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Core Curriculum for Specializing in Intellectual and Developmental Disability Wendy M. Nehring,2005 The first interdisciplinary text on

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Lewis, **Encyclopedia of Human Genetics and Disease** Evelyn B. Kelly, 2013-01-07 This two volume encyclopedia examines the history characteristics causes and treatment of genetic disease as well as the science of genetics itself Modern science has unlocked many of the mysteries of genetics providing a blueprint for understanding the origins behind previously mysterious ailments and conditions both common and uncommon A complete understanding remains elusive however geneticists are still refining theories about what causes chromosomes to mutate and genetic diseases remain difficult to diagnose and challenging to treat This fascinating reference explores the scientific and human aspects of this complex field of science Encyclopedia of Human Genetics and Disease features nearly 400 entries including well known genetic diseases rare and lesser known genetic diseases and the genetic factors that may contribute to common diseases and health conditions such as breast cancer and obesity The author presents in depth discussions of concepts essential to understanding genetic disease in 18 entries that provide background on key topics such as Genetics 101 the genome and the foundations of genetics genetic counseling and newborn screening Each of the 355 disorders profiled provides the history of the condition its prevalence causes treatment if any and further reading Interesting sidebars and compelling photos that help inform content accompany many entries *Vogel and Motulsky's Human Genetics* Friedrich Vogel, Arno G. Motulsky, 2013-04-17

The first two editions of this book published in 1979 and in 1986 were well received by the scientific community Translations into Italian Japanese and Russian suggest that this book was regarded useful in many parts of the world Meanwhile human genetics has seen dramatic developments and the molecular revolution has attracted thousands of scientists including many molecular biologists to this field About 3700 human genes have already been mapped to chromosomal sites Many such genes have been cloned and the various mutations causing disease have been identified Novel mutational mechanisms such as expanded trinucleotide repeats have been discovered in conditions such as Huntington's disease and the fragile X syndrome of mental retardation Gene action now can often be elucidated by studying the pathway from gene to phenotype following positional cloning rather than working in the opposite direction as was customarily done before the tools of new genetics were available In an increasing number of genetic diseases the pathogenic mechanisms have been elucidated with

positive consequences for prevention and treatment. It therefore became necessary to rewrite almost completely major portions of this book. These developments are now making genetics arguably the leading basic science for medicine as well as a recognized medical speciality. But all these changes do not mean that the entire framework of human genetics had to be reconstructed.

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Clinical Genetics in Nursing Practice Felissa R. Lashley, 2005-04-15 Designated a Doody's Core Title. The third edition of this award winning text provides new and updated knowledge about genetics issues relevant to nursing practice. Read in sequence or used as a reference, this is a comprehensive overview of how genetics affects the care that nurses provide. In addition to a summary of basic human genetics and discussion of the Human Genome Project, this new edition includes the latest research findings and implications about inheritance, major genetic disorders, cytogenetics or chromosomal inherited biochemical and congenital anomalies and genetics in twin studies. A consideration of the ethical impact of genetics on society and future generations as well as information on assisted reproduction round out the overview. Includes over 100 illustrations and photos of specific genetic disorders, tables and figures on the distribution of disease and an extensive appendix listing associations, organizations and websites relevant to genetics.

Human Genetics Case Workbook Ricki Lewis, 2004-02-16 Researching the Autism Spectrum Ilona Roth, Payam Rezaie, 2011-01-06 This selection of contemporary research provides up to date perspectives from leading investigators who are at the cutting edge of studies in autism spectrum disorders. The book allows readers to grasp new approaches to understanding the autism spectrum. Key areas of theory and research are covered from classification and diagnosis, genetics, neurology and biochemistry to socio-cognitive, developmental and educational perspectives essential to a broader understanding of the autism spectrum. In addition, it introduces new emphases on MEG

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