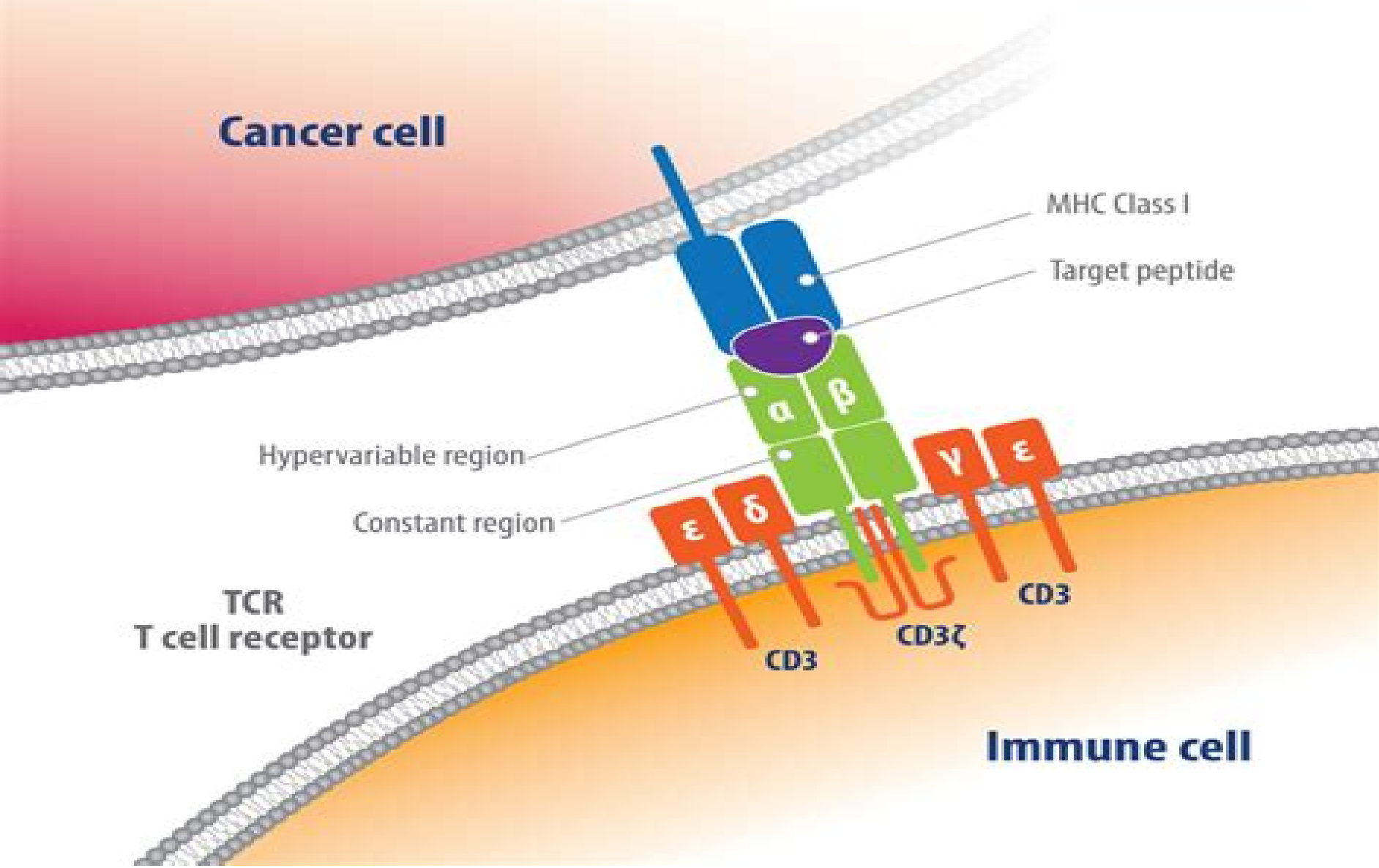


T Cell Receptors (TCR)



T Cell Receptors

Peter van den Elsen



T Cell Receptors:

The T-Cell Receptors Tak W. Mak, 2012-12-06 The importance of thymus dependent cells or T cells in the generation of a successful immune response was first realized in the early sixties In the following two decades a succession of elegant experiments established the antigen specificity of T cells and their ability to perform both as regulatory and effector cells T cells were shown to be essential in most immune reactions playing a crucial role in augmenting the activity of effector T and B cells against foreign antigen as well as in the suppression of effector activity against self antigens The means by which T cells differentiate foreign from self antigens is based on their recognition of antigen almost exclusively in the context of self major histocompatibility complex products unlike B cells which recognize an antigen alone It is this recognition mediated by the T cell receptor that sets into motion the diverse cell cell interactions which control the differentiation and regulation of the immune response Although its importance was well established the molecular nature of the T cell receptor remained elusive for two decades Many hypotheses as to its structure and precise function were put forward using immunoglobulin as a basis for conjecture but the Holy Grail of Immunology remained ephemeral until three years ago In the ensuing years both immunologists and molecular biologists have contributed to an explosion of data unsurpassed by any previous period in the field

T Cell Receptors John I. Bell, Michael J. Owen, M. J. Owen, Dr. Elizabeth Simpson, 1995 T cell receptor genes covers biochemical functional and genetic aspects of this key family of immunological molecules It should be essential reading for all those interested in the TCR and its function in health and disease It deals with the TCR protein its functional interactions with the signal transduction machinery of the T cell and with the structure and organization of the TCR genes It is the only comprehensive volume on the TCR and is written by leaders in the field Particularly useful is the tabulation and nomenclature of all the known TCR alpha and beta variable regions

The T Cell Receptor FactsBook Marie-Paule Lefranc, Gerard Lefranc, 2001-07-13 The T Cell Receptor FactsBook contains entries on all the 176 functional variable diversity joining and constant regions of the human T cell receptor including alpha beta gamma and delta loci Introductory chapters summarize information of T cell receptor chain synthesis chromosomal location and an overview of the human T cell receptor loci

Antigen Binding Molecules: Antibodies and T-Cell Receptors, 1996-12-13 Topics Covered Include X ray crystallography of ligands Catalytic antibodies Nature of the antigen Antibody binding sites Maturation of the immune response Computational biochemistry of antibodies and T cell receptors Antigen specific T cell receptors and their reactions

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Immunology Richard A. Goldsby, 2003 The new edition of the acclaimed bestseller always praised for offering cutting edge material in the context of landmark experiments in a student friendly format built on pedagogy not usually found in immunology texts

Antigen-specific T Cell Receptors and Factors, 1987 *The Human T-Cell*

Receptor Repertoire and Transplantation Peter van den Elsen, 2013-06-29 these analyses it became clear that the MHC class I molecule comprised a distinct groove on the external side of the molecule. The sides of the groove are formed by the α and β domains and a floor which is formed by 8 anti parallel β strands. The various polymorphic residues as determined from DNA sequence analysis are localized within these α helices and β plated sheets within the groove. More importantly these analyses also revealed the presence of electron dense material in the groove. This material was subsequently identified as a linear peptide of 8-10 amino acids long. High resolution crystallographic analyses of the class I MHC structure have revealed the existence of so called pockets within the grooves of the MHC class I molecules. These pockets designated A-F exhibited allele specificity and are directly involved in the binding of the peptide primarily through interaction with the dominant anchor residues as found in MHC class I associated peptides 6-7-9-11. The class II MHC antigens consist on the cell surface of a 34 kD α chain non covalently associated with a 28 kD β chain. With the exception of the DR α chain all other MHC class II α and β chains are polymorphic. Kuby Immunology Thomas J. Kindt, Richard A. Goldsby, Barbara A. Osborne, Janis Kuby, 2007 Originally authored by the award winning author Janis Kuby Immunology remains the best selling textbook for the undergraduate course. The first and only true textbook written by professors who teach the undergraduate course it presents the most current concepts in an experimental context with clinical advances highlighted in boxes supported by the kind of helpful pedagogical tools that other books do not provide.

Investigating and harnessing T-cell functions with engineered immune receptors and their ligands Bruno Laugel, 2015-01-22 T cells are an essential component of the immune system that provide protection against pathogen infections and cancer and are involved in the aetiology of numerous autoimmune and autoinflammatory pathologies. Their importance in disease, the relative ease to isolate, expand and manipulate them ex vivo have put T cells at the forefront of basic and translational research in immunology. Decades of study have shed some light on the unique way T cells integrate extrinsic environmental cues influencing an activation program triggered by interactions between peptide-MHC complexes and the antigen recognition machinery constituted of clonally distributed T cell receptors and their co-receptor CD4 or CD8. The manipulation of these molecular determinants in cellular systems or as recombinant proteins has considerably enhanced our ability to understand antigen specific T cell activation to monitor ongoing T cell responses and to exploit T cells for therapy. Even though these principles have given numerous insights in the biology of CD8 T cells that translate into promising therapeutic prospects as illustrated by recent breakthroughs in cancer therapy they have proven more challenging to apply to CD4 T cells. This Research Topic aims to provide a comprehensive view of the recent insights provided by the use of engineered antigen receptors and their ligands on T cell activation and how they have been or could be harnessed to design efficient immunotherapies. **Medical Subject Headings** National Library of Medicine (U.S.), 1998 **Life** David E. Sadava, 2008 This text aims to establish biology as a discipline not just a collection of facts. Life develops students

understanding of biological processes with scholarship a smooth narrative experimental contexts art and effective pedagogy
Cumulated Index Medicus ,1989 **Index Medicus** ,2004 Vols for 1963 include as pt 2 of the Jan issue Medical subject headings **Biomedical Index to PHS-supported Research: pt. A. Subject access A-H** ,1992 **Medical Subject Headings** ,2004 Medical Subject Headings ,1998 **Fundamental Immunology** William E. Paul,2008 Now thoroughly revised and updated this comprehensive up to date text is ideal for graduate students post doctoral fellows microbiologists infectious disease physicians and any physician who treats diseases in which immunologic mechanisms play a role

Translation of genetically engineered T cells in cancer immunotherapy Shao-An Xue,He Huang,Hakim Echchannaoui,Ralf-Holger Voss,2023-10-06 New Frontiers in Gene-Modified T Cell Technology Ignazio Caruana,Francesca Del Bufalo,Rayne Rouce,Shigeki Yagyu ,Paul G. Schlegel,2024-06-13 The development clinical translation and recent efficacy of novel gene therapies targeting refractory malignancies has led to research that extends this technology to a variety of infectious and rheumatological diseases Unlike conventional drugs or antibodies T cells have the potential to target and exert effector function in response to disease in a dynamic manner acting as a living drug The most efficacious form of gene modified T cells to date is the chimeric antigen receptor CAR modified T cell which redirects the specificity of T cells to an antigen expressed by tumor cells Clinical experience with autologous CAR T cells primarily in hematologic malignancies has underscored the feasibility and safety of the approach while also demonstrating dramatic and sustained antitumor effects through mechanisms orthogonal to those of traditional anticancer therapies However several challenging obstacles must be surmounted in order to improve the broader efficacy of this approach **Biomedical Index to PHS-supported Research** ,1993

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