

# Medical Ethics Policies Protocols Guidelines Programs

**Medical Ethics** John F. Monagle 1992 This manual is a compendium of various health care policies, guidelines, protocols, and programs that concern clinical issues with ethical implications. The collection of policies, guidelines, and procedures are helpful in drafting and reviewing institutional procedures and helping policymakers develop useful mechanisms for assuring ethical treatment of patients and staff.

**The Belmont Report** United States. National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research 1978

Reports on Managed Care American Medical Association. Council on Ethical and Judicial Affairs 1998 Ethics in the era of managed care This collection of AMA Council Reports from 1990 to 1997 examine a variety of ethical issues concerning managed care. Report topics include financial incentives to limit care, cost containment involving prescription drugs, restrictions on disclosure in managed care contracts, ethical issues in negotiating discounts for specialty care, capitation, and more. An analysis of current issues in medical ethics is also included.

**Code of Medical Ethics of the American Medical Association** American Medical Association 1897

Core Ethics for Health Professionals Robert F. Phalen 2017-08-10 A timely overview of ethics, emphasizing applications to biomedical researchers, health providers, and administrators There are no simple rules to guide ethical conduct in daily practice, health professionals must have a basic understanding of several topics including ethical theories; ethical scandals; laws, regulations, and institutional policies; and public perceptions. This book can be used for self-study, for classroom instruction, and as a refresher and update by practicing health professionals. The chapters have learning objectives, focused content, a summary of important points, a quiz, and a list of key references. Although the book is

arranged in a logical order, each chapter may be studied independently.

**Improving Healthcare Quality in Europe Characteristics, Effectiveness and Implementation of Different Strategies**

OECD 2019-10-17 This volume, developed by the Observatory together with OECD, provides an overall conceptual framework for understanding and applying strategies aimed at improving quality of care. Crucially, it summarizes available evidence on different quality strategies and provides recommendations for their implementation. This book is intended to help policy-makers to understand concepts of quality and to support them to evaluate single strategies and combinations of strategies.

*Code of Ethics for Nurses with Interpretive Statements* American Nurses Association 2001 Pamphlet is a succinct statement of the ethical obligations and duties of individuals who enter the nursing profession, the profession's nonnegotiable ethical standard, and an expression of nursing's own understanding of its commitment to society. Provides a framework for nurses to use in ethical analysis and decision-making.

**Complementary and Alternative Medicine in the United States** Institute of Medicine

2005-05-13 Integration of complementary and alternative medicine therapies (CAM) with conventional medicine is occurring in hospitals and physicians offices, health maintenance organizations (HMOs) are covering CAM therapies, insurance coverage for CAM is increasing, and integrative medicine centers and clinics are being established, many with close ties to medical schools and teaching hospitals. In determining what care to provide, the goal should be comprehensive care that uses the best scientific evidence available regarding benefits and harm, encourages a focus on healing, recognizes the importance of compassion and caring, emphasizes the centrality of relationship-based care, encourages patients to share in decision making about therapeutic options, and promotes choices in care that can include complementary therapies where appropriate.

Numerous approaches to delivering integrative medicine have evolved. Complementary and Alternative Medicine in the United States identifies an urgent need for health systems research that focuses on identifying the elements of these models, the outcomes of care delivered in these models, and whether these models are cost-effective when compared to conventional practice settings. It outlines areas of research in convention and CAM therapies, ways of integrating these therapies, development of curriculum that provides further education to health professionals, and an amendment of the Dietary Supplement Health and Education Act to improve quality, accurate labeling, research into use of supplements, incentives for privately funded research into their efficacy, and consumer protection against all potential hazards.

Medical Ethics John F. Monagle 1995

**International ethical guidelines for health-related research involving humans** Council for International Organizations of Medical Sciences (CIOMS) 2016-12-06 Progress towards a world where all can enjoy optimal health and health care is crucially dependent on all kinds of research including research involving humans. Involving humans in medical research is necessary to improve the knowledge base on which medicine should be based. At the same time, individuals participating in health-related research have individual human rights and have a right to be protected against the risks that research may bring to them. The tension between these two considerations has led the medical community to endorse ethical guidelines for health-related research. Research Ethics Committees can use these guidelines to evaluate whether a given research protocol is ethically acceptable or not. CIOMS, in association with the World Health Organization, started its work on ethics in health-related research in the late 1970s. Accordingly, CIOMS set out, in cooperation with WHO, to prepare guidelines to indicate how the ethical principles set forth in the Declaration of Helsinki of the World Medical Association, could be effectively applied, particularly in low-resource settings, given their socio-economic circumstances, laws and regulations, and executive and administrative arrangements. Since then revised editions of the

CIOMS ethical guidelines were published in 1993 and 2002. New developments in research have prompted CIOMS to again revise their ethical guidelines. The result is now available in this new publication. In the new 2016 version of the ethical Guidelines, CIOMS provides answers to a number of pressing issues in research ethics. The Council does so by stressing the need for research having scientific and social value, by providing special guidelines for health-related research in low-resource settings, by detailing the provisions for involving vulnerable groups in research and for describing under what conditions biological samples and health-related data can be used for research. In providing this new version, CIOMS hopes to ensure that the ethical Guidelines remain a living document that provides reasoned conditions for research in order to meet the challenges of modern research.

**Responsibility in Health Care** G.J. Agich 2012-12-06 Medicine is a complex social institution which includes biomedical research, clinical practice, and the administration and organization of health care delivery. As such, it is amenable to analysis from a number of disciplines and directions. The present volume is composed of revised papers on the theme of "Responsibility in Health Care" presented at the Eleventh Trans Disciplinary Symposium on Philosophy and Medicine, which was held in Springfield, Illinois on March 16-18, 1981. The collective focus of these essays is the clinical practice of medicine and the themes and issues related to questions of responsibility in that setting. Responsibility has three related dimensions which make it a suitable theme for an inquiry into clinical medicine: (a) an external dimension in legal and political analysis in which the State imposes penalties on individuals and groups and in which officials and governments are held accountable for policies; (b) an internal dimension in moral and ethical analysis in which individuals take into account the consequences of their actions and the criteria which bear upon their choices; and (c) a comprehensive dimension in social and cultural analysis in which values are ordered in the structure of a civilization ([8], p. 5). The title "Responsibility in Health Care" thus signifies a broad inquiry not only into the ethics of individual character and

actions, but the moral foundations of the cultural, legal, political, and social context of health care generally.

**Guidelines for Clinical Practice** Institute of Medicine 1992-02-01 Guidelines for the clinical practice of medicine have been proposed as the solution to the whole range of current health care problems. This new book presents the first balanced and highly practical view of guidelines—their strengths, their limitations, and how they can be used most effectively to benefit health care. The volume offers: Recommendations and a proposed framework for strengthening development and use of guidelines. Numerous examples of guidelines. A ready-to-use instrument for assessing the soundness of guidelines. Six case studies exploring issues involved when practitioners use guidelines on a daily basis. With a real-world outlook, the volume reviews efforts by agencies and organizations to disseminate guidelines and examines how well guidelines are functioning—exploring issues such as patient information, liability, costs, computerization, and the adaptation of national guidelines to local needs.

**Current Catalog** National Library of Medicine (U.S.) 1993

**Public Health Ethics: Cases Spanning the Globe** Drue H. Barrett 2016-04-20 This Open Access book highlights the ethical issues and dilemmas that arise in the practice of public health. It is also a tool to support instruction, debate, and dialogue regarding public health ethics.

Although the practice of public health has always included consideration of ethical issues, the field of public health ethics as a discipline is a relatively new and emerging area. There are few practical training resources for public health practitioners, especially resources which include discussion of realistic cases which are likely to arise in the practice of public health. This work discusses these issues on a case to case basis and helps create awareness and understanding of the ethics of public health care. The main audience for the casebook is public health practitioners, including front-line workers, field epidemiology trainers and trainees, managers, planners, and decision makers who have an interest in learning about how to integrate ethical analysis into their day to day public

health practice. The casebook is also useful to schools of public health and public health students as well as to academic ethicists who can use the book to teach public health ethics and distinguish it from clinical and research ethics.

**Hippocratic Writings** Hippocrates 2005-05-26 This work is a sampling of the Hippocratic Corpus, a collection of ancient Greek medical works. At the beginning, and interspersed throughout, there are discussions on the philosophy of being a physician. There is a large section about how to treat limb fractures, and the section called The Nature of Man describes the physiological theories of the time. The book ends with a discussion of embryology and a brief anatomical description of the heart.

**Medical Ethics Manual** John Reynold Williams 2005

**Clinical Practice Guidelines** Institute of Medicine 1990-02-01 The Alberta clinical practice guidelines program is supporting appropriate, effective and quality medical care in Alberta through promotion, development and implementation of evidence-based clinical practice guidelines.

**Clinical Ethics** Albert R. Jonsen 1992 Clinical Ethics introduces the four-topics method of approaching ethical problems (i.e., medical indications, patient preferences, quality of life, and contextual features). Each of the four chapters represents one of the topics. In each chapter, the authors discuss cases and provide comments and recommendations. The four-topics method is an organizational process by which clinicians can begin to understand the complexities involved in ethical cases and can proceed to find a solution for each case.

**Registries for Evaluating Patient Outcomes** Richard E. Gliklich 2014

**Responsible Conduct of Research** Adil E. Shamoo 2003 This is a comprehensive introduction to the ethical issues at stake in the conduct of biomedical research, with extensive use of case examples. Its content parallels the recommendations of the Commission on Research Integrity, and deals with ethical issues in the use of animals and humans in research. It includes chapters on intellectual property, authorship, peer review, and conflicts of interest. As of October 2000, all personnel

involved in research supported by the Public Health Service, including NIH, must receive the equivalent of 15 hours of training and education in research ethics. This book will be a convenient text for such short courses or seminars, and an excellent guidebook for all.

### **Institutional Protocols for Decisions about Life-sustaining Treatments 1988**

**Final Report: Sources and documentation**  
United States. Advisory Committee on Human Radiation Experiments 1995

*The Medical Ethics Advisor Sourcebook* 1996

### **Change Dynamics in Healthcare, Technological Innovations, and Complex Scenarios** Burrell, Darrell Norman 2024-02-26

In a world characterized by complexity and rapid change, the intersection of healthcare, social sciences, and technology presents a formidable challenge. The vast array of interconnected issues, ethical dilemmas, and technological advancements often evade comprehensive understanding within individual disciplines. The problem lies in the siloed approach to these critical domains, hindering our ability to navigate the complexities of our modern world effectively. *Change Dynamics in Healthcare, Technological Innovations, and Complex Scenarios* emerges as a transformative solution, offering a beacon of insight and knowledge to those grappling with the intricate dynamics of our interconnected society. *Change Dynamics in Healthcare, Technological Innovations, and Complex Scenarios* dives into organizational narratives, ethical challenges, and technological promises across healthcare, social sciences, and technology. It doesn't merely acknowledge the interplay between these disciplines; it celebrates their interconnectedness. By dissecting, analyzing, and synthesizing critical developments, this book serves as a compass, providing a rich resource for comprehending the multifaceted impacts of emerging changes.

Library of Congress Subject Headings Library of Congress 2013

*Advanced Practice Palliative Nursing* Constance Dahlin 2016-02-25 *Advanced Practice Palliative Nursing* is the first text devoted to advanced practice nursing care of the seriously ill and dying. This comprehensive work addresses all aspects of palliative care including physical, psychological, social, and spiritual needs.

Chapters include: symptoms common in serious illness, pediatric palliative care, spiritual and existential issues, issues around the role and function of the advanced practice nurse (APN), reimbursement, and nursing leadership on palliative care teams. Each chapter contains case examples and a strong evidence base to support the highest quality of care. The text is written by leaders in the field and includes authors who have pioneered the role of the advanced practice nurse in palliative care. This volume offers advanced practice content and practical resources for clinical practice across all settings of care and encompassing all ages, from pediatrics to geriatrics.

**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. *The Patient Self-Determination Act* Lawrence P. Ulrich 2001-07-18 The Patient Self-Determination Act of 1990 required medical facilities to provide patients with written notification of their right to refuse or consent to medical treatment. Using this Act as an important vehicle for improving the health care decisionmaking process, Lawrence P. Ulrich explains the social, legal, and ethical background to the Act by focusing on well-known cases such as those of Karen Quinlan and Nancy Cruzan, and he explores ways in which physicians and other caregivers can help patients face the complex issues in contemporary health care practices. According

to Ulrich, health care facilities often address the letter of the law in a merely perfunctory way, even though the Act integrates all the major ethical issues in health care today. Ulrich argues that well-designed conversations between clinicians and patients or their surrogates will not only assist in preserving patient dignity — which is at the heart of the Act—but will also help institutions to manage the liability issues that the Act may have introduced. He particularly emphasizes developing effective advance directives. Ulrich examines related issues, such as the negative effect of managed care on patient self-determination, and concludes with a seldom-discussed issue: the importance of being a responsible patient. Showing how the Patient Self-Determination Act can be a linchpin of more meaningful and effective communication between patient and caregiver, this book provides concrete guidance to health care professionals, medical ethicists, and patient-rights advocates.

**The American Medical Ethics Revolution**

Robert Baker 1999-12-13 D.--from the Introduction "Canadian Bulletin of Medical History"

Library of Congress Subject Headings Library of Congress. Subject Cataloging Division 1980

**Principles of Biomedical Ethics** James F. Childress 1983

*National Library of Medicine Current Catalog* National Library of Medicine (U.S.) 1993

**Advisory Committee on Human Radiation Experiments: Sources and documentation**

United States. Advisory Committee on Human Radiation Experiments 1995

**Medical 911** 1994

**Institutional protocols for decisions about life-sustaining treatments : special report.**

United States. Congress. Office of Technology Assessment 1988

Non-Heart-Beating Organ Transplantation

Institute of Medicine 1998-01-18 Non-heart-beating donors (individuals whose deaths are determined by cessation of heart and respiratory function rather than loss of whole brain function) could potentially be of major importance in reducing the gap between the demand for and available supply of organs for transplantation. Prompted by questions concerning the medical management of such

donors—specifically, whether interventions undertaken to enhance the supply and quality of potentially transplantable organs (i.e. the use of anticoagulants and vasodilators) were in the best interests of the donor patient—the U.S. Department of Health and Human Services asked the Institute of Medicine to examine from scientific and ethical points of view "alternative medical approaches that can be used to maximize the availability of organs from [a] donor [in an end-of-life situation] without violating prevailing ethical norms...." This book examines transplantation supply and demand, historical and modern conceptions of non-heart-beating donors, and organ procurement organizations and transplant program policies, and contains recommendations concerning the principles and ethical issues surrounding the topic.

Health Care Ethics Eileen E. Morrison

2009-10-06 This thorough revision of health care ethics brings the reader up to date on the most important issues in biomedical ethics today.

Interpretive Phenomenology Patricia Benner

1994-05-17 Patricia Benner's philosophical introduction to phenomenology develops the reader's understanding of the strategies and processes involved in this approach to human science. Contributors to the volume discuss the constitutive relationships between theory and practice, consider the possibility of a science of caring from a feminist perspective, introduce interpretive phenomenology for studying natural groups such as families, and suggest a ground for developing nursing ethics that is true to the caring and healing practices of nurses.

Following a thorough and engaging exposition of the field's theoretical and philosophical foundations, the work shifts focus to interpretive studies currently being undertaken within the scholarly community; the research tradition is then applied and reexamined as it relates to specific lines of inquiry. Interpretive Phenomenology presents an inclusive and well-integrated discussion of the many correlative topics within this subject area. Its relevance and accessibility will make this book an invaluable resource for advanced undergraduate and graduate students, as well as professional researchers in nursing and other health sciences. "The authors share a deep regard for



the primacy of practice and caring. This is not the ethereal caring of many current theorists. . . . Interpretive Phenomenology presents an inclusive and well-integrated discussion of the author's philosophical and practical approach, looking especially at the relationship between theory and practice." --Journal of Christian Nursing

Ethics in Health Administration Eileen E. Morrison 2008-03-20 New Edition Available 5/1/2013 Building on the wisdom and forward thinking of authors John Monagle and David Thomasa, this thorough revision of Health Care Ethics: Critical Issues for the 21st Century brings the reader up-to-date on the most important issues in biomedical ethics today. **Health Care Ethics: Critical Issues for the 21st Century** Eileen E. Morrison 2008-03-20 New Edition Available 5/1/2013 Building on the wisdom and forward thinking of authors John Monagle and David Thomasa, this thorough revision of Health Care Ethics: Critical Issues for the 21st Century brings the reader up-to-date on the most important issues in biomedical ethics today.

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