# Synthetic Peptides

Approaches to Biological Problems

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## Approaches to Biological Problems

Proceedings of a Glaxo — UCLA Colloquium held at Park City, Utah January 31 - February 4, 1988

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### **Preface**

The explosive growth in applications of synthetic peptides to the elucidation of biological problems provided the impetus to organize the UCLA Colloquium on Synthetic Peptides: Approaches to Biological Problems, held at Park City, Utah, January 31–February 4, 1988. Synthetic peptides, which now are widely accessible by the solid-phase synthesis method developed by Professor Bruce Merrifield 25 years ago, have become powerful tools with which to define the structure and functions of such biological activities as antigen—antibody, hormone—receptor, substrate—enzyme, and protein—DNA interactions. It is anticipated that, as the availability of amino acid sequences derived from genomic DNA increases, the understanding of the use of synthetic peptides—their syntheses, structures, and applications—will facilitate the approach to solving many biological problems.

The importance of peptides and synthetic chemistry on synthetic vaccines, mapping of immunodominant epitopes, and understanding of antigen—antibody interactions was reflected in the decision to hold this conference concurrently with the UCLA Symposium on Technological Advances in Vaccine Development. In his Keynote Address on Catalytic Antibodies, Dr. Richard Lerner outlined the influence of synthetic chemistry on biological problems. Approximately half of the contributions to this volume involve work related

to synthetic peptides in the field of immunology.

This proceedings volume, **Synthetic Peptides: Approaches to Biological Problems**, includes 25 articles in six sections, just as they appeared in the conference program: methods and applications, prediction of peptide and protein structures, synthetic peptide-based vaccines, determinations of antigenic domains, bioactive conformations of peptide hormones, and peptide hormones and growth factors.

During the preparation of this volume, I was greatly saddened by the untimely passing of my co-organizer for this meeting, Professor Tom Kaiser, at the age of 50, on July 18, 1988. Tom will be remembered as a valued colleague, an energetic scientist, and for his brilliant work on the structural bases of

biological activities of peptides and proteins.

We gratefully acknowledge Glaxo, Inc., for sponsorship of this meeting. Additional support was received from Smith Kline Beckman; Pharmacia LKB Biotechnology, Inc.; Merck Sharp & Dohme Research Laboratories; Applied Biosystems, Inc.; and O.C.S. Laboratories. In addition, we wish to thank the UCLA Symposia staff, especially Robin Yeaton and Betty Handy, for valuable and patient assistance.

James P. Tam

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