Informed Consent A Primer For Clinical Practice

Essential Ethics for Psychologists
Oxford Textbook of Palliative Medicine
Informed Consent
Research Design and Analysis
Self-determination in Health Care
Rethinking Informed Consent in Bioethics
Advances in Anesthesia, E-Book
2020 Clinical Dermatology Trials
101 Health and the Law
The Consent Primer
Promoting Legal and Ethical Awareness
Essential Clinical Procedures
E-Book
The Age of Informed Consent
Emerging Infectious Diseases
Informed Consent, Proxy Consent, and Catholic Bioethics
Mitochondrial Pathogenesis
Psychiatric Genetics
100 Activities for Teaching Research Methods
Cross-Cultural and Religious Critiques of Informed Consent
Legal Nurse Consulting Principles and Practices
Principles and Practice of Forensic Psychiatry, 2Ed
A Revised Consent Model for the Transplantation of Face and Upper Limbs:
Covenant Consent in Obesity Treatment
Ethical and Inclusive Research with Children
Pharmacogenomics: A Primer for Clinicians
Ethics Primer of the American Psychiatric Association
A Primer for Health Care Ethics
Ethical Ophthalmologist
Informed Consent for Human Subjects
Research
Informed Consent and Health Literacy
A Primer on Diseases of the Breast
The SAGE Encyclopedia of Economics and Society
Oxford Textbook of Obstetric Anaesthesia
Studying People
Oxford Textbook of Palliative Medicine
Cumulated Index Medicus
Pediatric Endoscopy, An Issue of Gastrointestinal Endoscopy Clinics of North America
The Narrative Approach to Informed Consent
Biostatistics and Epidemiology
Modern Pharmaceutical Industry

Written by foremost experts in the field, the 3rd Edition of Essential Clinical Procedures presents the latest common diagnostic and treatment-related procedures that you need to know as a physician assistant! Clear and concise, this medical reference book leads you step-by-step through over 70 techniques commonly seen in primary care and specialist settings, equipping you to deliver the best and safest care to your patients. Consult this title on your favorite e-reader, conduct rapid searches, and adjust font sizes for optimal readability. Visualize procedures and other important concepts clearly through over 300 high-quality illustrations. Ensure the safe execution of procedures with a focus on patient preparation and the proper use of instruments, as well as highlighted warnings to alert you to potential dangers that can occur when performing various treatments. Be fully prepared for any situation with guidance on the possible complications, rationales, and precautions behind many common procedures. Understand the ins and outs of nonprocedural issues, including informed consent, standard precautions, patient education, and procedure documentation. Get all the hands-on guidance you need to provide safe and effective care to your patients with Essential Clinical Procedures!

Understand the "must-know" aspects of the most commonly seen conditions with a systematic presentation featuring consistent headings and supporting color images. Quickly access core information through high-yield margin notes and consistent formatting across all chapters. Stay abreast of the latest procedures with a brand-new chapter and images on Reduction of the Shoulder/Finger Subluxations, plus many other updates throughout. For the new edition of Biostatistics and Epidemiology, Dr. Wassertheil-Smoller has included several new chapters (genetic statistics, molecular epidemiology, scientific integrity and research ethics) and a new appendix on the basic concepts of genetics and a glossary of genetic terminology. She has also expanded the coverage of multi-center trials (an important aspect of implementation of the standards of evidence-based medicine), controversies in screening for prostate, colon, breast, and other cancers. From Harry and Louise through the McCaughey septuplets, this book explains stories and issues in health care ethics that have appeared in the news media. Written for the general reader in a pluralistic society, it outlines and applies principles of justice from the Catholic tradition to contemporary problems that increasingly affect us all. This second edition contains extensive new material and new topics, including physician-assisted suicide, managed care, organ donation, genetic testing, cloning, and the question of futility. Aimed at a wide audience, this book will also be useful for introductory ethics courses in colleges and high schools. The literature on informed consent and its ethical-legal significance in clinical practice has grown rapidly in recent years. This unique book offers a practical description of the principles of informed consent and their application in daily clinical practice. Written by a team of experts in medical ethics and law, the chapters use a case-based approach to elucidate the essence of consent and highlight the ways in which individual patients and diverse situations can shape and even challenge the fundamental principles of informed consent. A range of situations in both primary and secondary care are covered and the content is arranged conceptually to help emphasise certain recurrent and related themes. An informative and rigorous yet accessible text, Informed Consent: A Primer for Clinical Practice is an essential resource for healthcare professionals working in all medical fields. Psychiatrists and other mental health professionals are increasingly confronted with questions about the genetics of psychiatric illness, and the clinical applications of new genetic findings. Psychiatric Genetics: A Primer for Clinical and Basic Scientists addresses these questions through a straightforward introduction to the essentials of psychiatric genetics, complementing more comprehensive textbooks that may seem overwhelming for those new to the field. Written and edited by leaders in the field and the International Society of Psychiatric Genetics (ISPG), the book covers basic epidemiology, recruitment for human studies, phenotyping strategies, formal genetic and molecular genetic studies, statistical genetics, bioinformatics and genomics, pharmacogenetics, the most relevant animal models, and biobanking. Each chapter begins with a list of "take home" points that summarizes content, followed by a brief overview of current knowledge and suggestions for further reading. This Primer is ideal for medical students, psychiatric residents, psychiatrists, and basic neuroscience researchers who are interested in learning about the key concepts and recent advances in the exciting field of psychiatric genetics.

In his latest book, Tom Nagy offers psychologists valuable lessons in ethical "orienteering." He begins by defining four points of our professional...
moral compass, competence, informed consent, confidentiality, and avoidance of harm and exploitation. Then he provides several decision-making maps and guides the reader through challenging, realistic ethical dilemmas in clinical, research, and education territories. This book reminds us that ethical practice is a career-long journey; I highly recommend it.

Janet L Sonne, PhD, Independent Clinical and Forensic Practice; Adjunct Professor of Psychology, Loma Linda University, Huntington Beach, CA; Fellow of the American Psychological Association Thomas Nagy's most recent book is replete with practical, accessible, and sound ethical guidance and wisdom. Straightforward explanations of basic ethical concepts provide a foundation on which he constructs sophisticated ethical analyses of issues arising in clinical, research, supervisory, training, and consultative settings. Compelling real-life vignettes illustrating applications of ethics codes and laws are seamlessly woven throughout the book, reflecting virtually every area of psychologists’ work. This book is a gift to students and seasoned professionals alike.

Janet T Thomas, PsyD, Independent Practice, Saint Paul, MN; Adjunct Faculty, Argosy University Twin Cities and the University of Saint Thomas, Minneapolis, MN; author, The Ethics of Supervision and Consultation: Practical Guidance for Mental Health Professionals

The second edition of this award-winning textbook has been thoroughly revised and updated throughout. Building on the success of the first edition, the book continues to address the History and Practice of Forensic Psychiatry, Legal Regulation of the Practice of Psychiatry, Psychiatry in relation to Civil Law, Criminal Law, and Family Law. Important sections such as Special Issues in Forensic Psychiatry, Law and the Legal System, and Landmark Cases in Mental Health Law are included. Designed to meet the needs of practitioners of forensic psychiatry, for residents in forensic psychiatry, and those preparing for the specialty examination in Forensic Psychiatry of the American Board of Psychiatry and Neurology, this volume will also answer the many questions faced by mental health professionals, mental health administrators, correctional health professionals and correctional health administrators, attorneys, judges, probation and parole officers and administrators all of whom, at one time or another, require a substantive presentation of the entire field of forensic psychiatry in the USA.

This work offers a comprehensive understanding rooted in Catholic anthropology and moral theory of the meaning and limits of informed and proxy consent to experimentation on human subjects. In particular, it seeks to articulate the rationale for proxy consent in both therapeutic and nontherapeutic settings. As to the former, the book proposes that the Golden Rule, recognizing the basic inclinations of human nature toward objective goods and goods of human persons, should underpin the notion of proxy consent to experimentation on humans. As to the latter, an additional scrutiny of the amount of risk involved is necessary, since the risk-benefit ratio frequently invoked to justify higher-risk therapeutic research does not exist in its nontherapeutic counterpart.

This study discusses a number of possible solutions to this question and develops a position that builds upon the objective notion of the human good.

Clinical Dermatology Trials 101 provides dermatologists with a handbook that allows them to become familiar with all aspects of clinical trials. Everything from obtaining the necessary tools and equipment, complying with local, federal, and international guidelines and regulations, and hiring and training staff for the safe and up-to-date conduct of dermatology clinical trials is covered. Written by leading experts in the field, Clinical Dermatology Trials 101 is the only clinical trial how-to available for dermatologists. With skin disease affecting nearly seventy percent of the population over a lifetime, and the rate of development of new drugs and devices for dermatologic use increasing at an exponential rate, there is a tremendous need for training and developing dermatology clinical research facilities to expedite the translation of basic and applied research, from bench to bedside. This is useful for practicing dermatologists, academic dermatologists, dermatology residents, clinical research fellows, dermatology fellows, research scientists, industry dermatologists, and medical students.

The definitive Oxford Textbook of Palliative Medicine now in its fifth edition, has again been thoroughly updated to offer a truly global perspective in this field of extraordinary talent and thoughtfulness. Updated to include new sections devoted to assessment tools, care of patients with cancer, and the management of issues in the very young and the very old, this leading textbook covers all the new and emerging topics since its original publication in 1993. In addition, the multi-disciplinary nature of palliative care is emphasized throughout the book, covering areas from ethical and communication issues, the treatment of symptoms and the management of pain. This fifth edition of the Oxford Textbook of Palliative Medicine is dedicated to the memory of Professor Geoffrey Hanks, pioneer in the field of palliative medicine, and co-editor of the previous four editions. Winner in the Medicine category of the British Medical Association Book Awards, this is a truly comprehensive text, no hospital, hospice, palliative care service, or medical library should be without this essential source of information.

This book supports the emerging field of vascularized composite allotransplantation (VCA) for face and upper-limb transplants by providing a revised, ethically appropriate consent model which takes into account what is actually required of facial and upper extremity transplant recipients. In place of consent as permission-giving, waiver, or autonomous authorization (the standard approaches), this book imagines consent as an ongoing mutual commitment, i.e. as covenant consent. The covenant consent model highlights the need for a durable personal relationship between the patient/subject and the care provider/researcher. Such a relationship is crucial given the recovery period of 5 years or more for VCA recipients. The case for covenant consent is made by first examining the field of vascularized composite allotransplantation, the history and present understandings of consent in health care, and the history and use of the covenant concept from its origins through its applications to health care ethics today. This book explains how standard approaches to consent are inadequate in light of the particular features of facial and upper limb transplantation. In contrast, use of the covenant concept creates a consent model that is more appropriate ethically for these very complex surgeries and long-term recoveries.

Over a short few decades, the field of pediatric endoscopy has matured from the exploratory to the routine. Performance of endoscopic procedures in children is now a fundamental aspect of the practice of more than 2000 pediatric gastroenterologists in North America, and endoscopic instruments are increasingly being developed with an eye to their...
pediatric applications. Ensuring safe and effective endoscopy in children requires specific medical knowledge and technical competency, in addition to appropriately designed equipment and settings. Obtaining consent from parents, as well as assent from patients, for the purposes of performing diagnostic and therapeutic gastrointestinal procedures begins with a deep understanding of risks and benefits that endoscopy affords and is typically gained through formal training in the field. Diagnostic endoscopy may help to confirm common pediatric conditions including eosinophilic esophagitis and inflammatory bowel disease, while therapeutic procedures to treat strictures in the GI tract may help children avoid more invasive surgeries. Using endoscopy in children to achieve hemostasis or to remove commonly swallowed foreign bodies, such as lithium batteries or high-powered magnets, can be lifesaving, and the insertion of feeding tubes can help medically complex patients to thrive. In short, pediatric endoscopy is an integral component of healthcare for children, and gaining and understanding of its best practices may help all clinicians to better recognize its role in pediatric disease outcomes. Designed as a ready-made teaching material, this book is divided into 12 chapters that each focus on one aspect of professional activity or behaviour, such as advertising, informed consent, the impaired physician, resource allocation and research. This book reviews quality definition, measurement, improvement, value, and accountability for obesity management. The interplay between quality, cost, access and satisfaction is fully depicted with a goal toward not only fulfilling current standards but also anticipating future needs. A thorough inventory of current best practices in all aspects of obesity care is catalogued with a gap analysis also employed for potential areas of improvement to be road mapped. All chapters are written by experts in their fields and include the most up-to-date scientific and clinical information, take home messages, and questions towards following the requirements of quality certification in obesity management. Quality in Obesity Treatment provides a comprehensive, contemporary review of this field and serves as a valuable resource for Bariatric Surgeons, Primary Care Physicians, Policy Makers, Insurance Administrators, Bariatricians, and any medical specialty interested in obesity quality management with likely candidates coming from GI, endocrinology, cardiology, sleep medicine and orthopedics. An invaluable resource to the rapidly emerging field of pharmacogenomics—complete with case studies, clinical pearls, and treatment recommendations—The aim of pharmacogenomics is to improve personalized medicine by taking into account how genes affect an individual's tolerability and response to drugs. Approaching pharmacogenomics from the current clinical, scientific, and implementation perspectives, this guide serves as an invaluable evidence-based resource to the subject. Reflecting the shift from genetics to genomics in the pharmaceutical sphere, the book covers pharmacogenomics fundamentals; genotyping tests and evidence; clinical implementation; ethical, legal, and social issues; and more. You'll also find illuminating case scenarios, clinical pearls, and evidence-based recommendations for treatments and alternatives based on CPIC, PharmGKB, and FDA guidelines. Consent is part of your life. Every day you interact with dozens, if not hundreds of people and consent plays a role every time. A role you're probably unaware of, at least until something goes wrong. This foundational book explores consent in a new way and will show you the fundamentals of Consent, how to use Consent in your relationships and sex life, and what to do when Consent goes wrong. This highly anticipated and comprehensive guide delves deep to explore the ins and outs of consent in our everyday lives. Regardless of whether you're brand new, or skilled at living a consensual life, this how-to guide is a beacon to set your course for better consent. In questions of psychiatric ethics, simple "yes" or "no" answers are rare, yet questions arise frequently in the clinical setting. "Should I accept a patient's invitation to a party?" "Is it OK to tell a patient that I, too, have had a depression?" "Can I release sensitive information about my patient without the patient's consent?" "Can I give a psychiatric opinion about someone I've never examined?" A shortage of ethics instruction from medical school through residency has left many psychiatrists facing the increasingly complex field of ethics without a clear guide to ethical decision making and conduct until now. Informed by some of the formal proceedings of the APA Ethics Committee, Ethics Primer of the American Psychiatric Association presents today's ethical dilemmas in eleven informative chapters -- brought to life by the clinical vignettes based on actual cases seen by this primer's distinguished contributors. Boundary violations between the doctor -- patient relationship and any other relationship (e.g., social, sexual/romantic, business) Issues such as informed consent, psychopharmacology, hospitalization, and psychotherapy with children, adolescents, and families such as confidentiality, competence and consent, quality of life, abuse and neglect, and use of restraints with geriatric populations. Involuntary hospitalization rife with conflict and controversy where many ethical principles clash (e.g., beneficence, autonomy, informed consent) Reconciling ethical conduct (as delineated in guidelines of the AMA, APA, and Sabin) with managed care's cost containment and rationing of medical services. Confidentiality, how this fundamental ethics principle can clash with the business ethics of managed care and insurance companies. Gifts from patients and industry when and why they are or are not acceptable. The often uncomfortable duty to report colleagues who engage in fraud or deception, from speaking privately with a colleague to referring a concern to a departmental committee, supervisor, or local APA ethics committee (and applicable legal mandates) Emergency care ethics how to ensure proper treatment for psychiatric patients who come to the emergency room with physical illnesses forensic psychiatry, including codes of ethics, boundary issues, and management of allegations of ethical misconduct When consultations and second opinions are needed. Including an appendix with selected material from The Principles of Medical Ethics With Annotations Especially Applicable to Psychiatry, this clinical guide and reference is sure to stimulate the discussion so integral to maintaining the dynamic tradition of ethics. As such, it is essential reading for every psychiatrist -- whether in training or in established clinical practice -- who aspires to a richer appreciation for the wisdom and subtleties of the guidelines and principles of medical ethics. Advances in Anesthesia highlights the year's significant medical advances, providing one source to review the essential information updates for the Anesthesiologist in that year. The distinguished editorial board, led by Dr. Thomas McLoughlin, includes Drs. Richard Dutton, Laurence Torsher, and Francis Salinas.
The board has assembled a first-rate volume for 2020, including topics on motor-sparing nerve blocks; whole blood; teaching airway management skills to non-anesthesiologists; endovascular procedures; regional anesthesia for cardiac surgery, ketamine; consent; medical cannabis; and more. Informed consent - the process of communication between a patient or research subject and a physician or researcher that results in the explicit agreement to undergo a specific medical intervention - is an ethical concept based on the principle that all patients and research subjects should understand and agree to the potential consequences of the clinical care they receive. Regulations that govern the attainment of informed consent for treatment and research are crucial to ensuring that medical care and research are conducted in an ethical manner and with the utmost respect for individual preferences and dignity. These regulations, however, often require - or are perceived to require - that informed consent documents and related materials contain language that is beyond the comprehension level of most patients and study participants. To explore what actions can be taken to help close the gap between what is required in the informed consent process and communicating it in a health-literate and meaningful manner to individuals, the Institute of Medicine’s Roundtable on Health Literacy convened a one-day public workshop featuring presentations and discussions that examine the implications of health literacy for informed consent for both research involving human subjects and treatment of patients. Topics covered in this workshop included an overview of the ethical imperative to gain informed consent from patients and research participants, a review of the current state and best practices for informed consent in research and treatment, the connection between poor informed consent processes and minority underrepresentation in research, new approaches to informed consent that reflect principles of health literacy, and the future of informed consent in the treatment and research settings. Informed Consent and Health Literacy is the summary of the presentations and discussion of the workshop. Economics is the nexus and engine that runs society, affecting societal well-being, raising standards of living when economies prosper or lowering citizens through class structures when economies perform poorly. Our society only has to witness the booms and busts of the past decade to see how economics profoundly affects the cores of societies around the world. From a household budget to international trade, economics ranges from the micro- to the macro-level. It relates to a breadth of social science disciplines that help describe the content of the proposed encyclopedia, which will explicitly approach economics through varied disciplinary lenses. Although there are encyclopedias of covering economics (especially classic economic theory and history), the SAGE Encyclopedia of Economics and Society emphasizes the contemporary world, contemporary issues, and society. Features: 4 volumes with approximately 800 signed articles ranging from 1,000 to 5,000 words each are presented in a choice of print or electronic editions Organized A-to-Z with a thematic Reader’s Guide in the front matter groups related entries Articles conclude with References & Future Readings to guide students to the next step on their research journeys Cross-references between and among articles combine with a thorough Index and the Reader's Guide to enhance search-and-browse in the electronic version Pedagogical elements include a Chronology of Economics and Society, Resource Guide, and Glossary This academic, multi-author reference work will serve as a general, non-technical resource for students and researchers within social science programs who seek to better understand economics through a contemporary lens. Modern Pharmaceutical Industry: A Primer comprehensively explains the broad range of divisions in the complex pharmaceutical industry. Experts actively involved in each component discuss their own contribution to a pharmaceutical company's work and success. Divisions include regulatory affairs, research and development, intellectual property, pricing, marketing, generics, OTC, and more. The seventeen chapters included in this resource offer a wide range of topics, from discovery and formulation to post-approval and legal. Readers will be given a detailed look at the structure of a contemporary drug company and a thorough understanding of what goes on behind the scenes. Modern Pharmaceutical Industry: A Primer is a valuable resource for all pharmacy students, new hires at pharmaceutical companies, drug company management, and academic health center libraries. No other text provides a comprehensive look at one of the most dynamic industries related to the modern healthcare system. Combining the best of author Ron Scott’s books, Promoting Legal Awareness in Physical and Occupational Therapy and Professional Ethics: A Guide for Rehabilitation Professionals, his newest text Promoting Legal and Ethical Awareness: A Primer for Health Professionals and Patients includes the latest case, regulatory, and statutory law. This valuable ethical and legal resource also includes an alphabetized section on HIPAA, current information on the reauthorized IDEA (Individuals with Disabilities Act), and expanded coverage of alternative dispute resolution and attorney-health professional-client relations. Cases and Questions allow you to apply key legal and ethical principles to a rehabilitation practice situation. Special Key Term boxes introduce and define important vocabulary to ensure your understanding of chapter content. Additional resource lists in each chapter include helpful sources for articles, books, and websites to further your learning. Case Examples let you put new ideas and concepts into practice by applying your knowledge to the example. Legal Foundations and Foundations chapters introduce the basic concepts of law, legal history, the court system, and ethics in the professional setting to provide a solid base for legal and ethical knowledge. An entire chapter devoted to healthcare malpractice provides vital information on practice problems that have legal implications, the claim process, and claim prevention. An extended discussion of the Americans with Disabilities Act informs you of your rights as an employee as well as the challenges faced in the workforce by your rehabilitation patients. Content on employment legal issues includes essential information for both employees and employers on patient interaction and the patient’s status in the workplace. Coverage of end-of-life issues and their legal and ethical implications provides important information for helping patients through end-of-life decisions and care. This outstanding text and reference provides health professionals & students with a balanced, comprehensive, highly readable survey of the legal concepts and controversies affecting them today. Avoiding unnecessarily technical language, it lucidly explains basic legal principles and theories, examines current issues and their implications, and probes future
legal trends. Throughout, each chapter offers a complete, self-contained introduction to a medicolegal topic—including invaluable endnotes that cite references, clarify perspectives, and suggest further readings. A unique appendix also explains how to use law library facilities to best advantage. The Narrative Approach to Informed Consent: Empowering Young Children’s Rights and Meaningful Participation is a practical guide for researchers who want to engage young children in rights-based, participatory research. This book presents the Narrative Approach, an original and innovative method to help children understand their participation in research. This approach moves away from traditional paper-based consent to tailor the informed consent process to the specific needs of young children. Through the Informing Story, which employs a combination of interaction, information and narrative, this method enables children to comprehend concepts through storytelling. Researchers are steered through the development of an Informing Story so that they can deliver accurate information to young children about what their participation in research is likely to involve. To further inform practice, the book documents the implementation of the Narrative Approach in four case studies demonstrating the variety of settings in which the method can be applied. The Narrative Approach to Informed Consent addresses the rights of young children to be properly researched, expands opportunities for their active and engaged research participation, and creates a unique conceptual ethical space within which meaningful informed consent can occur. This book will be an invaluable tool for novice and experienced researchers and is applicable to a wide range of education and non-education contexts. Written by an established, comprehensive, multidisciplinary focused, internationally-recognized team, the sixth edition has been fully revised and brought up to date with the inclusion of recent developments in the specialty, to ensure that it retains its reputation as the definitive reference on palliative care. A concise, straightforward overview of research design and analysis, helping readers form a general basis for designing and conducting research. The practice of designing and analyzing research continues to evolve with advances in technology that enable greater technical analysis of data—strengthening the ability of researchers to study the interventions and relationships of factors and assisting consumers of research to understand and evaluate research reports. Research Design and Analysis is an accessible, wide-ranging overview of how to design, conduct, analyze, interpret, and present research. This book helps those in the sciences conduct their own research without requiring expertise in statistics and related fields and enables informed reading of published research. Requiring no background in statistics, this book reviews the purpose, ethics, and rules of research, explains the fundamentals of research design and validity, and describes how to select and employ appropriate statistical techniques and reporting methods. Readers gain knowledge central to various research scenarios, from sifting through reports of meta-analyses and preparing a research paper for submission to a peer-reviewed journal to discussing, evaluating, and communicating research results. This book: Provides end-to-end guidance on the entire research design and analysis process Teaches readers how to both conduct their own research and evaluate the research of others Offers a clear, concise introduction to fundamental topics ideal for both reference and general education functions Presents information derived from the author’s experience teaching the subject in real-world classroom settings Includes a full array of learning tools including tables, examples, additional resource suggestions, complete references, and appendices that cover statistical analysis software and data sets Research Design and Analysis: A Primer for the Non-Statistician is a valuable source of information for students and trainees in medical and allied health professions, journalism, education, and those interested in reading and comprehending research literature. It is generally accepted in legal and bioethical discourse that the patient has a right to self-determination. In practice though, this is often not the case. Paternalism is waning and it is increasingly recognised that there are values other than medical factors which determine the choices that patients make. Unfortunately, these developments have not resulted in huge advances for patient self-determination, which is largely because the consent model has fundamental flaws that constrain its effectiveness. This book sets out to offer an alternative model to consent. In the property model proposed here, the patient’s bodily integrity is protected from unauthorised invasion, and their legitimate expectation to be provided with the relevant information to make an informed decision is taken to be a proprietary right. It is argued that the property model potentially overcomes the limitations of the consent model, including the obstacle caused by the requirement to prove causation in consent cases. The author proposes that this model could in the future provide an alternative or complementary approach for the courts to consider when dealing with cases relating to self-determination in health care. A sourcebook of exercises, games, scenarios and role plays, this practical, user-friendly guide provides a complete and valuable resource for research methods tutors, teachers and lecturers. Developed to complement and enhance existing course materials, the 100 ready-to-use activities encourage innovative and engaging classroom practice in seven areas: finding and using sources of information planning a research project conducting research using and analyzing data disseminating results acting ethically developing deeper research skills. Each of the activities is divided into a section on tutor notes and student handouts. Tutor notes contain clear guidance about the purpose, level and type of activity, along with a range of discussion notes that signpost key issues and research insights. Important terms, related activities and further reading suggestions are also included. Not only does the A4 format make the student handouts easy to photocopy, they are also available to download and print directly from the book’s companion website for easy distribution in class. Legal Nurse Consulting Principles and Practices, Fourth Edition, provides foundational knowledge on the specialty nursing practice of legal nurse consulting. Legal nurse consulting is defined, and essential information about the practice is discussed (history, certification, scope and standards of practice, and ethical and liability considerations). The essentials of the law and medical records are explored. Analysis of the various types of legal cases on which legal nurse consultants work is provided, as are other practice areas for legal nurse consultants. The various roles and skills of legal nurse consultants are explored, and the textbook concludes with discussion of the ways in which legal cases are adjudicated. This volume allows nurses
to bridge the gap from their clinical experience to the unfamiliar territory of the legal world, with practical advice on topics including tactics for being cross-examined in the courtroom and investigative and analytical techniques for medical records. Individual chapters by subject-matter experts focus on the full range of legal, medical, and business issues that new or experienced legal nurse consultants and nurse experts will encounter in their work. A nuanced look at the realities and complexities of toxic torts, medical malpractice cases, civil rights in correctional healthcare, ERISA and HMO litigation, and other practice areas is offered. Suitable for experienced nurses studying for certification as legal nurse consultants, and for expert witnesses, practitioners seeking to expand their current legal nurse roles, and other healthcare and legal practitioners. This volume re-evaluates informed consent in multicultural contexts and features perspectives from Buddhism, Confucianism, Hinduism, Christianity, Judaism and Islam. It is valuable reading for scholars interested in bioethics, healthcare ethics, research ethics, comparative religions, theology, human rights, law and sociology. From early conception until the postpartum period, anaesthetists are required to provide pregnant women with the highest standard of anaesthetic care and pain relief whilst negotiating challenges such as concurrent systemic disease, infertility, and obesity as well as practising in accordance with new developments in fetal medicine surgery, pharmacology, and imaging. The Oxford Textbook of Obstetric Anaesthesia provides an up-to-date summary of the scientific basis, assessment for and provision of anaesthesia throughout pregnancy and labour. This highly authoritative textbook is conceptually divided into nine sections, detailing maternal and fetal physiology, fetal and neonatal assessment and therapy, anaesthesia before and during pregnancy, labour and vaginal delivery, anaesthesia for caesarean delivery, anaesthetic and obstetric complications, as well as systemic disease. Individual chapters address topics such as simulation and ultrasound. Authored by an international team of expert anaesthetists this textbook reflects current world-wide practice and guidelines. Designed for consultants and trainees in anaesthesia, the Oxford Textbook of Obstetric Anaesthesia is the definitive source of expert knowledge for anaesthetists in this subspecialty. The imperative to include children and young people in educational research, and in more participatory ways, is educationally important when exploring policy and practice contexts. It is also critical to recognise that children have the right to contribute to debates, and can express their views through educational research, on matters that affect them. However, the freedom to research alongside young people is only afforded if we continue to unmask the illusion that well-intentioned research is always ethical. This book presents an international set of storied experiences, where researchers have been challenged and have changed the way they think, incorporating and exploring ethics in research. The contributors highlight the ethical dilemmas that can arise when children and young people are included in research agendas, and their reflexive approaches to these dilemmas include being responsive to the cultural, political and social contexts of the lives of the children and developing child-friendly research approaches to ensure their ‘voice’ is accessed in multiple ways. These solution-focused and local approaches facilitate a more ethical, deliberative process where the establishment of trust is central to an ethical engagement with young people and their families and where the expression of ethical dilemmas can improve research practice. This book is a critical resource for researchers and practitioners researching with and alongside children and young people. This book was originally published as a special issue of the International Journal of Inclusive Education. Informed consent is a central topic in contemporary biomedical ethics. Yet attempts to set defensible and feasible standards for consenting have led to persistent difficulties. In Rethinking Informed Consent in Bioethics, first published in 2007, Neil Manson and Onora O'Neill set debates about informed consent in medicine and research in a fresh light. They show why informed consent cannot be fully specific or fully explicit, and why more specific consent is not always ethically better. They argue that consent needs distinctive communicative transactions, by which other obligations, prohibitions, and rights can be waived or set aside in controlled and specific ways. Their book offers a coherent, wide-ranging and practical account of the role of consent in biomedicine which will be valuable to readers working in a range of areas in bioethics, medicine and law.