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KEY=EDITION - JONATHAN WEBER

HUMAN GENETICS

McGraw-Hill Education Today, human genetics is for everyone. It is about variation more than about illnesses, and increasingly about the common rather than about the rare. Once an obscure science or an occasional explanation for an odd collection of symptoms, human genetics is now part of everyday conversation. By coming to know genetic backgrounds, people can control their environments in more healthy ways. Genetic knowledge is, therefore, both informative and empowering. The 10th edition of Human Genetics: Concepts and Applications shows students how and why that is true. Users who purchase Connect Plus receive access to the full online ebook version of the textbook.

HUMAN HEREDITY: PRINCIPLES AND ISSUES

Cengage Learning HUMAN HEREDITY presents the concepts of human genetics in clear, concise language and provides relevant examples that you can apply to yourself, your family, and your work environment. Author Michael Cummings explains the origin, nature, and amount of genetic diversity present in the human population and how that diversity has been shaped by natural selection. The artwork and accompanying media visually support the material by teaching rather than merely illustrating the ideas under discussion. Examining the social, cultural, and ethical implications associated with the use of genetic technology, Cummings prepares you to become a well-informed consumer of genetic-based health care services or provider of health care services. Available with InfoTrac Student Collections <http://gocengage.com/infotrac>. Important Notice: Media content referenced within the product description or the product text may not be available in the ebook version.

THE FOREVER FIX

GENE THERAPY AND THE BOY WHO SAVED IT

St. Martin's Press Fascinating narrative science that explores the next frontier in medicine and genetics through the very personal prism of the children and families gene therapy has touched. Eight-year-old Corey Haas was nearly blind from a hereditary disorder when his sight was restored through a delicate procedure that made medical history. Like something from a science fiction novel, doctors carefully injected viruses bearing healing genes into the DNA of Corey's eyes—a few days later, Corey could see, his sight restored by gene therapy. THE FOREVER FIX is the first book to tell the fascinating story of gene therapy: how it works, the science behind it, how patients (mostly children) have been helped and harmed, and how scientists learned from each trial to get one step closer to its immense promise, the promise of a "forever fix," - a cure that, by fixing problems at their genetic root, does not need further surgery or medication. Told through the voices of the children and families who have been the inspiration, experimental subjects, and successes of genetic science, THE FOREVER FIX is compelling and engaging narrative science that tells explores the future of medicine as well as the families and scientists who are breaking new ground every day.

LEWIS, HUMAN GENETICS, NASTA EDITION

McGraw-Hill Education Today, human genetics is for everyone! It is about variation more than about illnesses, and increasingly about the common rather than the rare. Once an obscure science or an occasional explanation for an odd collection of symptoms, human genetics is now part of everyday conversation. By coming to know genetic backgrounds, people can control their environments in more healthful ways. Genetic knowledge is, therefore, both informative and empowering, and this new tenth edition of Human Genetics: Concepts and Applications shows students how and why that is true. New to the tenth edition, compelling stories and case studies include: A university encourages genetic testing of all incoming freshmen until stopped How are autism and schizophrenia related? How famine sowed the seeds of schizophrenia, thanks to epigenetics The Berlin patient: curing leukemia and HIV A new view of allergy, due to mutation Experimental evolution and the Gulf oil spill Human Genetics is now available with McGraw-Hill Connect Human Genetics, a web-based assignment and assessment platform that gives students the means to better connect with their coursework, with their teachers, and with the important concepts they will need to know for success now and in the future. Connect Human Genetics allows students to practice important skills at their own pace and on their own schedule. Students' assessment results and teacher's feedback are all saved online, so students can continually review their progress and plot their course to success. Teachers can also choose ConnectPlus Human Genetics. Like Connect, ConnectPlus provides online assignments and assessments, plus 24/7 access to an eBook - an online edition of the text - to aid them in successfully completing their work, wherever and whenever they choose.

HUMAN GENETICS

THE MOLECULAR REVOLUTION

Jones & Bartlett Learning Begins with molecular characterization of the human genome (rather than the conventional descriptions of Mendelian inheritance, pedigree analysis, and chromosome abnormalities), and maintains this emphasis on understanding human genetics in molecular terms throughout. Suitable as a text for biology

ESSENTIALS OF GENETICS, EBOOK, GLOBAL EDITION

Pearson Higher Ed For all introductory genetics courses A forward-looking exploration of essential genetics topics Known for its focus on conceptual understanding, problem solving, and practical applications, this bestseller strengthens problem-solving skills and explores the essential genetics topics that today's students need to understand. The 9th Edition maintains the text's brief, less-detailed coverage of core concepts and has been extensively updated with relevant, cutting-edge coverage of emerging topics in genetics. The full text downloaded to your computer With eBooks you can: search for key concepts, words and phrases make highlights and notes as you study share your notes with friends eBooks are downloaded to your computer and accessible either offline through the Bookshelf (available as a free download), available online and also via the iPad and Android apps. Upon purchase, you will receive via email the code and instructions on how to access this product. Time limit The eBooks products do not have an expiry date. You will continue to access your digital ebook products whilst you have your Bookshelf installed.

HUMAN GENETICS

CONCEPTS AND APPLICATIONS

McGraw-Hill Companies This work helps to make human genetics come alive through real-life examples and stories. It is designed for the non-science major taking an introductory level course.

ENCYCLOPEDIA OF HUMAN GENETICS AND DISEASE

ABC-CLIO This two-volume encyclopedia examines the history, characteristics, causes, and treatment of genetic disease, as well as the science of genetics itself. * Illustrations and photographs * An informative timeline of genetic study advancements * An annotated list of websites and helpful books

10TH INTERNATIONAL CONGRESS OF HUMAN GENETICS

ESSENTIALS OF GENETICS, GLOBAL EDITION.

CONCEPTS OF GENETICS

Benjamin-Cummings Publishing Company Concepts of Genetics is known for its focus on teaching core concepts and problem solving. This best-selling text has been extensively updated, with coverage on emerging topics in genetics, and problem-solving support has been enhanced.

BIOLOGY

McGraw-Hill Education Committed to Excellence in the Landmark Tenth Edition. This edition continues the evolution of Raven & Johnson's Biology. The author team is committed to continually improving the text, keeping the student and learning foremost. We have integrated new pedagogical features to expand the students' learning process and enhance their experience in the ebook. This latest edition of the text maintains the clear, accessible, and engaging writing style of past editions with the solid framework of pedagogy that highlights an emphasis on evolution and scientific inquiry that have made this a leading textbook for students majoring in biology and have been enhanced in this landmark Tenth edition. This emphasis on the organizing power of evolution is combined with an integration of the importance of cellular, molecular biology and genomics to offer our readers a text that is student friendly and current. Our author team is committed to producing the best possible text for both student and faculty. The lead author, Kenneth Mason, University of Iowa, has taught majors biology at three different major public universities for more than fifteen years. Jonathan Losos, Harvard University, is at the cutting edge of evolutionary biology research, and Susan Singer, Carleton College,, has been involved in science education policy issues on a national level. All three authors bring varied instructional and content expertise to the tenth edition of Biology.

CONCEPTS OF GENETICS: PEARSON NEW INTERNATIONAL EDITION

Pearson Higher Ed Concepts of Genetics is known for its focus on teaching core concepts and problem solving. This best-selling text has been extensively updated, with coverage on emerging topics in genetics, and problem-solving support has been enhanced.

ETHICS AND HUMAN GENETICS

A CROSS-CULTURAL PERSPECTIVE

Springer Science & Business Media Based in part on a survey of ethical decision-making among 682 medical geneticists worldwide, this book includes a chapter authored by a geneticist and an ethicist in 19 nations, describing genetic services, counselling, screening, prenatal diagnosis, and major ethical problems and social controversies faced by geneticists. The concluding chapter describes ethical and policy issues that exist worldwide, and offers some possible resolutions.

UNDERSTANDING GENETICS

A NEW YORK, MID-ATLANTIC GUIDE FOR PATIENTS AND HEALTH PROFESSIONALS

Lulu.com 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.

BEFORE WE ARE BORN, 10TH EDITION-SOUTH ASIA EDITION EBOOK

Elsevier India Covering the essentials of normal and abnormal human development for students in a variety of health science disciplines, Before We Are Born: Essentials of Embryology and Birth Defects, 10th Edition-South Asia Edition, reflects new research findings and current clinical practice through concise text and abundant illustrations. This edition has been fully updated by the world's foremost embryologists and is based on the popular text, The Developing Human, written by the same author team. It provides an easily accessible understanding of all of the latest advances in embryology, including normal and abnormal embryogenesis, causes of birth defects, and the role of genes in human development. Features streamlined content throughout, numerous photographs of common clinical cases and embryological explanations, didactic illustrations, and nearly 700 USMLE-style questions with full answers and explanations to help prepare for professional exams. Includes interactive clinical cases in every chapter that make important connections between human development and clinical practice--ideal for preparing for USMLE Step 1. Features access to 18 full-color, expertly narrated animations that guide students through key concepts in learning the complexities of embryologic development. Includes many new color photographs, new diagnostic images (3D ultrasound, CT scans, and MR images), an updated teratology section, revised and highlighted information on molecular aspects of developmental biology, and new information on the cellular and molecular basis of embryonic development. Follows the official international list of embryological terms (Terminologia Embryonica, 2013).

MEDICAL GENETICS AT A GLANCE

John Wiley & Sons Medical Genetics at a Glance covers the core scientific principles necessary for an understanding of medical genetics and its clinical applications, while also considering the social implications of genetic disorders. This third edition has been fully updated to include the latest developments in the field, covering the most common genetic anomalies, their diagnosis and management, in clear, concise and revision-friendly sections to complement any health science course. Medical Genetics at a Glance now has a completely revised structure, to make its content even more accessible. Other features include: • Three new chapters on Gene Identification, The Biology of Cancer, and Genomic Approaches to Cancer • A much extended treatment of Biochemical Genetics • A completely revised chapter on The Cell Cycle, explaining principles of biochemistry and genetics which are fundamental to understanding cancer causation • Two new chapters on Cardiac Developmental Pathology • An extended Case Studies section Providing a broad understanding of one of the most rapidly progressing topics in medicine, Medical Genetics at a Glance is perfect for students of medicine, molecular biology, genetics and genetic counselling, and is a previous winner of a BMA Award.

THE DEVELOPING HUMAN

CLINICALLY ORIENTED EMBRYOLOGY

Saunders Written by some of the world's most famous anatomists, the 10th edition of The Developing Human: Clinically Oriented Embryology continues to present medical students with a comprehensive and easily digestible review of this complex topic. Clearly written and well-structured descriptions include just the information that's needed, while nearly 600 illustrations help provide a clinically oriented guide to human development, utilizing a week-by-week and stage-by-stage approach to describe fetal organ and system development. Review questions and answers at the end of each chapter allow for effective exam preparation. Covers the latest advances in embryology, including normal and abnormal embryogenesis, causes of birth defects, and the role of genes in human development. Details how discoveries in molecular biology have affected clinical practice, including the development of sophisticated new techniques such as recumbent DNA technology and stem cell manipulation. Clinical case presentations, highlighted in special boxes, demonstrate how embryology concepts relate to clinical practice and are ideal for preparing for the USMLE Step 1. Three-dimensional animations - 2 new to this edition - help visual learners understand the subjects as discussed in the book as a whole. New and thoroughly revised assessment questions by Mark Torchia. Presents an authoritative description of human embryology through all stages of development. Rich illustrations correspond to the

text to enhance comprehension. Student Consult eBook version included with purchase. This enhanced eBook experience allows you to search all of the text, figures, references, and videos from the book on a variety of devices.

HUMAN HEREDITY: PRINCIPLES AND ISSUES

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MERCER'S TEXTBOOK OF ORTHOPAEDICS AND TRAUMA TENTH EDITION

CRC Press Highly Commended, BMA Medical Book Awards 2013 Orthopaedic problems account for over one-third of all medical and surgical problems. Mercer's Textbook of Orthopaedics and Trauma provides all the information required by the senior trainee or qualified specialist to improve understanding and management of any given condition or disease in this area. Si

HUMAN HEREDITY: PRINCIPLES AND ISSUES

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10TH MEETING OF THE IRISH SOCIETY OF HUMAN GENETICS, 2007

MASTERING PHARMACOGENOMICS: A NURSE'S HANDBOOK FOR SUCCESS

Sigma Theta Tau Pharmacogenomics, the study of the role of genetics in drug response, has developed into a critical new area in battling disease and improving outcomes. As our knowledge of the human genome grows, and with the advent of next-generation sequencing technologies, the role of genetic research is evolving to guide therapy and prevent many diseases and mitigate side-effects of current treatments. As a pivotal member of the healthcare team, nurses must acquire competency in pharmacogenomics to deliver optimal patient care in the 21st century. Mastering Pharmacogenomics provides nursing professionals with a foundational knowledge of human genetics and genomics that can be applied in optimizing drug therapy and patient care delivery. Coverage includes: Essential pharmacogenomics competencies for registered nurses Genomics technologies and resources Critical properties of drug response and disposition Covered specialties include obstetrics and prenatal patients, pediatrics, older adult patients, and oncology Ethics and patient care Future developments in technology and delivery

INTRODUCING GENETICS

FROM MENDEL TO MOLECULES

Garland Science The new edition of *Introducing Genetics* is a clear, concise, and accessible guide to inheritance and variation in individuals and populations. It first establishes the principles of Mendelian inheritance and the nature of chromosomes, before tackling quantitative and population genetics. The final three chapters introduce the molecular mechanisms t

PRESENT KNOWLEDGE IN NUTRITION

John Wiley & Sons *Present Knowledge in Nutrition, 10th Edition* provides comprehensive coverage of all aspects of human nutrition, including micronutrients, systems biology, immunity, public health, international nutrition, and diet and disease prevention. This definitive reference captures the current state of this vital and dynamic science from an international perspective, featuring nearly 140 expert authors from 14 countries around the world. Now condensed to a single volume, this 10th edition contains new chapters on topics such as epigenetics, metabolomics, and sports nutrition. The remaining chapters have been thoroughly updated to reflect recent developments. Suggested reading lists are now provided for readers wishing to delve further into specific subject areas. An accompanying website provides book owners with access to an image bank of tables and figures as well as any updates the authors may post to their chapters between editions. Now available in both print and electronic formats, the 10th edition will serve as a valuable reference for researchers, health professionals, and policy experts as well as educators and advanced nutrition students.

TEXTBOOK OF HUMAN GENETICS

Oxford University Press, USA This established text has been completely revised, reorganized, and updated to accommodate the vast changes that have occurred in human genetics during the last decade. Using numerous examples from human biology and medicine, the author reviews the basic laws of genetics, the common genetic diseases, and their causes in humans. He presents the latest data and techniques, including the molecular regulation of gene expression in humans and the use of restriction fragment length polymorphisms (RFLPs) to map human genes and identify genetic diseases. New chapters have been added on human immunogenetics, oncogenes, and cancer. The author explains complex concepts with clarity and uses an extensive array of diagrams and illustrations, including electron micrographs and clinical photographs of all common genetic diseases.

CELLULAR AND MOLECULAR IMMUNOLOGY E-BOOK

Elsevier Health Sciences *Cellular and Molecular Immunology* takes a comprehensive yet straightforward approach to the latest developments in this active and fast-changing field. Drs. Abul K. Abbas, Andrew H. Lichtman, and Shiv Pillai present sweeping updates in this new edition to cover antigen receptors and signal transduction in immune cells, mucosal and skin immunity, cytokines, leukocyte-endothelial interaction, and more. This reference is the up-to-date and readable textbook you need to master the complex subject of immunology. Recognize the clinical relevance of the immunology through discussions of the implications of immunologic science for the management of human disease. Grasp the details of experimental observations that form the basis for the science of immunology at the molecular, cellular, and whole-organism levels and draw the appropriate conclusions. Stay abreast of the latest advances in immunology and molecular biology through extensive updates that cover cytokines, innate immunity, leukocyte-endothelial interactions, signaling, costimulation, and more. Visualize immunologic processes more effectively through a completely revised art program with redrawn figures, a brighter color palette, and more 3-dimensional art. Find information more quickly and easily through a reorganized chapter structure and a more logical flow of material.

MATERNITY NURSING CARE (BOOK ONLY)

Cengage Learning MATERNITY NURSING CARE, 2nd Edition delivers all of the core components of obstetrical nursing, along with in-depth study features to help you develop and enhance analytical skills. Starting with women's health issues and pregnancy care, the chapters progress through labor and childbirth, postpartum care, and newborn development, addressing the nursing considerations for both normal and high-risk scenarios where appropriate. Written from a nursing perspective, MATERNITY NURSING CARE, 2ND Edition emphasizes overall patient care as opposed to medical treatment alone, and illustrates the critical importance of fostering the nurse-patient relationship. MATERNITY NURSING CARE, 2ND Edition is also bound with a helpful StudyWARE™ online, which includes case studies, NCLEX-style review questions, videos, activities, a glossary, and many other features to help you meet and exceed the highest standards of obstetrical nursing care. Important Notice: Media content referenced within the product description or the product text may not be available in the ebook version.

MEDELIAN INHERITANCE IN MAN

A CATALOG OF HUMAN GENES AND GENETIC DISORDERS

The twelfth edition of this classic reference work includes: • More than 2,000 new entries • A total of more than 9,000 entries • New features and enhancement of the familiar old features • Mapping information on more than 4,000 genes of known function • Information on specific point mutations responsible for more than 700 genetic disorders or neoplasms Mendelian Inheritance in Man (MIM) is a genetic knowledgebase that serves clinical medicine and biomedical research, including the Human Genome Project. It aims to be comprehensive (not only complete, but also collated, integrated, and interpreted), authoritative (not only accurate but also sound in its interpretations and judgements), and timely (not only up-to-date but also historically dimensioned). From a review of the eleventh edition, *Reproductive Toxicology*: "Even the convenience of computer-based forms of MIM cannot eliminate the need for MIM in book form. The preface provides a wonderful synopsis of human genetics. The information contained in this text serves as a concise review for those with a genetics background." From a review of the tenth edition, *New England Journal of Medicine*: "[Victor McKusick] has been for all these years the shepherd of the development of the field [of clinical genetics]. Perhaps his most important pragmatic achievement has been the 10 editions of Mendelian Inheritance in Man, which rapidly became and has remained the principal source of information on inherited diseases for all clinical geneticists. "In addition to the erudite entries in the books, the references given with each description represent a magnificent bibliography of clinical genetics. With McKusick's leadership and continued interest in gene mapping, the book also represents an important compendium of the location of genes on specific chromosomes. "The book is a magnificent security blanket for the clinical geneticist and should be in the libraries not only of these specialists, but also of all others who see patients with diseases that have genetic components."

VOGEL AND MOTULSKY'S HUMAN GENETICS

PROBLEMS AND APPROACHES

Springer Science & Business Media Provides information on the molecular basis of human genetics and outlines the principles of other epigenetic processes which together create the phenotype of a human being. This work also discusses the molecular basis for the concepts, methods and results in fields such as population genetics.

ADVANCES IN HUMAN GENETICS

Springer Science & Business Media From reviews of previous volumes in the series: 'Extremely valuable...thoroughly recommended.'-*Annals of Human Genetics* 'The most lucid and stimulating discussions of the topic to be found anywhere.'-*American Scientist*

MOLECULAR MEDICINE

GENOMICS TO PERSONALIZED HEALTHCARE

Elsevier Molecular medicine is the application of gene or DNA based knowledge to the modern practice of medicine. This book provides contemporary insights into how the genetic revolution is influencing medical thinking and practice on a broad front including clinical medicine, innovative therapies and forensic medicine. * Extensively revised just after the completion of the Human Genome Project, it provides the latest in molecular medicine developments * The only book in Molecular Medicine that has undergone 3 editions * Current practice as well as future developments identified * Extensive tables, well presented figures - resources for further understanding

PHARMACOGENETICS

Genes are important modifiers of human response to drugs, hormones, and toxins. Patients and healthy individuals alike display significant differences in response and suffer adverse effects as a result of exposure to many therapeutic agents as well as occupational chemicals. This introductory text brings together laboratory methods and epidemiologic studies for defining the role of heredity in human drug response. This book will benefit graduate students in pharmacology, genetics, epidemiology, nursing, and public health, and will serve as a handy reference for pharmacists, epidemiologists, and physicians responsible for the delivery and administration of drugs.

COLOR ATLAS OF GENETICS

Thieme From reviews of previous editions: A remarkable achievement concise but informative No geneticist or physician interested in genetic diseases should be without a copy -- *American Journal of Medical Genetics* Ever since the international Human Genome Project achieved its extraordinary goal of sequencing and mapping the entire human genome with far-reaching implications for understanding the causes and diagnosis of human genetic disorders progress in the field has been rapid. In the fourth edition of the bestselling *Color Atlas of Genetics*, readers will get a full overview of the field today, with an emphasis on the interface between fundamental principles and practical applications in medicine. The book utilizes the signature Flexibook format designed for easy visual learning and retention, and is invaluable for students, clinicians, and scientists interested in keeping current in this fast-moving area. New topics in the fully revised fourth edition of this highly praised atlas: Genetic signaling pathways involved in genetic disorders DNA repair systems Genomic disorders and genome-wide association studies Cancer genomes Ciliopathies, neurocristopathies, and other groups of causally related disorders Epigenetic changes in certain disorders Illustrated outline of human evolution With almost 200 stunning color plates concisely explained on facing pages, and including useful tables of data, a glossary of terms, key references, and online resources, this book makes every concept clear and accessible. It is an excellent introduction to genetics and basic genomics for students of medicine and biology, as well as an ideal teaching aid and refresher for investigators in any field of medicine or science.

DEVELOPMENTAL TOXICOLOGY

CRC Press The Second Edition of this highly regarded work provides a state-of-the-art review of developmental toxicology from basic science, clinical, epidemiological, and regulatory perspectives. This new edition highlights the latest approaches to understanding the mechanisms of developmental toxicity, testing pharmaceutical and environmental agents, and interpreting developmental toxicity data.; The contributors demonstrate how new information on molecular embryology and cell biology is being applied to problems in developmental toxicology. Chapters describe the effects of toxic exposure on the functional development of various organs, examine the relationship between maternal and developmental toxicity, and discuss current techniques for studying chemical disposition, metabolism, and placental transfer. Close attention is given to the use of mathematical and statistical techniques in data interpretation, as well as to the regulatory aspects of testing and risk assessment. Other chapters focus on pre- and post-conceptual clinical care and on genetic factors in clinical developmental toxicology.

HEALTH PROMOTION THROUGHOUT THE LIFE SPAN

Elsevier Health Sciences Up to date and easy to read, this textbook provides comprehensive coverage of all major concepts of health promotion and disease prevention. It highlights growth and development throughout the life span, emphasizing normal development as well as the specific problems and health promotion issues common to each stage. All population groups are addressed with separate chapters for individuals, families, and communities. -- Provided by publisher.

THE SELFISH GENE

Oxford University Press, USA An ethologist shows man to be a gene machine whose world is one of savage competition and deceit

AVERY'S NEONATOLOGY BOARD REVIEW

CERTIFICATION AND CLINICAL REFRESHER

Elsevier Health Sciences Prepare for success on the neonatology boards and in clinical practice with Avery's Neonatology Board Review: Certification and Clinical Refresher. This highly practical review tool follows the exam blueprint, is based on the trusted content found in Avery's Diseases of the Newborn, and has been carefully tailored for effective exam review by renowned neonatologist and educator, Patricia Chess, MD. Concise, relevant information is presented in a way that's easy to study and remember, giving you an important advantage on this challenging exam, as well as in the daily, fast-changing practice of neonatology. Equips residents, fellows, and physicians with an efficient, comprehensive system for study, designed specifically to help you perform at your best on the board exam. Presents information in a high-yield, outline format highlighted by key points, graphs, tables, images, and algorithms. Over 300 questions online Features board-style vignettes in every chapter with full, discursive answers online. Written by experts in the field of Neonatal-Perinatal Medicine, representing a wealth of trusted insight and guidance.

HOME CARE FOR THE HIGH-RISK INFANT

A FAMILY-CENTERED APPROACH

Jones & Bartlett Learning This revised, expanded edition addresses the increased interest in and demand for information about the nursing care of premature, technology-dependent infants once they are at home. In addition to clinical practice and research updates it includes care plans, assessment forms, sample care maps, and other hands-on materials which make it a valuable tool as well as a textbook.

KUMAR AND CLARK'S CLINICAL MEDICINE E-BOOK

Elsevier Health Sciences Edited by Adam Feather, MBBS, FRCP, FAcadMed; David Randall, MA, MRCP; and Mona Waterhouse, MA, MRCP Contributors comprise consultants at the top of their fields, paired with younger doctors closer to the exam experience, to ensure authority and relevance. Enhanced e-book accompanies the print book, for ease of transportation and use on the move. International Advisory Board, led by Professor Janaka de Silva and Professor Senaka Rajapakse, providing guidance for global coverage from across the world. Contributions to the e-book by members of the International Advisory Board to amplify areas of clinical importance in their parts of the world.