

Principles of Biomedical Ethics

> Page numbers reference Garrett et. al., Health Care Ethics. Prentice Hall, 2nd Edition, 1993, unless otherwise indicated.

Additional References:

1. Beauchamp and Childress, Principles of Biomedical Ethics, Fourth Edition. Oxford. 1994.
2. Engelhardt, H. Tristram Oxford: The Foundations of Bioethics, second edition, Oxford. 1996
3. Hall and Ellman, Health Care Law and Ethics, West. 1990.
4. Junkerman and Schiedermaier, Practical Ethics for Students, Interns, and Residents. University Publishing Group, 1998
5. American Medical Association Code of Medical Ethics, 1996
6. American College of Physicians Ethics Manual, Third Edition, 1993.

[These texts are available at the Erlanger Hospital Medical Library].

A standard approach to biomedical ethics, developed by Beauchamp and Childress in Principles of Biomedical Ethics, resolves ethical issues in terms of four ethical principles:

Autonomy [pages 3-6]

Beneficence & non-maleficence [pages 6-10]

Justice [pages 10-12]

Each of which need to be weighed and balanced in determining an optimal course of action.

All of these principles require a *conversation* about the needs and desires of the patient or, in the case of justice, members of community. The principles are intended to guide. In the case of autonomy, we are required to determine the wishes of the patient in order to protect his or her autonomy. In the case of beneficence and non-maleficence, we are required to determine the patient's views of what does and does not count as goods to be pursued or harms to be avoided. In the case of justice, we are required to follow due process in order to determine just limits on health care that will be generally accepted.

The demand for conversation arises out of what Engelhardt sees as the result of the break down of any traditional consensus about moral and religious goods. In particular, he argues that in the post-modern period, we no longer accept the possibility of constructing a content-full account of the human good. We engage one another, consequently, as moral strangers, who need to negotiate moral arrangements. Those negotiations are governed by the principles of autonomy (he calls it permission) and beneficence, but even in the case of beneficence we cannot presume a general agreement about human goods and their ranking. Engelhardt argues that the traditional distinctions used to distinguish moral categories of treatment assume a vision of the good that is not available to us.

For better or for worse, health care practitioners find themselves in a situation where they need to become moral negotiators with moral strangers.

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Mental Health among African Americans Erlanger A. Turner, 2019-09-25 In *Mental Health among African Americans* Innovations in Research and Practice Erlanger A Turner presents a new theoretical framework for understanding mental health disparities that emphasizes the need for culturally sensitive clinical practices and integration of Afrocentric values in order to address the lower rates of African Americans seeking treatment in the United States Turner traces this reluctance to the unethical scientific research practices that characterized experiments in recent history like the well known Tuskegee Syphilis study and stresses the need for providers to address race related stress **Library Journal** ,1962

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Ethics Manual American College of Physicians (2003-). Ethics and Human Rights Committee, Lois Snyder, 2005 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

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Table of Contents Erlanger Medical Ethics Orientation Manual May

1. Understanding the eBook Erlanger Medical Ethics Orientation Manual May
 - The Rise of Digital Reading Erlanger Medical Ethics Orientation Manual May
 - Advantages of eBooks Over Traditional Books
2. Identifying Erlanger Medical Ethics Orientation Manual May
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Erlanger Medical Ethics Orientation Manual May
 - User-Friendly Interface
4. Exploring eBook Recommendations from Erlanger Medical Ethics Orientation Manual May
 - Personalized Recommendations
 - Erlanger Medical Ethics Orientation Manual May User Reviews and Ratings
 - Erlanger Medical Ethics Orientation Manual May and Bestseller Lists
5. Accessing Erlanger Medical Ethics Orientation Manual May Free and Paid eBooks
 - Erlanger Medical Ethics Orientation Manual May Public Domain eBooks
 - Erlanger Medical Ethics Orientation Manual May eBook Subscription Services

- Erlanger Medical Ethics Orientation Manual May Budget-Friendly Options
- 6. Navigating Erlanger Medical Ethics Orientation Manual May eBook Formats
 - ePub, PDF, MOBI, and More
 - Erlanger Medical Ethics Orientation Manual May Compatibility with Devices
 - Erlanger Medical Ethics Orientation Manual May Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Erlanger Medical Ethics Orientation Manual May
 - Highlighting and Note-Taking Erlanger Medical Ethics Orientation Manual May
 - Interactive Elements Erlanger Medical Ethics Orientation Manual May
- 8. Staying Engaged with Erlanger Medical Ethics Orientation Manual May
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Erlanger Medical Ethics Orientation Manual May
- 9. Balancing eBooks and Physical Books Erlanger Medical Ethics Orientation Manual May
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Erlanger Medical Ethics Orientation Manual May
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Erlanger Medical Ethics Orientation Manual May
 - Setting Reading Goals Erlanger Medical Ethics Orientation Manual May
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Erlanger Medical Ethics Orientation Manual May
 - Fact-Checking eBook Content of Erlanger Medical Ethics Orientation Manual May
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
- 14. Embracing eBook Trends

- Integration of Multimedia Elements
- Interactive and Gamified eBooks

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