Health Care Ethics Committees

The Next Generation

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Health Care Ethics Committees Judith Wilson Ross, M.A., 1993-08-29 Approximately 85% of hospitals now have ethics committees But this statistic says little about the efficiency and importance of these committees in their institutions Frequently ethics committees exist more in name than in practice and are left without the guidance and help of their institution Health Care Ethics Committees provides a plethora of advice including possible projects and activities suggestions for making meetings more effective insights into policy making and models for mission statements and goals In addition this book gives leaders a panoramic view of the past present and future of ethics committees in health care **Ethics Committees** Judith Wilson Ross, 1993-01-01 Handbook for Health Care Ethics Committees Linda Farber Post, Jeffrey Blustein, 2015-06-30 How can dedicated ethics committees members fulfill their complex roles as moral analysts policy reviewers and clinical consultants The Joint Commission TJC accredits and certifies more than 19 000 health care organizations in the United States including hospitals nursing homes and home care agencies Each organization must have a standing health care ethics committee to maintain its status These interdisciplinary committees are composed of physicians nurses attorneys ethicists administrators and interested citizens Their main function is to review and provide resolutions for specific individual patient care problems Many of these committees are well meaning but may lack the information experience skills and formal background in bioethics needed to adequately negotiate the complex ethical issues that arise in clinical and organizational settings Handbook for Health Care Ethics Committees was the first book of its kind to address the myriad responsibilities faced by ethics committees including education case consultation and policy development Adopting an accessible tone and using a case study format the authors explore serious issues involving informed consent and refusal decision making and decisional capacity truth telling the end of life palliative care justice in and access to health care services and organizational ethics. The authors have thoroughly updated the content and expanded their focus in the second edition to include ethics committees in other clinical settings such as long term care facilities small community hospitals rehabilitation centers and hospices They have added three new chapters that address reproduction disability and the special needs of the elder population and they provide additional specialized policies and procedures on the book s website This guide is an essential resource for all health care ethics committee members Ethical Issues in Neurology James L. Bernat, 2008 Written by an eminent authority from the American Academy of Neurology's Committee on Ethics Law and Humanities this book is an excellent text for all clinicians interested in ethical decision making The book features outstanding presentations on dying and palliative care physician assisted suicide and voluntary active euthanasia medical futility and the relationship between ethics and the law New chapters in this edition discuss how clinicians resolve ethical dilemmas in practice and explore ethical issues in neuroscience research Other highlights include updated material on palliative sedation advance directives ICU withdrawal of life sustaining therapy gene therapy the very low birth weight premature infant the

developmentally disabled patient informed consent organizational ethics brain death controversies and fMRI and PET studies relating to persistent vegetative state **Health Care Ethics: Critical Issues for the 21st Century** Eileen E. Morrison, 2008-03-20 The Ethics of Rapid Tissue Donation (RTD) Chiara Mannelli, 2021-02-11 This book offers a reflection on the central role that the ethics of informed consent plays in Rapid Tissue Donation RTD RTD is an advanced oncology procedure that involves the procurement for research purposes of fresh tissues within two to six hours of a cancer patient s death Since RTD involves the retrieval of tissues after death and since the collected tissues are of great importance for medical research the need for any form of informed consent to regulate this procedure has been questioned This book argues for the necessity of informed consent to govern RTD and it provides the reader with a bespoke informed consent process applicable to cancer patients The analysis unfolds at the intersection between applied ethics public health ethics and clinical ethics and it is informed by philosophical theories of informed consent and by the social implications of individual choices By viewing medical issues relating to informed consent in oncology from an ethical perspective the book combines philosophical analysis with discussion of concrete cancer related issues As a result the book is suitable for readers interested in ethical reasoning as well as for those with a medical background It contributes to contemporary research by offering an original analysis that relies on a rigorous philosophical approach to address innovative issues at the cutting edge of medical research and policy making **Pharmacoethics** David A. Gettman, Dean Arneson, 2003-04-29 Due to the changing nature of the practice of pharmacy today s pharmacists pharmaceutical scientists and researchers are faced with an increasing amount of ethical dilemmas Pharmacoethics A Problem Based Approach not only introduces the current ethical issues it also provides decision making tools that can be applied to any ethical issue that Clinical Bioethics C. Viafora, 2005-07-19 A theory of Clinical Bioethics based on the integration of the moral logic of health care practice internal morality and the larger social concerns and processes external morality Clinical Bioethics A Search for the Foundations compares major theoretical models in the foundation of clinical bioethics and explains medicine as a normative practice. The goals of medicine are discussed with particular reference to the subjectivisation of health and the rationalisation of health care institutions This volume provides a consistent reconstruction of bioethical judgment both at the level of epistemological statute and institutional context i e clinical ethics committees and clinical ethics consultation Bioethical and Evolutionary Approaches to Medicine and the Law W. Noel Keyes, 2007 Bioethics is a multidisciplinary field of law and one that can not be ignored Bioethical and Evolutionary Approaches to Medicine and the Law is a comprehensive scholarly analysis of bioethics and the development of its standards The book is broken up into the following four parts Part I deals with scientific religious ethical and legal aspects of bioethics Part II evaluates 100 current bioethical issues and sets forth specific approaches for their resolution Part III focuses on medical legal and other problems from beginning of life overpopulation birth control in vitro fertilization etc through end of life physician assisted suicide advance directives euthanasia etc Part IV discusses the major bioethical issues

in genetics and genetic engineering Advance Health Care Directives Carol Krohm, Scott K. Summers, 2002 This volume deals with the difficult and sensitive issues faced by lawyers doctors nurses clerics and spiritual advisors in helping clients and patients plan write execute and implement personal contingency plans for health care decision making **Library of Medicine Current Catalog** National Library of Medicine (U.S.),1993 First multi year cumulation covers six Engineering Ethics Rosa Lynn B. Pinkus, 1997-05-13 Using the space shuttle programme as the framework this book examines ethical decision making in engineering **Protecting American Health Care Consumers** Eleanor D. Kinney, 2002-05-08 A contribution to health care studies and administrative law which offers a humane and practical alternative to the current process of reviewing consumer health care complaints Ethics by Committee D. Micah Hester, 2008 Ethics by Committee was developed for tens of thousands of people across the United States who serve on hospital ethics committees HECs Experts in bioethics clinical consultation health law and social psychology from across the country have contributed chapters on ethics consultation education and policy development The chapters discuss important considerations for HEC members such as promoting just and ethical organizations developing cultural and spiritual awareness and preparing for the forces of group dynamics in committee discussions and consensus building No other book on the market offers the diversity of perspectives and topics while remaining focused clear and useful Book jacket The Birth of Bioethics Albert R. Jonsen, 2003-08-28 Bioethics represents a dramatic revision of the centuries old professional ethics that governed the behavior of physicians and their relationships with patients This venerable ethics code was challenged in the years after World War II by the remarkable advances in the biomedical sciences and medicine that raised questions about the definition of death the use of life support systems organ transplantation and reproductive interventions In response philosophers and theologians lawyers and social scientists joined together with physicians and scientists to rethink and revise the old standards Governments established commissions to recommend policies Courts heard arguments and legislatures passed laws This book is the first broad history of the growing field of bioethics Covering the period 1947 1987 it examines the origin and evolution of the debates over human experimentation genetic engineering organ transplantation termination of life sustaining treatment and new reproductive technologies It assesses the contributions of philosophy theology law and the social sciences to the expanding discourse of bioethics Written by one of the field s founders The Birth of Bioethics is based on extensive archival research into sources that are difficult to obtain and on interviews with many of the leading figures in the moral debates in medicine A very readable and comprehensive account of the evolution of bioethics this book stresses the history of ideas but does not neglect the social and cultural context and the people involved It will serve the information needs of philosophers ethicists social historians and everyone interested in the origins of some of today s most hotly debated issues Clearinghouse Review ,1995 When Professionals Weep Renee S. Katz, Therese A. Johnson, 2016-01-22 When Professionals Weep speaks to the humbling and often transformational moments that clinicians

experience in their careers as caregivers and healers moments when it is often hard to separate the influence of our own emotional responses and worldviews from the patient s or family s When Professionals Weep addresses these poignant moments when the professional s personal experiences with trauma illness death and loss can subtly often stealthily surface and affect the helping process This edition like the first both validates clinicians experiences and also helps them process and productively address compassion fatigue burnout and secondary traumatic stress New material in the second edition includes increased emphasis on the burgeoning fields of hospice and palliative care organizational countertransference mindfulness and compassionate practice It includes thought provoking cases self assessments and exercises that can be used on an individual dvadic or group basis This volume is an invaluable handbook for practitioners in the fields of medicine mental health social work nursing chaplaincy the allied health sciences psychology and psychiatry **Legal Perspectives in Bioethics** Ana S. Iltis, Sandra H. Johnson, Barbara A. Hinze, 2007-12-19 Incisive and thought provoking this volume provides readers with a rich context for understanding the intersection between the law on bioethics and the central issues in Bioethics Mediation Nancy Neveloff Dubler, Carol B. Liebman, 2021-04-30 Bioethics Mediation offers stories bioethics about patients families and health care providers enmeshed in conflict as they wrestle with decisions about life and death It provides guidance for those charged with supporting the patient's traditional and religious commitments and personal wishes Today's medical system without intervention privileges those within shared cultures of communication and disadvantages those lacking power and position such as immigrants the poor and nonprofessionals This book gives clinical ethics consultants palliative care providers and physicians nurses and other medical staff the tools they need to understand and manage conflict while respecting the values of patients and family members Conflicts come in different guises and the key to successful resolution is early identification and intervention Every bioethics mediator needs to be prepared with skills to listen level the playing field identify individual interests explore options and help craft a principled resolution a consensus that identifies a plan aligned with accepted ethical principles legal stipulations and moral rules and that charts a clear course of future intervention The organization of the book makes it ideal for teaching or as a handbook for the practitioner It includes actual cases modified to protect the privacy of patients providers and institutions detailed case analyses tools for step by step mediation techniques for the mediator sample chart notes and a set of actual role plays with expert mediator and bioethics commentaries The role plays include discharge planning for a dying patient an at risk pregnancy HIV and postsurgical complications in the ICU treatment for a dying adolescent dialysis and multiple systems failure Expanded by two thirds from the 2004 edition the new edition features two new role plays a new chapter on how to write chart notes and a discussion of new understandings of the role of the clinical ethics consultant Taking Advance Directives Seriously Robert S. Olick, 2001-07-18 In the quarter century since the landmark Karen Ann Quinlan case an ethical legal and societal consensus supporting patients rights to refuse life sustaining treatment has become a cornerstone of bioethics Patients now

legally can write advance directives to govern their treatment decisions at a time of future incapacity yet in clinical practice their wishes often are ignored Examining the tension between incompetent patients prior wishes and their current best interests as well as other challenges to advance directives Robert S Olick offers a comprehensive argument for favoring advance instructions during the dying process He clarifies widespread confusion about the moral and legal weight of advance directives and he prescribes changes in law policy and practice that would not only ensure that directives count in the care of the dying but also would define narrow instances when directives should not be followed Olick also presents and develops an original theory of prospective autonomy that recasts and strengthens patient and family control While focusing largely on philosophical issues the book devotes substantial attention to legal and policy questions and includes case studies throughout An important resource for medical ethicists lawyers physicians nurses health care professionals and patients rights advocates it champions the practical ethical and humane duty of taking advance directives seriously where it matters most at the bedside of dying patients

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