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Medical ethics refers to moral principles usually applied in the medical practice to ensure that professional, ethical, and religious concepts are utilized effectively among health care practitioners. Health care practitioners should use medical ethics during the analysis of medical cases.

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1367 Words | 6 Pages. the principles of medical ethics. Medical ethics are a set of moral principles that apply values and judgments to the practice of medicine. The four values of autonomy, beneficence, non-maleficence, and justice are commonly considered to be the fundamentals within medical ethics. Medical ethics encompasses its practical application in clinical settings, and while these values do not give the answers as.

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Medical ethics refers to the discussion and application of moral values and responsibilities in the areas of medical practice and research. While questions of medical ethics have been debated since the beginnings of Western medicine in the fifth century B.C., medical ethics as a distinctive field came into prominence only since World War II.

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Begin your ethics in health care research paper by recognizing that the decisions made by healthcare professionals, be it physicians, nurses or medical staff, affect real people and may mean the difference between life and death, including abortion. The health and welfare of patients, along with the very serious aspect of treatment facilitation, requires that ethical standards be followed every step of the way for the health care professional.

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The American College of Physicians' Ethics Manual is designed to help facilitate the process of making ethical decisions in clinical practice and medical research. Some aspects of medicine are fundamental and timeless. Medical practice, however, does not stand still. The fifth edition of the Ethics Manual examines emerging issues in medical ethics faced by internists and their patients and revisits older issues that are still very pertinent. The distinction and potential conflicts between legal and ethical obligations when making clinical

decisions are discussed. New or expanded sections have been added including third party evaluations, confidentiality, complementary and alternative care, boundaries and privacy, gifts from patients, care of patients at the end of life, solid organ transplantation, physician-assisted suicide, the changing practice environment and managed care of ethics, physician-industry issues, selling products out of the office, health and human rights, patient safety, prisoners as patients, strikes and joint actions, consultation and shared care, and research ethics. Sections on genetic testing, the changing practice environment and managed care, organ donation, and disability certification have been added.

Bioethics has long been accepted as an interdisciplinary field. The recent 'empirical turn' in bioethics is, however, creating challenges that move beyond those of simple interdisciplinary collaboration, as researchers grapple with the methodological, empirical and meta-ethical challenges of combining the normative and the empirical, as well as navigating the difficulties that can arise from attempts to transcend traditional disciplinary boundaries. *Empirical Bioethics: Theoretical and Practical Perspectives* brings together contributions from leading experts in the field which speak to these challenges, providing insight into how they can be understood and suggestions for how they might be overcome. Combining discussions of meta-ethical challenges, examples of different methodologies for integrating empirical and normative research, and reflection on the challenges of conducting and publishing such work, this book will both introduce the novice to the field and challenge the expert.

#1 NEW YORK TIMES BESTSELLER □ “The story of modern medicine and bioethics—and, indeed, race relations—is refracted beautifully, and movingly.”—Entertainment Weekly NOW A MAJOR MOTION PICTURE FROM HBO® STARRING OPRAH WINFREY AND ROSE BYRNE □ ONE OF THE “MOST INFLUENTIAL” (CNN), “DEFINING” (LITHUB), AND “BEST” (THE PHILADELPHIA INQUIRER) BOOKS OF THE DECADE □ ONE OF ESSENCE’S 50 MOST IMPACTFUL BLACK BOOKS OF THE PAST 50 YEARS □ WINNER OF THE CHICAGO TRIBUNE HEARTLAND PRIZE FOR NONFICTION NAMED ONE OF THE BEST BOOKS OF THE YEAR BY The New York Times Book Review □ Entertainment Weekly □ O: The Oprah Magazine □ NPR □ Financial Times □ New York □ Independent (U.K.) □ Times (U.K.) □ Publishers Weekly □ Library Journal □ Kirkus Reviews □ Booklist □ Globe and Mail Her name was Henrietta Lacks, but scientists know her as HeLa. She was a poor Southern tobacco farmer who worked the same land as her slave ancestors, yet her cells—taken without her knowledge—became one of the most important tools in medicine: The first “immortal” human cells grown in culture, which are still alive today, though she has been dead for more than sixty years. HeLa cells were vital for developing the polio vaccine; uncovered secrets of cancer, viruses, and the atom bomb’s effects; helped lead to important advances like in vitro fertilization, cloning, and gene mapping; and have been bought and sold by the billions. Yet Henrietta Lacks remains virtually unknown, buried in an unmarked grave. Henrietta’s family did not learn of her “immortality” until more than twenty years after her death, when scientists investigating HeLa began using her husband and children in research without informed consent. And though the cells had launched a multimillion-dollar industry that sells human biological materials, her family never saw any of the profits. As Rebecca Skloot so brilliantly shows, the story of the Lacks family—past and present—is inextricably connected to the dark history of experimentation on African Americans, the birth of bioethics, and the legal battles over whether we control the stuff we are made of. Over the decade it took to uncover this story, Rebecca became enmeshed in the lives of the Lacks family—especially Henrietta’s daughter Deborah. Deborah was consumed with questions: Had scientists cloned her mother? Had they killed her to harvest her cells? And if her mother was so important to medicine, why couldn’t her children afford health insurance? Intimate in feeling, astonishing in scope, and impossible to put down, *The Immortal Life of Henrietta Lacks* captures the beauty and drama of scientific discovery, as well as its human consequences.

The goal of this open access book is to develop an approach to clinical health care ethics that is more accessible to, and usable by, health professionals than the now-dominant approaches that focus, for example, on the application of ethical principles. The book elaborates the view that health professionals have the emotional and intellectual resources to discuss and address ethical issues in clinical health care without needing to rely on the expertise of bioethicists. The early chapters review the history of bioethics and explain how academics from outside health care came to dominate the field of health care ethics, both in professional schools and in clinical health care. The middle chapters elaborate a series of concepts, drawn from philosophy and the social sciences, that set the stage for developing a framework that builds upon the individual moral experience of health professionals, that explains the discontinuities between the demands of bioethics and the experience and perceptions of health professionals, and that enables the articulation of a full theory of clinical ethics with clinicians themselves as the foundation. Against that background, the first of three chapters on professional education presents a general framework for teaching clinical ethics; the second discusses how to integrate ethics into formal health care curricula; and the third addresses the opportunities for teaching available in clinical settings. The final chapter, "Empowering Clinicians", brings together the various dimensions of the argument and anticipates potential questions about the framework developed in earlier chapters.

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.

Research universities are critical contributors to our national research enterprise. They are the principal source of a world-class labor force and fundamental discoveries that enhance our lives and the lives of others around the world. These institutions help to create an educated citizenry capable of making informed and crucial choices as participants in a democratic society. However many are concerned that the unintended cumulative effect of federal regulations undercuts the productivity of the research enterprise and diminishes the return on the federal investment in research. Optimizing the Nation's Investment in Academic Research reviews the regulatory framework as it currently exists, considers specific regulations that have placed undue and often unanticipated burdens on the research enterprise, and reassesses the process by which these regulations are created, reviewed, and retired. This review is critical to strengthen the partnership between the federal government and research institutions, to maximize the creation of new knowledge and products, to provide for the effective training and education of the next generation of scholars and workers, and to optimize the return on the federal investment in research for the benefit of the American people.

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