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Introduction To Genetics Study Guide

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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.

Clear, concise, and well-organized, the Cell and Molecular Biology Study Guide is an excellent learning tool for students of cellular and molecular biology. The sixteen chapters of the book follow a logical progression beginning with an introduction to cells and concluding with an overview of current techniques in cellular and molecular biology. Each brief chapter effectively separates core concepts, clarifying each individually and creating a set of building blocks that allow students to fully comprehend one aspect of the subject matter before moving on to the next. Topics in the guide include: Bioenergetics, Enzymes, and Metabolism The Plasma Membrane The Cytoskeleton and Cell Motility DNA Replication and Repair Cell Signaling and Signal Transduction The book also covers aerobic respiration and mitochondria, photosynthesis, and the chloroplast, the nature of the gene and genome, gene expression, and cellular reproduction. Accessible and informative, Cell and Molecular Biology Study Guide can be used as a companion to standard textbooks in the field. It is also a useful reference tool for students new to the discipline or those looking for a quick review of the subject matter. Mark Running earned his Ph.D. in genetics at the California Institute of Technology and completed postdoctoral research at the University of California, Berkeley. Dr. Running is an assistant professor in the Department of Biology at the University of Louisville in Kentucky where he teaches courses in developmental, cellular, and molecular biology. In addition to his teaching, he serves on the Undergraduate Curriculum Committee. Dr. Running is the recipient of numerous grants from the National Science Foundation, and was a Howard Hughes Predoctoral Fellow and a Damon Runyon-Walter Winchell Cancer Research Postdoctoral Fellow.

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It has been recognized for almost 200 years that certain families seem to inherit cancer. It is only in the past decade, however, that molecular genetics and epidemiology have combined to define the role of inheritance in cancer more clearly, and to identify some of the genes involved. The causative genes can be tracked through cancer-prone families via genetic linkage and positional cloning. Several of the genes discovered have subsequently been proved to play critical roles in normal growth and development. There are also implications for the families themselves in terms of genetic testing with its attendant dilemmas, if it is not clear that useful action will result. The chapters in The Genetics of Cancer illustrate what has already been achieved and take a critical look at the future directions of this research and its potential clinical applications.

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