

Sex Chromosomes Turner Syndrome

Sex Chromosomes Ursula Mittwoch 2014-06-28 *Sex Chromosomes* focuses on the study of sex chromosomes, including human chromosomal abnormalities, behavior and characteristics of chromosomes, and cell division. The book first offers information on the chromosomal basis of sex determination, as well as development of the cell theory, mitosis, fertilization, meiosis, and discovery of sex chromosomes. The publication also ponders on the mitosis, meiosis, and formation of gametes. Discussions focus on the special characteristics of sex chromosomes, abnormalities of cell division, and sexual differentiation. The manuscript reviews sex chromosomes in plants, *Drosophila*, and *Lepidoptera*. The book also examines sex-chromosome mechanisms that differ the classic type; sex chromosomes in fishes, amphibia, reptiles, and birds; and sex chromosomes in man. Discussions focus on normal human sex chromosomes, Turner's syndrome, Klinefelter's syndrome, true hermaphrodites, testicular feminization, and pseudohermaphrodites. Sex chromosomes in mammals other than man, including monotremata, marsupialia, insectivora, rodentia, and carnivora, are discussed. The publication is a dependable reference for readers interested in the study of sex chromosomes.

Human Chromosome Methodology Jorge J. Yunis 2016-01-22 *Human Chromosome Methodology* fills the need for an authoritative and up-to-date treatise which would serve as a text and reference for advances in human cytogenetics. The book includes readily comprehensible chapters that cover each phase of laboratory investigation from the preparation of materials for sex chromatin and chromosome techniques for bone marrow, blood, skin, and gonadal specimens to the subject of autoradiography and chromosome identification. Included also are guides to microscopy and photomicrography as well as an up-to-date treatment of chromosomes in disease. It is hoped that this volume will serve as an adequate guide to laboratory techniques and their applications for research workers, students of genetics, and members of the medical profession involved in setting up a laboratory of cytogenetics.

Turner Syndrome Theola Stubby 2021-03-31 Turner syndrome, a condition that affects only females, results when one of the X chromosomes (sex chromosomes) is missing or partially missing. Turner syndrome can cause a variety of medical and developmental problems, including short height, failure of the ovaries to develop, and heart defects. This book is storytelling about a girl who found out she has a form of Turner syndrome when she was 16. It was difficult to see how anything positive was going to come out of the new journey she was faced with. The journey has not always been easy but, through everything, she has become a stronger person and learned a lot.

Sex Hormone Replacement Therapy Henry Burger 2013-03-09 *Sex Hormone Replacement Therapy* addresses important issues in contemporary endocrinology. Its major emphasis is on the consequences of the menopause and on androgen deficiency in the absence of overt disease of the hypothalamus and/or pituitary in men. The use of hormone therapy for the primary prevention of cardiovascular disease in post-menopausal women has become an area of major debate, and the pros and cons are examined in detail. Secondary prevention also remains controversial and is addressed in this volume. An issue of particular concern is the treatment of estrogen deficiency symptoms in women with the diagnosis of breast cancer. Whether older men with some androgen deficiency symptoms and equivocally low plasma testosterone concentrations should be treated with androgens is also a matter of great debate, as is the efficacy and safety of plant estrogens in the treatment of menopause-related complaints. *Sex Hormone Replacement Therapy* will be of interest to students and non-specialists, as well as a wide range of medical practitioners including endocrinologists, gynecologists, rheumatologists and cardiologists.

The 7 Sexes Elof Axel Carlson 2013 Few of us know much about the biology of sex determination, but what could be more interesting than to

discover how we are shaped into males and females? In this book, Elof Carlson tells the incredible story of the difficult quest to understand how the body forms girls and boys. Carlson's history takes us from antiquity to the present day to detail how each component of human reproduction and sexuality was identified and studied, how this knowledge enlarged our understanding of sex determination, and how it was employed to interpret such little understood aspects of human biology as the origin of intersex births.

Aneurysms-Osteoarthritis Syndrome Denise van der Linde 2016-10-03 *Aneurysms-Osteoarthritis Syndrome: SMAD3 Gene Mutations* is a first-of-its-kind compilation of the genetic discovery, research, and care associated with AOS. With the field of genetically triggered aortopathies growing, this important reference will compile the newest discoveries in this field, allowing cardiologists, cardio-thoracic surgeons, clinical geneticists, vascular surgeons, orthopedic surgeons, and researchers to gain the knowledge they need without having to gather the data from various sources. Coverage includes genotype and phenotype correlations, the functional role of SMAD3, and insights into the role of TGFbeta signaling in aortic disease. The book will increase knowledge about AOS, providing awareness and better patient care for this aggressive disease. Covers Aneurysms-Osteoarthritis Syndrome, from genetic discovery to patient care Contains clinical management guidance on optimal cardiovascular treatments and surgery Explains the autosomal dominant syndromes caused by mutations in the SMAD3 gene Identifies the key features of this syndrome, including arterial aneurysms and tortuosity, early onset arthritis, and mild craniofacial features

Genetic Disorders of Human Sexual Development Leonard Pinsky 1999 There have been many advances in understanding mammalian sex determination and differentiation during the last decade. Using these advances to elucidate clinical conditions of abnormal sexual development, the authors of this book bring together expertise in molecular endocrinology, molecular genetics, and dysmorphology. This is not a conventional textbook. It is seldom dogmatic, frequently presents alternatives, highlights speculation, raises questions, and attempts to provide answers. This book will be welcomed by medical geneticists, genetic counselors, endocrinologists, gynecologists, urologists, and students who need fully-referenced information about the genetic aspects of human sexual maldevelopment in order to better manage their patients and their patients' families.

The Chromosome Disorders Gordon Howard Valentine 1969

Chromosomes of Man Hans Zellweger 1977

The Danish Cytogenetic Central Register Johannes Nielsen 1980

Sex Chromosome Abnormalities And Human Behavior Daniel B Berch 2019-07-11 This volume is based on a symposium, "Cognitive and Psychosocial Dysfunctions Associated with Sex Chromosome Abnormalities," presented at the 1986 Annual Meeting of the American Association for the Advancement of Science. It contains reports from individual research groups and a psychological study.

Chromosome Abnormalities and Genetic Counseling R. J. McKinlay Gardner 2003-08-28 Chromosomal abnormalities can cause disability in children, and reproductive difficulty in parents. Many parents and couples seek genetic counseling in order to learn why they, or a relative, may have had a child with a particular collection of medical problems and/or intellectual disability. There may have been a history of multiple miscarriage, or infertility. They may want to know the outlook for a pregnancy, and what the risks might be. These and other questions concerning chromosome abnormalities are addressed in this standard text, which will be of interest to genetic counselors, medical geneticists, pediatricians and obstetricians, infertility specialists, and laboratory cytogeneticists. This third edition has been thoroughly updated, and is richly illustrated and fully referenced. New chapters have been written on preimplantation diagnosis and on reproductive risks due to environmental agents. The practical applications of recent advances in molecular cytogenetics are noted. The book will give counselors the information that will enable them to help concerned parents accommodate to their particular "chromosomal situation", and to determine what may be, for them, the best course of action.

Symposium on Nuclear Sex D. Robertson Smith 2013-10-22 Symposium on Nuclear Sex covers the proceedings of the 1957 Symposium on Nuclear Sex, held at King's College Hospital Medical School. This book is organized into three sections encompassing 22 chapters that consider the problem of developmental sex anomalies and certain cytological aspects. After a brief presentation of the history of chromosomal sex and sex anomalies, this book goes on exploring the genetic and cytological observations related to nuclear sex, with a particular emphasis on the genetic implications of nuclear sexing. The second part deals first with the fundamentals of sex chromosomes and the development of human intersexes. The discussion then shifts to problems in intersexuality, children of homosexuals, transvestism, sexual development disturbances, and the nature of the genetic defect in Klinefelter's syndrome. The third part covers studies of the nuclear sex of a group of teratomas and other sex tumors. This book will prove useful to workers and researchers in the fields related to nuclear sexing including anatomy, biochemistry, biology, clinical medicine, cytology, endocrinology, genetics, pathology, physiology, psychology, and surgery.

Somatropin (Genotropin) for Subcutaneous Injection 2014 Turner syndrome (TS) is characterized by the absence of all or part of a normal second sex chromosome in females. Around 50% of patients with TS have sex chromosome abnormalities, while the remaining half have one sex chromosome. Mutations of chromosomes in patients with TS lead to a range of clinical features including, but not limited to, short stature. Adult height of untreated women with TS is approximately 20 cm shorter than that of adult women in the general population, with the average height being around 143 cm. Recombinant human growth hormone (rhGH), also called somatropin, is used to accelerate short-term growth in girls with TS. Genotropin is one of several somatropin products available in Canada and is indicated for the treatment of short stature associated with TS in patients whose epiphyses are not closed at a dose of 0.33 mg/kg per week, divided into six to seven doses. The objective of this report was to conduct a systematic review of the benefits and harms of Genotropin compared with other available somatropin products for the treatment of short stature associated with TS.

Diabetes Associated with Single Gene Defects and Chromosomal Abnormalities F. Barbetti 2017-03-21 This volume, written by renowned experts, provides complete coverage of the main genetic conditions associated with diabetes. Divided into five sections, it offers insights into genetic defects involving the pancreatic beta cell, extreme insulin resistance, ciliopathies, obesity and glucose metabolism, chromosomal defects, and other genetic conditions associated with increased susceptibility to diabetes. Other topics include the various subtypes of monogenic diabetes, such as the neonatal form and the Wolfram syndrome, as well as chromosomal defects leading to complex conditions affiliated with diabetes, like Trisomy 21 or Prader-Willi syndrome. There are also chapters dedicated to the poorly explored relationships between metabolism and neurodegenerative disorders like Friedreich's ataxia and muscular dystrophy. This book is a reference for every pediatric and adult endocrinologist and diabetologist, even experienced ones, with an interest in the intricacies and protean aspects of disorders of glucose metabolism secondary to genetic diseases.

Human Cytogenetics John L. Hamerton 2013-10-22 Human Cytogenetics: Clinical Cytogenetics, Volume II presents the general theoretical principles and clinical aspects of cytogenetics, a branch of genetics that deals specifically with the study of the chromosomes. The volume focuses on the clinical cytogenetics of human. It discusses the sex chromosomes and their abnormalities and the abnormalities of sexual development and differentiation; mechanism of sex determination in mammals; major autosomal abnormalities found in human populations; and chromosome abnormalities in relation to human pregnancy wastage and chromosome changes in neoplasia. The book will be a great reference book for geneticists, cytogeneticists, pathologists, clinicians, and medical students.

Genetics of Sex Determination R.S. Verma 1996-04-23 The Genetical Theory of Natural Selection by R.A. Fisher (1930) dictated that sexual dimorphisms may depend upon a single medelian factor. This could be true for some species but his suggestion could not take off the ground as

gender in *Drosophila* is determined by the number of X chromosomes. Technical advances in molecular biology have revived the initial thinking of Fisher and dictate that TDF or SRY genes in humans or Tdy in mice are sex determining genes. The fortuitous findings of XX males and XY female, which are generally termed sex reversal phenomenon, are quite bewildering traits that have caused much amazement concerning the pairing mechanism(s) of the pseudoautosomal regions of human X and Y chromosomes at meiosis. These findings have opened new avenues to explore further the genetic basis of sex determination at the single gene level. The aim of the fourth volume, titled Genetics of Sex Determination is to reflect on the latest advances and future investigative directions, encompassing 10 chapters. Commissioned several distinguished scientists, all pre-eminent authorities in each field to shed their thoughts concisely but epitomise their chapters with an extended bibliography. Obviously, during the past 60 years, the meteoric advances are voluminous and to cover every account of genes, chromosomes, and sex in a single volume format would be a herculean task. Therefore, a few specific topics are chosen, which may be of great interest to scientists and clinicians. The seasoned scientists who love to inquire about the role of genes in sex determination should find the original work of these notable contributors very enlightening. This volume is intended for advanced students who want to keep abreast as well as for those who indulge in the search for genes of sex determination.

The Nature and Origin of X Chromosome Aberrations in Turner's Syndrome Jan Lindsten 1963

Exploring the Biological Contributions to Human Health Institute of Medicine 2001-07-02 It's obvious why only men develop prostate cancer and why only women get ovarian cancer. But it is not obvious why women are more likely to recover language ability after a stroke than men or why women are more apt to develop autoimmune diseases such as lupus. Sex differences in health throughout the lifespan have been documented. Exploring the Biological Contributions to Human Health begins to snap the pieces of the puzzle into place so that this knowledge can be used to improve health for both sexes. From behavior and cognition to metabolism and response to chemicals and infectious organisms, this book explores the health impact of sex (being male or female, according to reproductive organs and chromosomes) and gender (one's sense of self as male or female in society). Exploring the Biological Contributions to Human Health discusses basic biochemical differences in the cells of males and females and health variability between the sexes from conception throughout life. The book identifies key research needs and opportunities and addresses barriers to research. Exploring the Biological Contributions to Human Health will be important to health policy makers, basic, applied, and clinical researchers, educators, providers, and journalists-while being very accessible to interested lay readers.

Chromosome Abnormalities and Genetic Counseling R.J. MKinlay Gardner 2011-11-11 Advances in cytogenetics continue to crop up in wonderful ways, and we know exponentially more about chromosomes now than mere decades ago. Likewise, the necessary skills in offering genetic counseling continue to evolve. This new edition of Chromosome Abnormalities in Genetic Counseling offers a practical, up-to-date guide for the genetic counselor to marshal cytogenetic data and analysis clearly and effectively to families.

Management of Genetic Syndromes Suzanne B. Cassidy 2011-09-20 The bestselling guide to the medical management of common genetic syndromes—now fully revised and expanded A review in the American Journal of Medical Genetics heralded the first edition of Management of Genetic Syndromes as an "unparalleled collection of knowledge." Since publication of the first edition, improvements in the molecular diagnostic testing of genetic conditions have greatly facilitated the identification of affected individuals. This thorough revision of the critically acclaimed bestseller offers original insights into the medical management of sixty common genetic syndromes seen in children and adults, and incorporates new research findings and the latest advances in diagnosis and treatment of these disorders. Expanded to cover five new syndromes, this comprehensive new edition also features updates of chapters from the previous editions. Each chapter is written by an expert with extensive direct professional experience with that disorder and incorporates thoroughly updated material on new genetic findings, consensus diagnostic criteria, and management

strategies. Edited by two of the field's most highly esteemed experts, this landmark volume provides: A precise reference of the physical manifestations of common genetic syndromes, clearly written for professionals and families Extensive updates, particularly in sections on diagnostic criteria and diagnostic testing, pathogenesis, and management A tried-and-tested, user-friendly format, with each chapter including information on incidence, etiology and pathogenesis, diagnostic criteria and testing, and differential diagnosis Up-to-date and well-written summaries of the manifestations followed by comprehensive management guidelines, with specific advice on evaluation and treatment for each system affected, including references to original studies and reviews A list of family support organizations and resources for professionals and families Management of Genetic Syndromes, Third Edition is a premier source to guide family physicians, pediatricians, internists, medical geneticists, and genetic counselors in the clinical evaluation and treatment of syndromes. It is also the reference of choice for ancillary health professionals, educators, and families of affected individuals looking to understand appropriate guidelines for the management of these disorders. From a review of the first edition: "An unparalleled collection of knowledge . . . unique, offering a gold mine of information." —American Journal of Medical Genetics

Human Afflictions and Chromosomal Aberrations Raymond Turpin 2013-10-22 Modern Trends in Physiological Sciences, Volume 32: Human Afflictions and Chromosomal Aberrations presents the study of the links between chromosome aberrations and physical and mental congenital anomalies. This book discusses the possibilities of human cytogenetic research as well as its difficulties. Organized into 15 chapters, this volume begins with an overview of the development of human chromosome investigations. This text then explains the methods for studying human chromosomes, which can be applied without controlling the atmosphere of the incubator. Other chapters describe the structural features of the normal human karyotype. This book discusses as well the early appearance of a chromosome aberration that produces a change in the hereditary patrimony manifest in a constitutional disorder of the individual. The final chapter deals with the biochemical effects that correspond to numerical or structural anomalies in chromosome 21. This book is a valuable resource for genetecists, cytogeneticists, physicians, and clinical researchers.

ABNORMAL KARYOTYPES Sana Nimer Abu Shihab 2013-10 In my first book (Your Easy Way To Chromosomes), the main topic was about the human chromosomes, their structures, abnormalities, syndromes, and chromosome analysis. In this book I focused on abnormal karyotypes and how chromosomal abnormalities happen. A karyotype is a picture of a person's chromosomes from body cells (blood, hair, or any other tissue), photographing them through a microscope and arranging them in pairs, ordered by size and position of centromere for chromosomes of the same size. Karyotype test (alternative names are Chromosome Analysis, Chromosomal Analysis) plays a role in: diagnosis genetic diseases which are related to chromosomal abnormalities, diagnosis some birth defects, and provides clinical utility in the diagnosis and treatment of hematologic malignancies. On the other hand some genetic abnormalities cannot be detected by karyotype analysis such as microdeletions. Karyotype helps clinical cytogeneticist to identify abnormalities by: Counting the number of chromosomes and looking for extra chromosome such as in trisomy 21 or missing chromosome in a karyotype such as in Turner syndrome. Looking for changes in chromosome structure such as chromosomal deletions, duplications, translocations, insertions, inversions and other chromosomal abnormalities. Writing a book related to your field shows your passion and commitment to your job. Sana Nimer sananimer1@gmail.com sananimer1@hotmail.com

Understanding Genetics Genetic Alliance 2009 The purpose of this manual is to provide an educational genetics resource for individuals, families, and health professionals in the New York - Mid-Atlantic region and increase awareness of specialty care in genetics. The manual begins with a basic introduction to genetics concepts, followed by a description of the different types and applications of genetic tests. It also provides information about diagnosis of genetic disease, family history, newborn screening, and genetic counseling. Resources are included to assist in patient care, patient and professional education, and identification of specialty genetics services within the New York - Mid-Atlantic region. At the end of each section, a list of

references is provided for additional information. Appendices can be copied for reference and offered to patients. These take-home resources are critical to helping both providers and patients understand some of the basic concepts and applications of genetics and genomics.

Sex Determination Shanmugaratnam K. 1961

Genetic Disorders, Syndromology and Prenatal Diagnosis T.V.N. Persaud 2012-12-06 Birth defects have assumed an importance even greater now than in the past because infant mortality rates attributed to congenital anomalies have declined far less than those for other causes of death, such as infectious and nutritional diseases. As many as 50 % of all pregnancies terminate as miscarriages, and in the majority of cases this is the result of faulty intrauterine development. Major congenital malformations are present in at least 2 % of all liveborn infants, and 22 % of all stillbirths and infant deaths are associated with severe congenital anomalies. Not surprisingly, there has been a great proliferation of research into the problems of developmental abnormalities over the past few decades. This series, *Advances in the Study of Birth Defects*, was conceived in order to provide a comprehensive focal source of up-to-date information for physicians concerned with the health of the unborn child and for research workers in the fields of fetal medicine and birth defects. The first four volumes featured recent experimental work on selected areas of high priority and intensive investigation, including mechanisms of teratogenesis, teratological evaluation, molecular and cellular aspects of abnormal development, and neural and behavioural teratology. It seems logical and timely that the clinical aspects should now be presented. Accordingly, leading experts were invited to review a broad range of common problems from the standpoint of embryology, aetiology, clinical manifestations, diagnosis and management. This volume deals with genetic disorders and prenatal diagnosis.

Females Are Mosaics Barbara Migeon 2007-03-15 Women can be described as genetic mosaics because they have two distinctly different types of cells throughout their bodies. Unlike males, who have one X chromosome (inherited from their mother), females have two X chromosomes in every cell (one from each parent). The fathers copy works in some cells, while the mothers copy works in others. These two X chromosomes often function differently, especially if one carries a defective gene. Much has been written about the Y chromosome and its role in inducing maleness. This will be the first book about the X chromosome as a key to female development and the role of X-related factors in the etiology of sex differences in human disease. Barbara Migeon, from the renowned McKusick-Nathan Institute at Johns Hopkins, is a major figure in clinical genetics and is eminently qualified to write this book, and she writes clearly and effectively. She describes both the underlying molecular mechanisms and the remarkable genetic consequences of X inactivation and its role in determining the biological concepts characteristic of women. *Females are Mosaics* will be valuable to geneticists, biologists, and all health professionals interested in women's health.

Gender and Genes Klasien Horstman 2013 This Yearbook of Women's History (Jaarboek voor Vrouwengeschiedenis) is dedicated to Gender and Genes. Intruding upon our everyday lives, the world of DNA, genes and genomics has become a challenging field of research, both clinical and biomedical as well as socio-cultural. It is also a challenging topic for a Yearbook which traditionally focuses on women and gender from a historical point of view. Gender issues are part and parcel of genes and genomics in scientific research and socio-cultural discourses and representations. Current literature on genes and genomics does not abound in analyses of biomedical and socio-cultural realms where gender aspects are played out and exchanged. This Yearbook may thus contribute to a field of analysis which contextualizes history from the viewpoint of current biotechnological developments. This volume contains articles on medical cases (reproductive testing and the case of the sex chromosomes, and framing cancer risk in women and men), cultural representations, a portrait of female scientist Rosalind Franklin and interviews with feminist science philosophers Katarina Karkazis and Donna Dickenson.

MRCOG Part One Alison Fiander 2016-10-13 A fully updated and illustrated handbook providing comprehensive coverage of all curriculum areas

covered by the MRCOG Part 1 examination.

Human Chromosomes Sajirō Makino 1975

Human Chromosome Methodology Jorge J. Yunis 2012-12-02 Human Chromosome Methodology serves as an authoritative guide to cytogenetic techniques. This book presents each phase of laboratory work from preparation of materials for the X and Y bodies to application of other laboratory techniques including chromosome identification, autoradiography, and dermatoglyphics. The text also describes the structure and molecular organization of chromosomes and the advances in the automation of chromosome analysis. It provides a thorough review of the clinical manifestations of chromosome disorders. Organized into 13 chapters, the book presents the illustrated and diagrammatic examples and discussions of the subject matter and detailed tables and charts for learning efficiency. It also provides outlined presentation of cytogenetic procedures and notes and comments for each procedure that will assist readers in erroneous work phases. Moreover, it gives thorough lists of references in each chapter for further reading. This reference will be useful for research professionals, lecturers, genetics and molecular biology students, and members of the medical profession involved in genetics.

Turner Syndrome Mary Golanna 2022-12-01 Turner syndrome is a sex chromosomal condition in women that has an incidence rate of 1 in 2000 to 5000 live female births. (1) Girls and women who have it either have an absent X chromosome or there is an abnormality in one of the two X chromosomes. The exact cause for why the second X chromosome is missing or partially missing from cells is unknown. It is believed to result in a random cell division error while the reproductive cells of the parent are being formed. (2) Girls and women with Turner syndrome may be born with typical female genitalia or the genitalia can appear slightly abnormal. Some girls are diagnosed at birth because of these abnormalities, while others have no noticeable symptoms and are not diagnosed until later when they fail to begin puberty or menstruate. In most cases, the cause of Turner syndrome is unknown, although in some cases it does run in the family. In this quick start guide, you will discover... All there is to know about Turner syndrome Signs and symptoms of Turner syndrome Causes and types of Turner syndrome Ways to deal and treat this condition Information specifically for parents A 3-Step plan to help manage this condition

Gardner and Sutherland's Chromosome Abnormalities and Genetic Counseling R. J. M. Gardner 2018 Preceded by Chromosome abnormalities and genetic counseling / R.J. McKinlay Gardner, Grant R. Sutherland, Lisa G. Shaffer. c2012.

The X in Sex David. BAINBRIDGE 2009-06-30 A tiny scrap of genetic information determines our sex; it also consigns many of us to a life of disease, directs or disrupts the everyday working of our bodies, and forces women to live as genetic chimeras. The culprit--so necessary and yet the source of such upheaval--is the X chromosome, and this is its story. An enlightening and entertaining tour of the cultural and natural history of this intriguing member of the genome, *The X in Sex* traces the journey toward our current understanding of the nature of X. From its chance discovery in the nineteenth century to the promise and implications of ongoing research, David Bainbridge shows how the X evolved and where it and its counterpart Y are going, how it helps assign developing human babies their sex--and maybe even their sexuality--and how it affects our lives in infinitely complex and subtle ways. X offers cures for disease, challenges our cultural, ethical, and scientific assumptions about maleness and femaleness, and has even reshaped our views of human evolution and human nature. Table of Contents: Prologue 1. Making a Difference Interlude: What Is It, Exactly? 2. The Duke of Kent's Testicles Interlude: How Sexy Is X? 3. The Double Life of Women Epilogue: The Chosen One Further Reading Glossary Index Reviews of this book: The author of *Making Babies* takes a lively, witty tour of the X chromosome, creator of "a delicious symmetry between men and women"...Entertaining and informative...A fine demonstration of science made accessible. --Kirkus Reviews Reviews of this book: A well-written, well-researched, easy-to-read study that explains what has been learned about the X and Y chromosomes using DNA sequencing and other molecular

biology techniques. British biologist Bainbridge...has pulled together historical and current scientific research about how the X and Y chromosomes affect us and what the genes on these chromosomes actually do, like causing sex-linked diseases and color blindness...An excellent example of good science writing...Recommended. --Margaret Henderson, Library Journal Reviews of this book: Bainbridge is an essentialist, interested in understanding what aspects of gender are biologically driven, and why...He has a central question he wants to answer. The question is not so much why men and women are different (a worn topic that's the subject of too many Mars-and-Venus bestsellers) but, far more specific and far more interesting: Why are men and women more different than they need to be? --Liza Mundy, Washington Post Reviews of this book: Bainbridge summarizes our knowledge of the genetic information that determines one's sex by recounting the ancients' speculations about the genesis of gender, following with modern biologists' discovery of the X and Y chromosomes about a century ago, and of the sex-determining gene Sry in the 1990s. In a discussion rich with history, evolution, and philosophy, Bainbridge points out the dramatic effect that gender selection has on people's lives...A fascinating, often humorous analysis of the science of sexuality. --Gilbert Taylor, Booklist Reviews of this book: In *The X in Sex*, David Bainbridge explains the far-reaching effects of X. Bainbridge...moves with ease between straightforward accounts of biology and historical stories about its effect, like the chapter describing the progression of hemophilia through the royal houses of Europe. Bainbridge discusses cultural history as well as natural history, and his wit enlivens every page. --Christine Kenneally, New York Times Book Review Reviews of this book: There are many literary stars (such as Stephen Jay Gould, Richard Dawkins and Matt Ridley) in the firmament of writers on evolution, and to a man they write with dash and persuasive logic. David Bainbridge is one such and in his latest book he takes the reader through the glories of the X chromosome at a cracking pace. --Miriam Stoppard, Times Higher Education Supplement (UK) Reviews of this book: The truth is that the behaviours of [chromosomes] X and Y are inextricably linked. Bainbridge explores this link in a compelling tale that takes in how the sex chromosomes became sex chromosomes, and the very different consequences of this for women and men. Along the way we encounter the Duke of Kent's testicles, calico cats and non-identical identical twin girls. His story weaves science, history and the history of science (with a little religion for good measure) in a straightforward, anecdotal fashion that will appeal to scientists and non-scientists alike. --Mark T. Ross, New Scientist (UK) Reviews of this book: In his structure/function analysis of the X chromosome, Bainbridge provides a tongue-in-cheek, yet informative, description of one of the two human sex chromosomes. --R. Adler, Choice Reviews of this book: If you have ever been intrigued by some of the puzzles of genetics--why boys tend to get haemophilia or colour blindness while girls are more likely to have an identical twin or to develop rheumatoid arthritis later in life--then *The X in Sex* is for you. --Chris Tyler-Smith, Times Literary Supplement David Bainbridge takes us on a fascinating tour of X chromosomes and explains what the possession of these intricately folded, infinitesimally narrow, two-inch long strings of genetic codes weighing almost nothing, means for their bearers--that is for each one of us, male and female. History and personal anecdotes are woven together with up-to-date summaries of the science, punctuated with Bainbridge's zany--and very British--humor, so that this information-packed book is pure pleasure to read. --Sarah Blaffer Hrdy author of *Mother Nature: A History of Mothers, Infants, and Natural Selection* *The X in Sex* is absolutely fascinating, so intriguing, in fact, that I found myself unwilling to put it down. David Bainbridge surveys an astonishing amount of new information from recent genomic studies of the X chromosome, clearly explaining the findings in a way the average person can easily follow. The science is presented via amusing and highly appropriate metaphors and clever turns of phrase, all of which serve to brighten the prose and present the reader with catchy ways to think about complex ideas. This is an informative, authoritative, and thoroughly enjoyable read: one of the best books I have read in recent years. --Jane Lancaster, University of New Mexico This is wonderful stuff--beautifully written, clear, jargon-free, with anecdotes sure to hold the attention. --other hupauthor Tim Birkhead, author of *Promiscuity: An Evolutionary History of Sperm Competition*

Cassidy and Allanson's Management of Genetic Syndromes John C. Carey 2021-01-27 MANAGEMENT OF GENETIC SYNDROMES THE MOST RECENT UPDATE TO ONE OF THE MOST ESSENTIAL REFERENCES ON MEDICAL GENETICS Cassidy and Allanson's Management of Genetic Syndromes, Fourth Edition is the latest version of a classic text in medical genetics. With newly covered disorders and cutting-edge, up-to-date information, this resource remains the most crucial reference on the management of genetic syndromes in the field of medical genetics for students, clinicians, caregivers, and researchers. The fourth edition includes current information on the identification of genetic syndromes (including newly developed diagnostic criteria), the genetic basis (including diagnostic testing), and the routine care and management for more than 60 genetic disorders. Written by experts, each chapter includes sections on: Incidence Diagnostic criteria Etiology, pathogenesis and genetics Diagnostic testing Differential diagnosis Manifestations and Management (by system) The book focuses on genetic syndromes, primarily those involving developmental disabilities and congenital defects. The chapter sections dealing with Manifestations and Management represents the centerpiece of each entry and is unmatched by other genetic syndrome references. Management of Genetic Syndromes is perfect for medical geneticists, genetic counselors, primary care physicians and all healthcare professionals seeking to stay current on the routine care and management of individuals with genetic disorders.

Sex-determination Francis Albert Eley Crew 1965

Transition of Care M. Polak 2018-06-11 Pediatricians face many questions and challenges during patients' transition from childhood to adulthood. Chronic diseases justify long-term medical, psychological, educational, and social management. The establishment of networks with adult endocrinologists is, therefore, very important to achieving optimal conditions for a successful transition period. In this book, leading experts discuss major public health issues such as diabetes and obesity. They also review genetic diseases, including Turner syndrome and congenital adrenal hyperplasia. All clinical cases are presented in a focused way to highlight any issues which might be discussed by the doctor with patients and family. Psychological approaches are debated as are proposals for defining specific tools to encourage autonomy during this life period. This book is a valuable tool and will be helpful in answering many questions for endocrinologists, adult and pediatric, dedicated to the long-term management of patients with chronic diseases.

The Y Chromosome: Clinical aspects of Y chromosome abnormalities 1985

On the X Chromosome of Man Victor Almon McKusick 1964

Molecular Genetics of Sex Determination Stephen S. Wachtel 2014-07-23 In this era of accelerated discovery and prolific output, Molecular Genetics of Sex Determination keeps readers abreast of this field's fast-moving biology. Its chapters were completed by experts in each area only months before publication. The text is organized into two parts. First, it reviews the basic biology of sex determination and summarizes groundbreaking work in mouse, marsupial, and Drosophila systems. Second, it covers current human genetics, clinical studies, and the syndromes of abnormal sex differentiation. With chapters by preeminent reproductive biologists, this is a capital work. Ohno's law is described by Ohno; the Lyon hypothesis, by Lyon; Sinclair tells how he cloned the testis-determining gene; and so on. Molecular Genetics of Sex Determination is authoritative, comprehensive, and current. It is prime reading for geneticists, developmental biologists, graduate students in these and related fields, clinical researchers, physicians, and medical students. Reviews the genetics of sex determination in 19 up-to-date chapters Features research on sex chromosomes and sex-determining genes Includes abnormalities of sex determination and clinical genetics Written by scientists who pioneered work in this field

Sex Chromosomes Turner Syndrome

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