

# A Casebook Of Medical Ethics

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 The Oxford Textbook of Clinical Research Ethics  
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 An Introduction  
 A Casebook of Ethical Challenges in Neuropsychology  
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 Clinical Ethics Casebook  
 Clinical Ethics  
 Well and Good  
 A Graphic Medicine Casebook

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## SANTOS HEATH

**A Casebook** Oxford University Press, USA

This casebook consists of hundreds of case vignettes drawn from a variety of areas of clinical practice. It presents over 200 brief, concrete examples along with specific questions and alternatives that highlight the challenging ethical problems involved. This book can be read independently or can serve as a supplement to any core text in clinical/medical ethics.

**Muted Consent** Vanderbilt University Press

I. Defining "research"--II. Issues in study design . -- III. Harm and benefit -- IV. Voluntary informed consent -- V. Standard of care -- VI. Obligations to participants and communities -- VII. Privacy and confidentiality -- VIII. Professional ethics.

*Beyond a Western Bioethics* OUP USA

Interesting and important ethical questions confront researchers, regulators, institutional review boards, support personnel, and research participants committed to the ethical conduct of human subjects research at all stages of research. Questions encompass - but are not limited to - study design, enrolling participants, balancing the clinical needs of participants against the research agenda, ending trials, discharging post-trial obligations, and resolving conflicts. Straightforward solutions to these types of questions are often not found in regulations, ethics codes, or the bioethics literature. These resources may leave room for interpretation, offer conflicting guidance, or simply fail to address particular questions. Ethics consultation, which has been offered in clinical care settings with regularity since the 1980s, has since the turn of the century increasingly been sought in the clinical research context. Because there has only lately been recognition that ethics consultants can play a valuable role helping the research community conduct research in the most ethically informed way, there are many open questions in the field of research ethics consultation including the appropriate role of consultants and the best methods of consultation. The Clinical Center Bioethics Consultation Service has been serving the NIH community of researchers, administrators, healthcare providers, and research participants for more than a decade, conducting nearly 1,000 consultations in that time. In this book, members of the Bioethics Consultation Service reflect on this long track-record and unparalleled range of research ethics consultations to share a collection of their most interesting and informative research ethics consultations and to start a dialogue on remaining open questions. Although the NIH experience is unique, this book focuses on cases - and associated lessons - that are generalizable and valuable for the entire clinical research community. This book will be valuable to ethics consultants, clinical investigators, students and teachers, and others desiring insight into clinical research ethics and ethics consultation.

*The Oxford Textbook of Clinical Research Ethics* Penn State Press

Every accredited American hospital is required to have a mechanism for handling ethical concerns; most hospitals satisfy this requirement by constituting an institutional healthcare ethics committee (HEC), a pattern which is repeated in most western countries. This text provides definitive, comprehensive guidance for members of healthcare ethics committees who find themselves confronted with ethically challenging situations. Each chapter includes learning objectives, clinical case studies and questions to stimulate discussion among committee members. Particular emphasis is given to consultation, as this often presents the greatest challenges to committee members. Each chapter stands alone as a teaching module, as well as forming part of a comprehensive volume. Written and edited by nationally and internationally recognized experts in bioethics, this is essential reading for every member of a healthcare ethics committee.

**A Casebook from the Files of Experts** Wadsworth Publishing Company

Ethical Issues in International Biomedical Research is the definitive book on the ethics of research

involving human subjects in developing countries. Using 21 actual case studies, it covers the most controversial topics, including the ethics of placebo research in Africa, what benefits should be provided to the community after completion of a research trial, how to address conflicts between IRBs in developed and developing countries, and undue inducement of poor people in developing countries. Each case is accompanied by two expert commentaries, written by many of the worlds leading experts in bioethics as well as new voices with research experience in developing countries. No other volume has this scope. Students in bioethics, public and international health, and ethics will find this book particularly useful.

*Cases and Materials* Purdue University Press

#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.

*A Clinical Introduction to Medical Ethics Systems Theory* Springer

You decide! Learn to identify and effectively evaluate ethical problems in health care. The hundreds of brief, readable cases in this book will help you do just that. Author Peter Horn, a philosopher and applied ethicist who has taught clinical ethics for 20 years, includes cases drawn from a variety of areas of clinical practice. Discussion questions following each case help you appreciate the moral complexity of the situations and offer guidance for structuring possible answers.

**Tough Decisions** Amer Psychological Assn

When we are patients, few of us understand the implications and risks of the complex procedures modern medicine has developed for curing diseases and altering consciousness and human biology. Here is a book that attempts to clarify the issues raised by such complexities. The work is a primer



- [The Wager: A Tale Of Shipwreck, Mutiny And Murder By David Grann](#)
- [Goodnight Moon By Margaret Wise Brown](#)
- [Lessons In Chemistry: A Novel By Bonnie Garmus](#)
- [November 9: A Novel](#)
- [Things We Hide From The Light \(knockemout Series, 2\) By Lucy Score](#)
- [The Covenant Of Water \(oprah's Book Club\)](#)
- [Little Blue Truck's Springtime: An Easter And Springtime Book For Kids By Alice Schertle](#)