pharmaceutical medicine covering the major therapeutic targets HIV 1 protease renin beta secretase gamma secretase plasmepsins and fungal proteases A ready reference aimed primarily at professionals in the pharmaceutical industry as well as for anyone studying proteases and their function **National Library of Medicine Current Catalog** National Library of Medicine (U.S.), **The Macrophage as Therapeutic Target** J. M. Aerts, 2003-07-10 with contributions by numerous experts

Biomedical Index to PHS-supported Research: pt. A. Subject access A-H, 1992 Understanding Viruses Teri Shors, 2009 Combining the molecular clinical and historical aspects of virology Understanding Viruses is a textbook for the modern undergraduate virology course The text provides an introduction to human viral diseases Additional chapters on viral diseases of animals the history of clinical trials gene therapy and xenotransplantation prions and viroids plant viruses and bacteriophages add to the coverage Jacket Biophysical and Computational Tools in Drug Discovery Anil Kumar Saxena, 2021-10-18 This book reviews recent physicochemical and biophysical techniques applied in drug discovery research

and it outlines the latest advances in computational drug design. Divided into 10 chapters, the book discusses about the role of structural biology in drug discovery and offers useful application cases of several biophysical and computational methods including time-resolved fluorometry, TRF with Förster resonance energy transfer (FRET), X-Ray crystallography, nuclear magnetic resonance spectroscopy, mass spectroscopy, generative machine learning for inverse molecular design, quantum mechanics, molecular mechanics, QM/MM, ONIOM, and quantum molecular dynamics (QMT) methods. Particular attention is given to computational search techniques applied to peptide vaccines using novel mathematical descriptors and structure and ligand-based virtual screening techniques in drug discovery research. Given its scope, the book is a valuable resource for students, researchers, and professionals from pharmaceutical industry interested in drug design and discovery.

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