



THE PATIENT- PHYSICIAN RELATION

The Patient as Partner, Part 2

Robert M. Veatch

Patient Physician Relation Pt 2 The Patient As Partner

Robert M. Veatch



Patient Physician Relation Pt 2 The Patient As Partner:

The Patient-Physician Relation Robert M. Veatch,1991-01-22 Throughout the past two decades when medical ethics has had a renaissance Robert Veatch has been a leading contributor to its dialogue and advance This collection of his work shows the breadth and the cogency of his thinking it is a book worth having Journal of the American Medical Association a fascinating dissection of almost every aspect of the doctor patient relationship strongly recommended reading for all health care workers interested in this rapidly evolving field Queen s Quarterly This outstanding discussion of important current medical issues is a valuable addition to academic and professional libraries Choice an important contribution to bioethics certain to provoke controversy in the field Medical Humanities Review Lucid and well argued Religious Studies Review This book heralds the imminent demise of doctor knows best In it Robert M Veatch proposes a postmodern medicine in which decisions about patient care will routinely involve both doctor and patient not only in ethically complex cases such as the termination of life sustaining treatment but in everyday care as well

The Nature of the Doctor-Patient Relationship Pierre Mallia,2012-08-01 This book serves to unite biomedical principles which have been criticized as a model for solving moral dilemmas by inserting them and understanding them through the perspective of the phenomenon of health care relationship Consequently it attributes a possible unification of virtue based and principle based approaches

Informed Choice of Medical Services: Is the Law Just? Marj Milburn,2017-09-29 This title was first published in 2001 This work is a uniquely multi disciplinary contribution to the existing bioethical literature on the topic of informed choice of medical services It is also the first comprehensive bioethical text to confront the central issue of power in the clinical encounter and to argue for statutory protection of the right to informed choice While the majority of bioethicists argue for a conciliatory rather than adversarial approach to the chronic problem of uninformed consent the author of this work argues that the external regulation of medicine is essential if the right to informed choice is to be protected This argument is based upon an extensive review of the bioethical legal political medical sociological and philosophical literature as well as a wide range of empirical and anecdotal evidence evolving from a detailed exploration of power and the limits of rationality in the clinical encounter

Conflicts of Interest and the Future of Medicine Marc A. Rodwin,2013-10-03 In this book Marc Rodwin examines the development of conflicts of interest in the health care systems of the US France and Japan He shows that national differences in the organization of medical practice and the interplay of organized medicine the market and the state give rise to variations in the type and prevalence of such conflicts and then analyzes the strategies that each nation employs to cope with them Drawing on the experiences of these three nations Conflicts of Interest and the Future of Medicine demonstrates that we can mitigate these problems with carefully planned reform and regulation

The Basics of Bioethics Laura K. Guidry-Grimes,Robert M. Veatch,2019-10-18 The Basics of Bioethics Fourth Edition offers an easy to follow introduction to this dynamic field intended for healthcare professionals teachers students and anyone interested in bioethics

Accessible and enjoyable for readers of all backgrounds the book contains numerous cases including ones that recently have dominated international headlines to help anchor the broader discussion The text is suitable for use in short courses in schools of medicine nursing and other health professions continuing professional education various undergraduate departments and adult education Chapters are organized around common moral themes in order to help readers understand the values and other connections that tie together different positions in bioethics This fourth edition adds a new chapter on alternative frameworks in bioethics including narrative ethics and casuistry feminist approaches care ethics and virtue ethics Due to significant advances in genetics and reproductive possibilities this new edition devotes a full chapter to each The combined teaching research and clinical experience of the two authors helps make this edition current with the evolving field of bioethics while still embedding the major issues in a systematic framework that allows readers easily to navigate the larger field

Key Changes to the Fourth Edition An added chapter on new and emerging approaches in bioethics including those based on virtue ethics casuistry and narrative ethics feminist ethics and care ethics Updates throughout the book based on developments in ethical theory and new medical research Revisions and updates to the Learning Objectives Key Terms Bibliographies and URLs The addition of multiple recent case studies including Jahi McMath an undocumented patient who needs a rule bent a pediatrician who turns away unvaccinated patients a minor eligible for pediatric bariatric surgery a daughter suing a hospital for non disclosure of her father s Huntington s diagnosis CRISPR edited newborn babies

Theological Analyses of the Clinical Encounter G.P. McKenny, J.R. Sande, 2013-04-17 Efforts to evaluate the clinical encounter in terms of autonomous agents governed by rationally justified moral principles continue to be criticised These essays written by physicians ethicists theologians and philosophers examine various models of the clinical encounter emerging out of these criticisms and explore the prospects they offer for theological and religious discourse Individual essays focus on the reformulation of covenant models revisions of principles approaches and topics such as power authority narrative rhetoric dialogue and alterity The essays display a range of conclusions about whether theology articulates generally accessible religious insights or is a tradition specific discipline Hence the volume reflects current debates in theology while analysing current models of the clinical encounter Students professionals and scholars who find themselves at the intersection of theology and medicine will welcome these voices in an ongoing conversation

Making Medical Decisions for the Profoundly Mentally Disabled Norman L. Cantor, 2009-08-21 A legal and moral analysis of medical decision making on behalf of those with such severe cognitive impairments that they cannot exercise self determination In this book Norman Cantor analyzes the legal and moral status of people with profound mental disabilities those with extreme cognitive impairments that prevent their exercise of medical self determination He proposes a legal and moral framework for surrogate medical decision making on their behalf The issues Cantor explores will be of interest to professionals in law medicine psychology philosophy and ethics as well as to parents guardians and health care providers who face perplexing

issues in the context of surrogate medical decision making The profoundly mentally disabled are thought by some moral philosophers to lack the minimum cognitive ability for personhood Countering this position Cantor advances both theoretical and practical arguments for according them full legal and moral status He also argues that the concept of intrinsic human dignity should have an integral role in shaping the bounds of surrogate decision making Thus he claims while profoundly mentally disabled persons are not entitled to make their own medical decisions respect for intrinsic human dignity dictates their right to have a conscientious surrogate make medical decisions on their behalf Cantor discusses the criteria that bind such surrogates He asserts contrary to popular wisdom that the best interests of the disabled person are not always the determinative standard the interests of family or others can sometimes be considered Surrogates may even consistent with the intrinsic human dignity standard sometimes authorize tissue donation or participation in nontherapeutic medical research by profoundly disabled persons Intrinsic human dignity limits the occasions for such decisions and dictates close attention to the preferences and feelings of the profoundly disabled persons themselves Cantor also analyzes the underlying philosophical rationale that makes these decision making criteria consistent with law and morals

Wondergenes Maxwell J. Mehlman, 2003-09-16 Wondergenes not only imagines a future world in which genetic enhancement is the norm but asserts that this future has already begun Genetically engineered substances are already in use by athletes in vitro fertilization already provides the primitive means by which parents can select an embryo and the ability to create new forms of genetically engineered human beings is not far off What happens when gene therapy becomes gene enhancement Who will benefit and who might be left behind What are the costs to our values and beliefs and to the future of our society To answer these questions Maxwell J Mehlman provides an overview of the scientific advances that have led to the present state of genetic enhancement and explains how these advances will be used in the future to redefine what we think of as a normal human being He explores the ethical dilemmas already facing researchers and medical practitioners and the dilemmas we will all be expected to face In his forecast of the dangers inherent in this technology he is particularly concerned with the emergence of a genobility made up of those able to afford increasingly expensive enhancement Wondergenes is a serious accessible introduction to the social and personal implications of genetic engineering Mehlman weighs the social and economic costs of the many proposals to regulate or limit genetic engineering and provides six concrete policy recommendations from professional licensing to a ban on germ line enhancement that propose to make the future of genetic enhancement more equitable and safe

Health Reform Daniel Drache, Terry Sullivan, 2005-08-18 Health Reform explores the challenges facing health care provision in the advanced economies The book exposes the limitations of market led health reform and demonstrates the indispensable role of a vibrant public authority in the renewal of modern health care systems Issues covered include cost containment and privatisation strategies in an international perspective the role of business and the private sector in setting the agenda for health care reform the restructuring of Anglo Saxon health systems and the shift

in state market boundaries in Canada the USA the UK and Australia the frontier of health care reform in terms of health and social cohesion the role of patient choice in health care reform Conflicts of Conscience in Health Care Holly Fernandez Lynch, 2010-08-13 A balanced proposal that protects both a patient's access to care and a physician's ability to refuse to provide certain services for reasons of conscience Physicians in the United States who refuse to perform a variety of legally permissible medical services because of their own moral objections are often protected by conscience clauses These laws on the books in nearly every state since the legalization of abortion by Roe v Wade shield physicians and other health professionals from such potential consequences of refusal as liability and dismissal While some praise conscience clauses as protecting important freedoms opponents concerned with patient access to care argue that professional refusals should be tolerated only when they are based on valid medical grounds In Conflicts of Conscience in Health Care Holly Fernandez Lynch finds a way around the polarizing rhetoric associated with this issue by proposing a compromise that protects both a patient's access to care and a physician's ability to refuse This focus on compromise is crucial as new uses of medical technology expand the controversy beyond abortion and contraception to reach an increasing number of doctors and patients Lynch argues that doctor patient matching on the basis of personal moral values would eliminate or at least minimize many conflicts of conscience and suggests that state licensing boards facilitate this goal Licensing boards would be responsible for balancing the interests of doctors and patients by ensuring a sufficient number of willing physicians such that no physician's refusal leaves a patient entirely without access to desired medical services This proposed solution Lynch argues accommodates patients freedoms while leaving important room in the profession for individuals who find some of the capabilities of medical technology to be ethically objectionable

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