

Ahmad S. Teebi
Editor

Genetic Disorders Among Arab Populations

Second Edition

 Springer

Genetic Disorders Among Arab Populations

Sabine Zange



Genetic Disorders Among Arab Populations:

Genetic Disorders Among Arab Populations Ahmad S. Teebi, 2010-07-30 Arab populations have their own genetic disorders both universal and particular Genetic diversity within these source populations along with the fact that the rates of inbreeding are often high and family sizes are often large constitute conditions that facilitate the emergence and detection of phenotypes explained notably by autosomal recessive inheritance in which case the use of homozygosity gene mapping can facilitate the discovery of the corresponding genes The present book includes 5 parts dealing with various aspects that relate to the genetic structure of Arabs and minorities within the Arab world as well as genetic disorders prevalent in this part of the world It includes updated reviews of the genetic disorders in various Arab countries and geographic regions The focus is primarily but not exclusively on the group of single gene disorders with particular emphasis on autosomal recessive conditions It further includes epidemiological and clinical data as well as inheritance patterns mutation and polymorphism data and available haplotype analysis data The ethnic and genetic diversity of the Arab populations is discussed as well as aspects of genetic counseling practice in this region together with a proposal for an ethical framework for genetic research and prevention of genetic disorders The target audience of this book includes human and medical geneticists genetic counselors researchers medical specialists dealing with Arab patients or practicing in Arab countries medical and genetic counseling students and nurses

[The Gale Encyclopedia of Genetic Disorders](#) Stacey Blachford, 2002 Presents nearly four hundred articles describing genetic disorders conditions tests and treatments including high profile diseases such as Alzheimer s breast cancer and heart disease

Genetic Diseases of the Eye Elias I. Traboulsi, 1998 This authoritative reference book deals with the clinical aspects molecular genetics and management of both common and rare genetic disorders of the eye Topics include ocular embryology the role of developmental genes color vision and electrophysiologic testing in retinal disorders The management of the patient with low vision from inherited eye disease is also discussed Thoroughly referenced with almost 500 illustrations the book provides comprehensive up to date coverage of ophthalmic genetics

The Genetics of Neurological Disorders Michael Baraitser, 1997 BACK COVER COPY This third edition of *The Genetics of Neurological Disorders* incorporates the recent advances in genetics and molecular genetics into this comprehensive classic work The classification of diseases within the text has been updated in line with the change in practice necessitated by these recent advances Invaluable for both the geneticist and the neurologist this book provides up to date information about counselling recurrence risks across the wide range of conditions seen by both the adult and paediatric neurologist Each condition is covered in a way that helps the clinician reach a diagnosis before counselling and carrier detection and prenatal diagnosis are also discussed for many conditions Reviews of previous editions coherent well organised extremely well documented work This is a fine book to take to the clinic as well as the library *New England Journal of Medicine* No neurologist will care to function from now on without ready access to this book and it will be consulted by all

clinical geneticists *Annals of Human Genetics* The strength of this book lies in the scholarly treatment of both old and new literature pertaining to familial neurological diseases This book can heartily be recommended to the clinician seeking information on the genetics of neurological disease *Brain* As a comprehensive source of reference it is unrivalled a unique compilation of great importance which sets out all the current information about genetic elements markers and calculations of risk No neurologist can afford to be without a copy close to hand *Journal of Neurology Neurosurgery and Psychiatry* NOT on Back cover Additional reviews This edition has grown by 257 worthwhile pages *British Medical Journal* on 2e The book is very successful indeed combining a rigorous academic analysis of evidence for any genetic statement with clear English *Archives of Disease in Childhood* on 1e encyclopaedic coverage of the literature ranging from major studies of larger populations to single families exhibiting obscure conditions *Journal of Nervous and Mental Diseases* on 1e The book is a comprehensive compendium which presents the facts clearly *Developmental Medicine and Child Neurology* on 1e the best of its kind with wonderfully succinct summaries and replete with over 200pp of references outstanding value it will deservedly find itself very widely used and referred to with pleasure providing as it does so lucid and informative a guide to a complex and painful subject matter *Genetic Research* Cambridge on 2e *Population Structure and Genetic Disorders* Aldur W. Eriksson,1980 Research Report American University of Beirut,1995 Hereditary Hearing Loss and Its Syndromes Helga V. Toriello,William Reardon,Robert J. Gorlin,2004 This unique textbook aims to assist clinicians dealing with deaf patients and families by critically reviewing all relevant published material on genetics pathology clinical presentation diagnosis and laboratory findings Thoroughly revised the Second Edition has been updated throughout and includes a new chapter on hearing loss with cardiovascular disorders It reflects all the progress on the molecular understanding of deafness made in recent years and integrates these findings into clinical practice **Genetic Skin Disorders** Virginia P. Sybert,1997 A comprehensive survey of the subject this scholarly work describes in full detail 327 distinct inherited dermatologic conditions Each disease entry follows a consistent format with sections devoted to major and minor dermatologic features associated clinical abnormalities histopathology including both light and electron microscopy biochemical and molecular information treatment mode of inheritance and recurrence risks prenatal diagnosis differential diagnosis support group listing and annotated references **Annals of Saudi Medicine** ,2012 **The Genetics of Renal Disease** Frances Flinter,E. R. Maher,Anand Saggarr-Malik,2003 Renal genetic disorders may become apparent at any time in life and recognition of specific renal disorders is important in allowing accurate genetic counselling Active counselling requires knowledge about the natural history of the disorder an understanding of the genetic component and mode of inheritance risk of the disorder in other family members information about patient attitudes towards family planning and the possibility of pre natal diagnosis *The Genetics of Renal Disease* provides a comprehensive account of both the hereditary nephropathies and more generalised disorders which may affect the renal tract It includes comprehensive guidelines from an international

group of authors on renal function and development chromosomal disorders dysmorphic syndromes inherited predisposition to kidney cancer and gene therapy for renal cancer Specific chapters deal with individual renal tract disorders such as an extensive discussion of Alport s disease cystic renal diseases the Bardet Biedl and Alstrom syndromes Each section gives a brief description of the clinical and pathological features of a particular disorder followed by a review of the genetics of the disorder including incidence inheritance of familial occurrence genetic linkage and gene assignment and prenatal diagnosis and carrier detection Original studies are extensively quoted and are referenced at the end of each section There are a great many dysmorphic syndromes that involve the urinary tract These are incorporated within the book *A Genetic Analysis of Arodermatitis Enteropathica* Shari L. Griffen,1998

Overgrowth Syndromes Meyer Michael Cohen,Giovanni Neri,Rosanna Weksberg,2002 Overgrowth Syndromes presents a broad yet in depth discussion of children who are large at birth or experience excessive postnatal growth or some combination of increased weight length and head circumference Many of these syndromes are associated with an increased frequency of tumors The book is important because of the ever increasing number of newly identified overgrowth syndromes and the rapid progression of molecular knowledge of these conditions It covers Beckwith Wiedemann syndrome Simpson Golabi Behmel syndrome Sotos syndrome Proteus syndrome Bannayan Riley Ruvalcaba syndrome Klippel Trenaunay syndrome neurofibromatosis and fragile X syndrome among other topics Each chapter provides a historical perspective and deals with epidemiology etiology and molecular biology when known clinical and pathological features diagnostic criteria and differential diagnosis The book is encyclopedic in scope It will be of value to pediatricians medical geneticists oncologists hematologists surgeons pathologists radiologists dermatologists nephrologists and molecular biologists

The Lancet ,2006 *X-linked Mental Retardation* Roger E. Stevenson,Charles E. Schwartz,Richard J. Schroer,2000 This is a comprehensive and up to date survey of mental retardation caused by genes on the X chromosome Clinical and laboratory data on 130 syndromes are presented in a concise and consistent manner Each syndrome is defined and information is provided on somatic features growth and development neurological signs cognitive performance imaging and other laboratory findings and when possible the nature and localization of the responsible gene Craniofacial and other somatic findings are extensively illustrated A differential matrix accompanies each syndrome description to assist the reader in identifying other X linked syndromes with overlapping features As background the X chromosome and the emergence of the concept of X linked mental retardation are discussed X chromosome maps show the location and mapping limits of the responsible genes The authors have extensive experience in the clinical and laboratory delineation of X linked mental retardation They have described new syndromes regionally mapped disease loci on the X chromosome and in several cases isolated the genes responsible for X linked syndromes

The Child with Multiple Birth Defects Meyer Michael Cohen,1997 A medical genetics book which came into existence because of the lack of literature on multiple congenital anomaly syndromes this study incorporates an approach to the patient with multiple

anomalies It deals with the underpinnings of the field of syndromology Designed for students residents fellows and physicians with an interest in syndromology this book assumes some general understanding of the rudiments of both medicine and genetics The author emphasizes the conceptual aspects of syndromology rather than provide an encyclopaedic treatment on specific syndromes atopic covered extensively in other textbooks Since 1982 when the first edition of this monograph came out the field has advanced dramatically That is why Cohen has rewritten the entire book and added new chapters on genetics teratogens and facial dysmorphology Medical Principles and Practice ,2007 *Inborn Errors of Development* Charles J. Epstein,Robert P. Erickson,Anthony Joseph Wynshaw-Boris,2008 Honored by the Association of American Publishers as the Best Medical Book of 2004 Inborn Errors is the definitive work on genetically caused abnormalities of human development Despite the explosion in genetic advances the causes of two thirds of all birth defects remain unknown However we are on the brink of a revolution in this area and Inborn Errors is at the forefront It is the first book to connect the disease causing gene to its biochemical pathway and to the structural functional disorder Mutations of the gene the clinical picture genetic counseling and prognosis and any known treatments are discussed For medical geneticists genetic counselors pediatricians and developmental biologists the book is a unique and groundbreaking reference

Environmental Justice for Arab Americans? Gia Grier,Amy MacDonald,Cari Varner,Kimberly Wetzel,Lauren Zajac,2005 **Preservation of Human Progeny in the Age of Biotechnology** Mohd. Saifuddeen,2015 **Annual Review of Genomics and Human Genetics** ,2009

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