

STEROID HORMONE RESISTANCE

Mechanisms and Clinical Aspects

Steroid Hormone Resistance Mechanisms And Clinical Aspects

J. Larry Jameson



Steroid Hormone Resistance Mechanisms And Clinical Aspects:

Steroid Hormone Resistance George P. Chrousos, D. Lynn Loriaux, Mortimer B. Lipsett, 2012-12-06 This volume represents the first attempt to present in one place the clinical syndromes and the pathophysiologic basis for the resistance states to each of the classes of steroid hormones Glucocorticoids mineralocorticoids androgens estrogens progesterone and vitamin D have widely diverse roles ranging from the control of homeostasis to reproduction and bone formation They are similar in that they share a chemical structure and that their action is in the cell nucleus where they induce transcription of specific genes leading to synthesis of function specific proteins Clinical syndromes of steroid hormone resistance to androgens complete and partial testicular feminization aldosterone pseudo hypoaldosteronism and vitamin D vitamin D dependent rickets type II have been known for many years Progesterone and glucocorticoid resistance syndromes have been described only recently Resistance to estrogens has not been reported in man or in animals It is hoped that a detailed reexamination of what is known about each of these conditions at the clinical and molecular levels will enhance our understanding of the function of these hormones and their mechanisms of action New insight and research initiatives should result G P Chrousos D L Loriaux M B Lipsett vii **ACKNOWLEDGMENTS** The contents of this volume are based in part on the proceedings of an International Conference held in Bethesda in the summer of 1984 This conference was sponsored by the National Institute of Child Health and Human Development Bethesda Maryland

Hormone Resistance and Other Endocrine Paradoxes Margo P. Cohen, Piero P. Foa, 2012-12-06 Endocrinology and Metabolism Progress in Research and Clinical Practice is a new series that has been designed to present timely critical reviews of constantly evolving fields to provide practical and up to date guidance in the solution of pertinent clinical problems to offer an alternative to the laborious search of the literature and the often frustrating reading of highly technical articles and to translate the language of the laboratory into that of the practice of medicine We think that this volume and those to come will prove useful to physicians and to physicians in training as well as to investigators in a wide variety of specialties in short to anyone who seeks answers to questions in endocrinology and metabolism The first chapter of this volume could well serve as a general introduction to the entire series It points out how our growing understanding of the molecular basis of biologic communication has led to the discovery of a growing number of clinical syndromes as well as to the realization that phenotypically similar diseases may have radically different pathogenetic mechanisms and thus may require radically different therapeutic strategies

National Library of Medicine Current Catalog National Library of Medicine (U.S.), Current Catalog National Library of Medicine (U.S.), First multi year cumulation covers six years 1965 70

Kinins IV Lowell Greenbaum, 2012-12-06 Scientists from 25 countries came together at the Hyatt Regency Hotel on October 21 25 1984 for the 4th International Kinin Congress in the beautiful city of Savannah Georgia Many of the delegates enjoyed southern hospitality for the first time The friendly city with its streets lined with the Live Oak Tree symbol of the Congress the balmy

weather and the excellent facilities of the hotel set the stage for scientific events and exchange that proved so successful The organization of the meeting was the result of many hours days and weeks of effort by many including from Augusta Drs James H Sutherland John Catravas William Davis Jr and Hiroshi Okamoto and from Charleston SC Julie Chao Ronald Mayfield and Donald Miller Special thanks go to Ms Cher Cornett of the Department of Medical Illustration at the Medical College of Georgia for her talent in the design of the logo as well as the graphics of the program and abstracts We are indeed indebted to Dr Paul Brucker Director of the Division of Health Communications at the Medical College of Georgia for his considerable aid in spending time with us to arrange the programming format and continuity Ms Sandra Usry provided very excellent and devoted secretarial help for several years while the Congress was in the making Myocardial and Skeletal Muscle Bioenergetics Nachman Brautbar,2013-03-08 **Biochemical Basis of Functional Neuroteratology** ,1988-07-01 Biochemical Basis of Functional Neuroteratology **Scientific Directory and Annual Bibliography** National Institutes of Health (U.S.),1987 **Oxygen Transport to Tissue VIII** Ian S. Longmuir,2012-12-06 In this volume the policy of review by anonymous referees and minor correction by the editor has been continued but perhaps should not be extended without an agreed policy statement by the Society Our choice is minimal revision with rapid publication or proper review with some delay in publication The editor wishes to express his gratitude to Ann Richardson Joann Fish Lance Johnson and Philip Weinbrecht for their invaluable help in the preparation of this volume Ian S Longmuir v BRIEF HISTORY OF ISOTT The Society endured a long gestation period During the 1960s its formation was discussed at a number of international meetings devoted to oxygen in biological systems Prominent among a great number of such gatherings were those held at the Institute of Diseases of the Chest London 1960 Bedford College London 1963 Queen Elizabeth College London 1963 and the Seventh Bad Oeynhausen Conference 1967 At first reservations were expressed about the desirability of forming a highly specialized society which might not achieve the critical mass necessary for its continued existence However the 1971 meeting in the Max Planck Institut fur Arbeitsphysiologie Dortmund answered these doubts and Dr Melvin Knisely commenced planning a very successful inaugural meeting in South Carolina At this meeting in 1973 in Charleston and Clemson the Society was formed with the customary remit of the promotion of scientific exchanges *Hormone Resistance Syndromes* J. Larry Jameson,1999-01-26 Leading clinicians review in depth the clinical and molecular aspects of the hormone resistance syndromes caused by receptor mutations These mutations are involved in a large number of disorders including dwarfism Laron syndrome pseudohypoparathyroidism insulin resistance resistance to thyroid hormone glucocorticoid resistance hereditary resistance to vitamin D and estrogen resistance Throughout the book s distinguished contributors not only summarize the clinical features of these and related disorders but also illuminate their molecular bases *Hormone Resistance Syndromes* provides an authoritative summary and standard reference for all those clinicians and researchers actively studying these mutations today and affords a unique opportunity to understand hormone action in normal physiology

Recent Progress in Hormone Research Roy O. Greep, 2013-10-22 Recent Progress in Hormone Research Volume 41 covers the proceedings of a Laurentian Hormone Conference held in late August 1984 at the Homestead in Hot Springs Virginia The book presents papers on the hormone receptor interactions the biosynthesis secretion metabolism and mechanism of action of the steroid hormones and the mechanisms of action of thyroid stimulating hormone TSH and TRH The text also includes papers on the molecular characterization of a brain specific mRNA and the factors affecting changes in frequency and amplitude of GnRH pulses and the resulting functional consequences in various mammals including humans with disorders of fertility Papers on the biological heritage of mammalian endocrinology such as the actions of urotensin I in mammals and fishes and the clinical implications of the glycosylation and posttranslational processing of the TSH are also encompassed Endocrinologists neuroscientists biochemists biophysicists and scientists involved in hormone research will find the book invaluable

Receptor Mediated Antisteroid Action M. K. Agarwal, 2019-05-20 No detailed description available for Receptor Mediated Antisteroid Action

Lipoprotein Deficiency Syndromes Aubie Angel, 2012-12-06 Current interest in lipoprotein deficiency states stems from the growing realization of their importance in the etiology of premature coronary heart disease While hypercholesterolemia and coronary heart disease risk are strongly correlated in their etiologic relationship it is becoming equally clear that deficiencies in HDL whether congenital or acquired also enhance the risk for the future development of coronary atherosclerosis This has led to renewed attention to the lipid hypothesis and realization of the fact that each lipoprotein class and apoprotein species has specific functions in the transport and cellular uptake of various lipids It is a truism that a biochemical correlate of disease once identified is subsequently recognized with increasing frequency in clinical medicine The story of HDL was no exception Indeed hypoalphalipoproteinemia appears to be a disease of high prevalence approaching and perhaps even exceeding that of familial hypercholesterolemia Its clinical significance escaped our notice for many years largely due to a heavy emphasis on hypercholesterolemia and to difficulties in measuring HDL reliably

Molecular Structure and Biological Activity of Steroids Martin Bohl, 2018-01-18 Molecular Structure and Biological Activity of Steroids focuses on the identification of steroid structural features that control particular biological effects Steroid geometry plays a critical role in steroid activity therefore the results of X ray crystallographic NMR spectroscopic and theoretical studies on structure activity relationships form an integral part of the book Well established observations and proposed models are presented summarizing knowledge of molecular features e g steroid backbone structures side chain constitutions substitution sites configurations hydrogen bonding patterns that can affect steroid action The book also provides overviews about recent progress in various frontier fields of steroid research including such areas as sex steroids glucocorticoids aromatase inhibitors vitamin D metabolites brassinosteroids neuromuscular blocking agents and cardiotonic steroids Molecular Structure and Biological Activity of Steroids is a must have resource for biochemists pharmacologists endocrinologists biophysicists and others actively involved in steroid research

Nutritional and

Toxicological Significance of Enzyme Inhibitors in Foods Mendel Friedman, 2013-03-09 Soybean proteins are widely used in human foods in a variety of forms including baby formulas flour soy protein concentrates soy protein isolates soy sauces textured soy fibers and tofu The presence of inhibitors of digestive enzymes in soy proteins impairs nutritional quality and possible safety of this important legume Normal processing conditions based on the use of heat do not completely inactivate these inhibitors so that residual amounts of plant protease inhibitors are consumed by animals and man Inhibitors of digestive enzymes are present not only in legumes such as soybeans lima beans and kidney beans but also in nearly all plant foods including cereals and potatoes albeit in much smaller amounts The antinutritional effects of inhibitors of proteolytic enzymes have been widely studied and can be ameliorated by processing and or sulfur amino acid fortification A more urgent concern is reports that rats fed diets containing even low levels of soybean derived inhibitors which are found in foods such as soy based baby formulas may develop over their lifespan pancreatic lesions leading eventually to neoplasia or tumor formation On the other hand recent studies suggest that certain enzyme inhibitors from plant foods may prevent cancer formation in other tissues A key question therefore is whether inhibitors from plant foods constitute a human health hazard

Physician's Guide to the Laboratory Diagnosis of Metabolic Diseases N. Blau, M. Duran, M.E. Blaskovics, K.M. Gibson, 2012-12-06 Delay and mistakes in the diagnosis of inherited metabolic diseases may have devastating consequences Reference laboratory data are scattered and clinical descriptions of rare conditions are hard to locate This book describes 298 disorders grouped into 35 chapters according to the type of condition Within each group of disorders chapters provide tables of pertinent clinical findings as well as reference and pathological values for crucial metabolites Relevant metabolic pathways and diagnostic flow charts are included There are four indices to make the book as user friendly as possible Disorders index Signs and symptoms index Organs index and Tests index The Physician's Guide provides paediatricians and other physicians with a unique aid to help them select the correct diagnosis from a bewildering array of complex clinical and laboratory data

Inheritance of Kidney and Urinary Tract Diseases Adrian Spitzer, Ellis D. Avner, 2012-12-06 Genetic disorders have emerged as a prominent cause of morbidity and mortality among infants and adults As many as 10% to 20% of hospital admissions and at least 10% of the mortality in this age group are due to inherited diseases There are at least two factors that have brought genetic disorders into the forefront of pediatrics One is a great reduction in childhood mortality due to infections and nutritional deficiency states and the other is the rapid progress made in the identification of genetic defects Amniocentesis chorionic villus sampling and recombinant DNA technology have already had a tremendous impact on the practice of medicine This is why the first two chapters of this volume are dedicated to general principles of molecular genetics and to a description of the techniques used to diagnose genetic disorders at the DNA level The relevance of this new area of science to the study of inherited renal diseases is reflected in the large body of knowledge that has been generated regarding the association between various glomerular nephritides and genetic markers such as the HLA system and even

more impressively in the direct or indirect identification of abnormal genes or gene products in Alport's syndrome, autosomal dominant polycystic kidney disease and Lowe's syndrome. These discoveries figure prominently in the pages of this book. Yet the progress we have made has barely scratched the surface of the problem.

Recent Progress in Hormone Research

James H. Clark, 2013-10-22. Recent Progress in Hormone Research Volume 46 provides a superior summary of the developments in the field of hormone research. The book discusses the molecular basis of androgen insensitivity, the tissue-specific expression of the growth hormone gene and its control by growth hormone factor 1 and the molecular characterization of mammalian tachykinin receptors and a possible epithelial potassium channel. The text also describes the properties of the guanylate cyclase receptor family, insulin-like growth factor binding proteins and growth hormone receptor and binding protein. Mutations in the insulin receptor gene in genetic forms of insulin resistance, the characteristics of the cAMP response unit and the role and secretion of inhibin in the rat are also considered. The book further tackles the structure of the lutropin/choriogonadotropin receptor and the gene for multiple endocrine neoplasia type 2a. Endocrinologists, physiologists and biochemists will find the text invaluable.

Cumulated Index Medicus, 1996. *Physician's Guide to the Diagnosis, Treatment, and Follow-Up of Inherited Metabolic Diseases*. Nenad Blau, Marinus Duran, K. Michael Gibson, Carlo Dionisi-Vici, 2014-07-08. This book combining and updating two previous editions is a unique source of information on the diagnosis, treatment and follow-up of metabolic diseases. The clinical and laboratory data characteristic of rare metabolic conditions can be bewildering for both clinicians and laboratory personnel. Reference laboratory data are scattered and clinical descriptions may be obscure. The Physician's Guide documents the features of more than five hundred conditions grouped according to type of disorder, organ system affected (e.g. liver, kidney, etc.) or phenotype (e.g. neurological, hepatic, etc.). Relevant clinical findings are provided and pathological values for diagnostic metabolites highlighted. Guidance on appropriate biochemical/genetic testing is provided. Established experimental/therapeutic protocols are described with recommendations on follow-up and monitoring. The authors are acknowledged experts and the book will be a valuable desk reference for all who deal with inherited metabolic diseases.

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