

# GENETIC TESTING IN NEONATES AND CHILDREN



# The Genetic Testing Of Children

**Carol Levine**



## **The Genetic Testing Of Children:**

*The Genetic Testing of Children* Angus Clarke, 2020-07-26 This book written by a leading geneticist examines the ethical and social issues raised by the genetic testing of children The opinions of geneticists ethicists and affected families are all included to give a balanced view of this controversial field Issues covered include confidentiality potential abuses of genetic information eg the use of test results by insurance companies and the value of predictive genetic testing The aim of the book is to improve awareness of the complexity of the issues raised and provide suggestions as to how the discussions must develop it therefore raises new questions as well as answering those that already exist The Genetic Testing of Children

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University Institute. Law Department, 2009 Genetics, Society and Clinical Practice Peter S. Harper, Angus Clarke, 1997 This book examines from both ethical and clinical perspectives the impact of increasing genetic knowledge on individuals and families The authors use their experience of working at the coal face of clinical genetics to describe the social implications arising from the ability to test for an increasing number of genetic diseases Current controversial issues such as genetic testing and insurance early diagnosis of late onset disorders population screening and abuses of genetics are all described with great clarity Several of the chapters are based on lead articles and editorials the authors have been asked to write for The Lancet and British Medical Journal The book provides a concise introduction to the effects on individuals of the practical issues associated with genetic testing and therefore should be essential reading for all involved in clinical genetics

*Children, Medicine and the Law* Michael D. A. Freeman, 2005 This volume collects in convenient form the most influential articles analysing the key issues surrounding children medicine and the law today Issues examined include the implications of assisted reproduction for children neonatal intensive care health care HIV testing of new born children **Genetic Disorders Sourcebook** Kathy Massimini, 2000 Presents consumer health information gathered from sources including government reports medical publications and patient organizations to help readers understand the basic causes treatments and health care currently available for many kinds of genetic disorders Includes information on screening tests lifestyle expectations and optimal disease management through medications diets and physical therapy Concludes with a directory of resources advice about caring for children with chronic illness and a glossary of genetic terminology Annotation copyrighted

by Book News Inc Portland OR      *Genetics* Lori B. Andrews, Maxwell J. Mehlman, Mark A. Rothstein, 2006 This is the revised edition of the casebook *Genetics Ethics Law and Policy* which has been used successfully in law schools in both the seminar and course context It is authored by three of the nation's leading experts on genetic ethics law and policy Students enjoy the course because of the topicality of the subjects many of which they hear about in the news gene discoveries embryo stem cell research Faculty members enjoy teaching from the book because of the excellent teaching manual and because they can link it to other topics the casebook covers issues in health law employment law insurance law criminal law family law and other fields The casebook is supplemented regularly on the TWEN website so that it is always current A background in genetics is not required for either students or teachers The casebook and teachers manual are written so that the casebook can be used for undergraduate courses or courses for the health professions for public health or for public policy      **The Acts & Proceedings of the ... Regular Session of the General Synod, Reformed Church in America** Reformed Church in America. General Synod, 2001 Issues for 1868 include index      **Meeting Children's Psychosocial Needs Across the Health-care Continuum** Judy Rollins, Judy Holt Rollins, Rosemary Bolig, Carmel C. Mahan, 2005      Page's Ohio Revised Code Annotated Ohio, 1992      Handbook of Genomics and the Family Kenneth P. Tercyak, 2010 Today's consumer can send a company a DNA sample and receive a detailed set of genetic test results in return all without ever visiting a health care provider Although knowing one's personal risk for serious disease may lead some individuals to make more informed health choices an ever growing set of questions remains Are predictive genetic tests meaningful Can the results be harmful as well as helpful In what ways can genetic information be used by health care providers to predict disease risk and optimize medical management within concerned families Most important how might the landscape of genetic testing affect the care of children's health Although there are no easy answers the *Handbook of Genomics and the Family* details in one authoritative volume the challenges entailed by the latest genetic advances and offers insights into the potential translation of this knowledge in pediatric and family practice and public health This timely and comprehensive reference introduces readers to the study of how genes singly and in combination with each other and the environment affect health and behavior summarizes current findings on genetics and genomics in disease etiology and prevention and provides family focused perspectives relating to genetic counseling and education Key features of the *Handbook* include In depth background in genetics and genomics in relationship to disease etiology and epidemiology Integrative frameworks for understanding genetics and genomics in the context of families Considerations for communicating genetic and genomic information Individual chapters on genetic and familial risks for specific diseases and conditions including cancer obesity and Type 1 diabetes among children and adolescents Training ethics and other emerging issues and controversies Contemporary policies on genetic testing in children and families The *Handbook of Genomics and the Family* is essential reading for clinical child and pediatric psychologists and psychiatrists pediatricians and nurses clinical geneticists and genetic counselors and public health

specialists as advances in genetics lead the way to more personalized models of health *The Lancet*, 1938 **Southern Educational Journal**, 1898 **Genetics in the Clinic** Mary Briody Mahowald, 2001 *Genetics in the Clinic* provides a broad understanding of advances in genetics and their implications for primary care practice. The book begins with basic concepts in genetics diagnosis and management of genetic disease cancer genetics and prenatal testing. Part Two deals with topics relevant to all areas of primary care such as cultural and ethnic differences the role of the nurse and genetic counselling. Part Three addresses ethical or social issues applicable to specific practices. The book's final section considers regulatory matters including the impact of managed care and insurance and employment issues. Practitioners who are knowledgeable about health care but who have no special training in genetics will find this book especially useful as they learn to deal with the clinical ethical and social implications of advances in genetics for their practice. Thoroughly explains how to obtain a useful family history what kind of genetic tests to order and when how to counsel parents about testing of children how to counsel for late onset or susceptibility conditions and how to identify those for whom cancer risk or other risk testing is appropriate. Expert perspectives on disability ethnic gender and class differences reflecting the wide variety of patients seen by today's practitioners. Glossary of genetic terms and extensive bibliography of print and electronic sources useful for clinicians who want to update their practice or for explaining complex terminology to patients. Dr McKusick is world renowned for his expertise in the field of human genetics and specific genetic conditions. Contributors have been trained in various areas of primary care obstetrics paediatrics family medicine internal medicine and nursing **Genes, Society and the Future, Volume II, Part Five - Genetic Testing of Competent Minors** Deborah Lawson, 2014 Genetic testing raises new issues from those involved in other medical contexts particularly for children. Most of the concerns relevant to minors are prompted by the familial and predictive aspects of genetic information. It is vital that GPs and other health professionals know more about genetic testing and genetics services in New Zealand so that they can better facilitate informed consent recognise and acknowledge any limitations in their expertise particularly as they will influence their patients when discussing testing possibilities know when to refer patients for genetic testing and can offer some degree of genetic counselling if required. Genetic testing of children who lack capacity to consent to genetic testing for non medical reasons should be treated with caution. Many adults choose not to discover their own genetic risk status and the threat to the child's autonomy and right to confidentiality are the reasons for this caution. Also where there is a lack of evidence about what the test results may signify for the child's health this uncertainty is best dealt with by waiting until the child is able to make personal choices. A register should be established to facilitate disclosure to persons who have reached the age of sixteen or eighteen years or earlier if they are competent and personally seek access to the information of the fact that they underwent genetic testing as children. Initially the minors may be informed either that they underwent predictive or carrier testing as children or that some information is available about genetic risk status should they wish to access it. Such a register is the appropriate method for

ensuring that people who undergo testing as children are informed of the fact for the following reasons Firstly it would encourage parents and health professionals to disclose test results to children as the fact of testing will be disclosed to them anyway Secondly it gives the person tested a choice regarding whether or not to access the information assuming that he or she has not already been told Thirdly it avoids the difficulties of imposing a new disclosure duty that may have unwieldy and undesirable consequences in terms of monitoring enforcement and sanctions Genetic counselling would be required to assist minors in deciding whether to access their test results and to support them whatever their choice The privacy of the register and its information must be strictly maintained      **DNA-based Tests** Jerroo S. Kotval,1994      **The New Child Study** ,1926

*Taking Sides* Carol Levine,1997 *Taking Sides Clashing Views on Controversial Bioethical Issues* 10th edition is a debate style reader designed to introduce students to controversies in bioethics The readings which represent the arguments of leading philosophers and social commentators reflect a variety of viewpoints and have been selected for their liveliness and substance and because of their value in a debate framework For each issue the editor provides a concise introduction and postscript summary The introduction sets the stage for the debate as it is argued in the yes and no readings and the postscript briefly reviews the opposing views and suggests additional readings on the controversial issue under discussion By requiring students to analyze opposing viewpoints and reach considered judgments *Taking Sides* actively develops students critical thinking skills It is this development of critical thinking skills that is the ultimate purpose of each of the volumes in the widely acclaimed *Taking Sides* program Book jacket      Truth and Consequences of the Genetic Revolution California. Legislature. Senate. Select Committee on Genetics and Public Policy,1996

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