

Autonomy And Human Rights In Health Care An International Perspective International Library Of Ethics

Moral Blindness in Business Jacob Dahl Rendtorff.2020-08-03 In this book, Jacob Dahl Rendtorff investigates moral blindness in business and public administration based on Hannah Arendt's concept of banality of evil in her famous report on the Nazi-criminal Adolf Eichmann trial in Jerusalem in 1961. Moral blindness and evil in management is instrumental wrongdoing inflicted upon human beings as a violation of their dignity and humanity. Organizational evil in business, bureaucracies and public administration is analysed with focus on obedience to authority and systemic role conformity of managers and administrators. This includes the critical question about how concepts of banality of evil and moral blindness can explain ethical insensibility and lack of moral understanding in business and administration. Rendtorff proposes a humanistic vision of management and ethical leadership. Moral thinking, responsibility and moral judgment is essential in management and governance in business and administration. This book is a must-read for academics and practitioners studying and working in philosophy of management, business ethics, political philosophy, administration ethics and corporate social responsibility.

The Elderly Martin Lyon Levine.2017-05-15 Aging is a public health priority that is becoming

increasingly important in both developed and less developed nations, with individual health care providers and law-makers each facing difficult ethical and policy dilemmas. The complex issues physicians deal with include informed consent and patient decision-making capacity, use of advance care planning and decision-making by family and medical staff, and withdrawing and withholding life-sustaining interventions. Broader questions include: has aging been over medicalized? Is it ethical for older patients to receive less medical care than younger ones, through unspoken practice or formal rationing? Is there inevitable conflict between the generations over scarce medical resources? How should physician, patient and family confront end-of-life decisions? How have different nations responded to increasing numbers of the elderly? Have social values changed as to family responsibility and individual autonomy? This volume brings together the most significant published essays in the field.

New Perspectives on Paternalism and Health Care Thomas Schramme. 2016-10-09 This work sets the stage regarding debates about paternalism and health care for years to come. The anthology is organized around four parts: i) The concept of paternalism and theoretical issues regarding the idea of anti-paternalism, ii) strategies for justifying different forms of paternalism, iii) paternalism in psychiatry and psychotherapy, iv) paternalism and public health, and v) paternalism and reproductive medicine. Medical paternalism was arguably one of the main drivers of debates in medical ethics and has led to a wide acknowledgement of the value of patient autonomy. However, more recent developments in health care, such as the increasing significance of public health measures and the commercialization of medical services, have led to new social circumstances and hence to the need to rethink issues regarding paternalism. This work provides an invaluable source for many scholars and practitioners, since it deals in new and original ways with one of the main and

oldest issue in health care ethics.

Health Policy and Ethics Roger Worthington, Robert Rohrbaugh. 2017-11-22 This book demonstrates the utility of healthcare law, policy and professional standards in analysing the ethical issues that arise in the provision of health services. Rejecting moral absolutes, its examination of health law, policy and professional standards and of how societies codify beliefs is pertinent to ethical analysis - and also offers the possibility of practical solutions to healthcare challenges across the globe. Comparing and contrasting ethical and policy issues from countries around the world (with a focus on Asia, Europe and the USA), this book addresses such issues as conflict of interest, the balance between healthcare quality and cost, and the effect of geography and demographics on access to healthcare. Critique and discussion are tempered with suggestions for the evaluation of policy and systems; its pragmatic approach suggests how theory can and should inform practice. *Health Policy and Ethics* offers refreshing reading for professionals and academics in healthcare, medical ethics and policy. Researchers and students with an interest in healthcare delivery, comparative healthcare policy analysis, and health and human rights will also find much of interest. 'This book offers an unusual and welcome perspective on the rights and wrongs of health policy, with comparisons across space and time, from China to Italy, from Malaysia to India, from pharmaceutical ethics to the overarching topic of rationing.' Theodore Marmor, Professor Emeritus, Yale University School of Management Ethical analysis in this domain is not easy, as things held sacred will at times conflict, and of course within a given society there will be variability in values and priorities between individuals and over time. The anticipated difficulties serve to further emphasize that the ongoing ethical analysis should include input from those with experience and skill at that task. This book successfully demonstrates that point. In addition, *Health Policy and Ethics* is a welcome bridge

between these two fields, and a very worthwhile read for individuals whose primary interest lies in either one. - From the Foreword by Mark R. Mercurio

Health Rights Thomas Pogge.2017-05-15 Health Rights is a multidisciplinary collection of seminal papers examining ethical, legal, and empirical questions regarding the human right to health or health care. The volume discusses what obligations health rights entail for governments and other actors, how they relate to and potentially conflict with other rights and values, and how cultural diversity bears on the formulation and implementation of health rights. The paramount importance of such questions is illustrated, among other things, by the catastrophic health situation in developing countries and current debates about the TRIPS Agreement and health care reform in the United States. The volume is divided into five main parts which focus on philosophical questions about the bases for the right to health or health care; links between health and human rights; global bioethics and public health ethics; intellectual property rights in pharmaceuticals; and finally health rights issues arising in specific contexts such as HIV/AIDS, tuberculosis, and gender.

Autonomy and Human Rights in Health Care David N. Weisstub,Guillermo Díaz Pintos.2007-12-20 This book offers a group of essays published in memory of David Thomasma, one of the leading humanists in the field of bioethics during the twentieth century. The authors represent many different countries and disciplines throughout the globe. The volume deals with the pressing issue of how to ground a universal bioethics in the context of the conflicted world of combative cultures and perspectives.

Care in Healthcare Franziska Krause,Joachim Boldt.2017-10-24 This book is open access under a CC BY 4.0 license. This book examines the concept of care and care practices in healthcare from the interdisciplinary perspectives of continental philosophy, care ethics, the social sciences, and

anthropology. Areas addressed include dementia care, midwifery, diabetes care, psychiatry, and reproductive medicine. Special attention is paid to ambivalences and tensions within both the concept of care and care practices. Contributions in the first section of the book explore phenomenological and hermeneutic approaches to care and reveal historical precursors to care ethics. Empirical case studies and reflections on care in institutionalised and standardised settings form the second section of the book. The concluding chapter, jointly written by many of the contributors, points at recurring challenges of understanding and practicing care that open up the field for further research and discussion. This collection will be of great value to scholars and practitioners of medicine, ethics, philosophy, social science and history.

Informed Consent and Health Thierry Vanswevelt, Nicola Glover-Thomas. 2020-04-24 Informed consent is the legal instrument that purports to protect an individual's autonomy and defends against medical arbitrariness. This illuminating book investigates our evolving understanding of informed consent from a range of comparative and international perspectives, demonstrating the diversity of its interpretations around the world. Chapters offer a nuanced analysis of the problems that impede the understanding and implementation of the concept of informed consent and explore the contemporary challenges that continue to hinder both the patient and the medical community.

Healthcare as a Universal Human Right Rui Nunes. 2022-02-07 This important book outlines how, despite varying levels of global socio-economic development, governments around the world can guarantee their citizens' fundamental right to basic healthcare. Grounded in the philosophical position that healthcare is an essential element to human dignity, the book moves beyond this theoretical principle to offer policy-makers a basis for health policies based on public accountability and social responsiveness. Also emphasizing the importance of global co-operation, particularly in

the area of health promotion and communication, it addresses, too, the issue of financial sustainability, suggesting robust mechanisms of economic and social regulation. New opportunities created by e-health, evidence-based data and artificial intelligence are all highlighted and discussed, as is the issue of patient rights. Students and researchers across bioethics, public health and medical sociology will find this book fascinating reading, as will policy-makers in the field.

Healthcare as a Human Rights Issue Sabine Klotz, Heiner Bielefeldt, Martina

Schmidhuber, Andreas Frewer. 2017-11-30 This book deals with various facets of the human right to health: its normative profile as a universal right, current political and legal conflicts and contextualized implementation in different healthcare systems. The authors come from different countries and disciplines - law, political science, ethics, medicine etc. - and bring together a broad variety of academic and practical perspectives. The volume contains selected contributions of the international conference *The Right to Health - an Empty Promise?* held in September 2015 in Berlin and organized by the Emerging Field Initiative Project Human Rights in Healthcare (University of Erlangen-Nürnberg).

Bioethics in a European Perspective H.A. Ten Have, Bert Gordijn. 2013-03-09 This book gives an overview of the most salient themes in present-day bioethics. The book focuses on perspectives typical for the European context. This highlights not only particular bioethical themes such as social justice, choices in health care, and health policy (e.g., in post-communist countries), it also emphasizes specific approaches in ethical theory, in relation to Continental philosophies such as phenomenology and hermeneutics.

Bioethics and the Holocaust Stacy Gallin, Ira Bedzow. 2022-07-07 This open access book offers a framework for understanding how the Holocaust has shaped and continues to shape medical ethics,

health policy, and questions related to human rights around the world. The field of bioethics continues to face questions of social and medical controversy that have their roots in the lessons of the Holocaust, such as debates over beginning-of-life and medical genetics, end-of-life matters such as medical aid in dying, the development of ethical codes and regulations to guide human subject research, and human rights abuses in vulnerable populations. As the only example of medically sanctioned genocide in history, and one that used medicine and science to fundamentally undermine human dignity and the moral foundation of society, the Holocaust provides an invaluable framework for exploring current issues in bioethics and society today. This book, therefore, is of great value to all current and future ethicists, medical practitioners and policymakers – as well as laypeople.

Politics Between Nations Adebowale Akande. 2023-06-27 This multidisciplinary volume examines the meaning of global conflict and cooperation by international actors that can be caused by dis- or misinformation to people and discusses how to build diplomacy for peace and regional cooperation. The book further identifies boundaries of the relationships among the various governments of the world, transatlantic alliances, international organizations, non-governmental organizations, transnational corporations, and the overall interdependence of nations in the making of the modern world. Topics discussed in this volume include diplomacy, international relations theory, Eurasia politics, European Union, Brexit, Taliban taking over of Kabul government, and the ongoing Afghanistan conflict, terrorism, ISIS and Al Qaeda, international law, international organizations, interstate and intrastate war, threats and challenges, global civil society, religion, and culture. The volume advances contemporary theories and concepts to explain these issues concerning peoples and cultures in the complex world we live in. The book is a must-read for students, researchers, and scholars of international relations, political science, political history, political geography, economics,

and law in general, as well as diplomacy, political communication, and security studies in particular.
Human Rights, Dignity and Autonomy in Health Care and Social Services: Nordic Perspectives
Henriette Sinding Aasen.2010

Bioethical Decision Making and Argumentation Pedro Serna, José-Antonio Seoane.2016-09-26
This book clarifies the meaning of the most important and pervasive concepts and tools in bioethical argumentation (principles, values, dignity, rights, duties, deliberation, prudence) and assesses the methodological suitability of the main methods for clinical decision-making and argumentation. The first part of the book is devoted to the most developed or promising approaches regarding bioethical argumentation, namely those based on principles, values and human rights. The authors then continue to deal with the contributions and shortcomings of these approaches and suggest further developments by means of substantive and procedural elements and concepts from practical philosophy, normative systems theory, theory of action, human rights and legal argumentation. Furthermore, new models of biomedical and health care decision-making, which overcome the aforementioned criticism and stress the relevance of the argumentative responsibility, are included.
Autonomy, Consent and the Law Sheila A.M. McLean.2009-09-10
Autonomy is often said to be the dominant ethical principle in modern bioethics, and it is also important in law. Respect for autonomy is said to underpin the law of consent, which is theoretically designed to protect the right of patients to make decisions based on their own values and for their own reasons. The notion that consent underpins beneficent and lawful medical intervention is deeply rooted in the jurisprudence of countries throughout the world. However, Autonomy, Consent and the Law challenges the relationship between consent rules and autonomy, arguing that the very nature of the legal process inhibits its ability to respect autonomy, specifically in cases where patients argue that their ability to

act autonomously has been reduced or denied as a result of the withholding of information which they would have wanted to receive. Sheila McLean further argues that the bioethical debate about the true nature of autonomy - while rich and challenging - has had little if any impact on the law. Using the alleged distinction between the individualistic and the relational models of autonomy as a template, the author proposes that, while it might be assumed that the version ostensibly preferred by law - roughly equivalent to the individualistic model - would be transparently and consistently applied, in fact courts have vacillated between the two to achieve policy-based objectives. This is highlighted by examination of four specific areas of the law which most readily lend themselves to consideration of the application of the autonomy principle: namely refusal of life-sustaining treatment and assisted dying, maternal/foetal issues, genetics and transplantation. This book will be of great interest to scholars of medical law and bioethics.

Personhood and Health Care David C. Thomasma, David N. Weisstub, Christian Hervé. 2013-04-17
PERSONHOOD AND HEALTH CARE This book arose as a result of a pre-conference devoted to the topic held June 28, 1999 in Paris, France. The pre-conference preceded the Annual Congress of the International Academy of Law and Mental Health. Other chapters were solicited after the conference in order to more completely explore the relation of personhood to health care. The pre conference was held in honor of Yves Pelicier who led so many of our French colleagues in medicine, philosophy, and ethics as Christian Herve notes in his Tribute. As health care is aimed at healing persons, it is important to realize how difficult it is to construct a theory of personhood for health care, and thus, a theory of how healing in health care comes about or ought to occur. The book is divided into four parts, Concepts of the Person, Theories of Personhood in Relation to Health Care and Bioethics, Person and Identity, and Personhood and His Relations. Each section explores a

critical arena in constructing the relation of personhood to health care. Although no exploration of this nature can be exhaustive, every effort was made to present both conflicting and complementary views of personhood from within similar and different philosophical and religious traditions. PART ONE: CONCEPTS OF THE PERSON Tracing the origins of the concept of person from antiquity through present day, Jean Delemeau provides an historical sketch of the development of a wide range of meanings.

Ethics of Care Barnes, Marian, Brannelly, Tula, Lizzie Ward, Nicki Ward. 2015-10-28 Over the last twenty years, research on feminist care ethics has flourished, and this collection makes a unique contribution to that body of work. Drawing on a wealth of practical experience across eight different disciplinary fields, the international contributors demonstrate the significance of care ethics as a transformative way of thinking across diverse geographical, political, and interpersonal contexts. From an analysis of global responsibilities to a reimagining of care from the perspective of people with learning disabilities, each chapter highlights the necessity of thinking about the ethics of care within policies and practice.

Moral Equality, Bioethics, and the Child Claudia Wiesemann. 2016-08-18 Presenting real life cases from clinical practice, this book claims that children can be conceived of as moral equals without ignoring the fact that they still are children and in need of strong family relationships. Drawing upon recent advances in childhood studies and its key feature, the 'agentic child', it uncovers the ideology of adultism which has seeped into much what has been written about childhood ethics. However, this book also critically examines those positions that do accord moral equality to children but on grounds not strong enough to support their claim. It lays the groundwork for a theory of moral equality by assessing the concepts of parenthood, family, best interest,

paternalism, and, above all, autonomy and trust which are so important in envisioning what we owe the child. It does not only show how children – like adults – should be considered moral agents from infancy but also how ethical theories addressing adults can significantly profit from recognizing this. The analysis takes into account contributions from European as well as American scholars and makes use of a wide range of ethical, psychological, cultural, and social-scientific research.

Globalization, Human Rights and Populism Adebowale Akande. 2023-06-16 This book offers a comprehensive and multidisciplinary global overview of populism and human rights in the light of globalization. It examines why the dominant (neo)liberal paradigm of the last decades resulted in major economic and social inequalities which resulted in the surge of national populism, led by the election success of right-wing parties, movements, and leaders across the world. It discusses, among other topics, the success of Brexit in Britain and the election success of Donald Trump and Marine Le Pen and explains why there is a need for a dialogue on human rights and globalization in this era of populism. Further contributions analyze various important topics of the field, including cross-culturalism, globalization, human rights, challenges and threats, diversity, curbing global corruption, sustainable development, populism, the decline of free speech, the new nationalism, internationalization, global regime of human rights, leadership theory, global management competencies, gender, quality management, individualism-collectivism, and examples of new initiatives in global organizations. This makes the book a valuable and useful resource for students, researchers, and scholars of international relations, political science, sociology, political psychology, law, diplomatic studies, Communication and media studies, economics, education and management, as well as practitioners and policy-makers interested in a better understanding of globalization, populism, and human rights.

Autonomy and Clinical Medicine J. Bergsma, David C. Thomasma. 2013-03-14 This book arises from a two-fold conviction. The first is that autonomy, despite recent critiques about its importance in bioethics and philosophy of medicine, and the traditional resistance of medicine to its intrusion into the doctor-patient relation, is a fundamental building block of an individual's identity and mechanisms for dealing with illness, disease, and incapacity. As such it is an essential component in the health care professional's armamentarium employed to bring about healing. Furthermore, it functions in a similar way to assist the health professional in his or her relations to the sick and injured. The second conviction follows from the first. Autonomy is far more complex than appears from the philosophical use of the concept. In this conviction we join those who have criticized the over-reliance on autonomy in modern, secular bioethics originating in the United States, but gaining ascendancy in other cultures. This critique relies on appeals to the richer contexts of persons' lives. Elsewhere the contemporary critique of autonomy appears in a variety of alternative ethical models like narrative ethics, casuist ethics, and contextualism. Indeed, postmodern criticism of all bioethics argues that there is no defensible foundation for claims that one ought to respect autonomy or any other principle as a way of ensuring that one is ethical.

Human Rights, Dignity and Autonomy in Health Care and Social Services Henriette Sinding Aasen, Rune Halvorsen, António Barbosa da Silva. 2009 The overall theme of this volume is the understanding of human dignity, autonomy, and human rights in health care and social services in modern welfare states, with special reference to the Nordic countries. Focus is put on vulnerable groups such as children, individuals with cognitive impairment or mental illness, and persons with physical disabilities. Experts from different disciplines identify the ethical and legal dilemmas in modern welfare services and describe how basic values and/or rights come in conflict in concrete

situations. Of particular interest is how the human rights perspective challenges the policies and regulations of modern welfare states while at the same time providing the overall normative direction for solving ethical, legal, and social conflicts or shortcomings. Although the human rights perspective is the most dominant, insights from philosophy and the social sciences provide both a necessary and fruitful supplement to the legal approach. The volume will be of interest for academics, researchers, and students in the field of health care ethics, human rights, and welfare state policies. It presents a challenging outlook on dilemmas that are characteristic for the modern welfare state in general, and for the Nordic countries in particular, and it will give the reader important insights and references for further studies.

Advance Directives Peter Lack, Nikola Biller-Andorno, Susanne Brauer. 2013-10-21 This volume gives an overview on the currently debated ethical issues regarding advance directives from an international perspective. It focuses on a wider understanding of the known and widely accepted concept of patient self-determination for future situations. Although advance directives have been widely discussed since the 1980s, the ethical bases of advance directives still remain a matter of heated debates. The book aims to contribute to these controversial debates by integrating fundamental ethical issues on advance directives with practical matters of their implementation. Cultural, national and professional differences in how advance directives are understood by health care professions and by patients, as well as in laws and regulations, are pinpointed.

Advance Directives: Rethinking Regulation, Autonomy & Healthcare Decision-Making Hui Yun Chan. 2018-10-03 This book offers a new perspective on advance directives through a combined legal, ethical and philosophical inquiry. In addition to making a significant and novel theoretical contribution to the field, the book has an interdisciplinary and international appeal. The book will

help academics, healthcare professionals, legal practitioners and the educated reader to understand the challenges of creating and implementing advance directives, anticipate clinical realities, and preparing advance directives that reflect a higher degree of assurance in terms of implementation.

Routledge Handbook of Global Health Rights Clayton Ó Néill, Charles Foster, Jonathan Herring, John Tingle. 2021-05-30 This book examines the idea of a fundamental entitlement to health and healthcare from a human rights perspective. The volume is based on a particular conceptual reasoning that balances critical thinking and pragmatism in the context of a universal right to health. Thus, the primary focus of the book is the relationship or contrast between rights-based discourse/jurisprudential arguments and real-life healthcare contexts. The work sets out the constraints that are imposed on a universal right to health by practical realities such as economic hardship in countries, lack of appropriate governance, and lack of support for the implementation of this right through appropriate resource allocation. It queries the degree to which the existence of this legally enshrined right and its application in instruments such as the International Covenant on Economic, Social and Cultural Rights (ICESCR) and the Universal Declaration of Human Rights (UDHR) can be more than an ephemeral aspiration but can, actually, sustain, promote, and instil good practice. It further asks if social reality and the inequalities that present themselves therein impede the implementation of laudable human rights, particularly within marginalised communities and cadres of people. It deliberates on what states and global bodies do, or could do, in practical terms to ensure that such rights are moved beyond the aspirational and become attainable and implementable. Divided into three parts, the first analyses the notion of a universal inalienable right to health(care) from jurisprudential, anthropological, legal, and ethical perspectives. The second part considers the translation of international human rights norms into specific jurisdictional

healthcare contexts. With a global perspective it includes countries with very different legal, economic, and social contexts. Finally, the third part summarises the lessons learnt and provides a pathway for future action. The book will be an invaluable resource for students, academics, and policymakers working in the areas of health law and policy, and international human rights law. *Healthy Thoughts* Reidar Krummradt Lie.2002 This book, edited by a team of leading European bioethicists, is in all respects an innovative publication. As part of the core materials project of the European Ethics Network, this book collects European perspectives on health care ethics reflecting both the rich philosophical tradition and the broad interdisciplinary network in the field of European health care ethics. In the first part of the book on the physician-patient relationship, the authors present different views on the integration of patient autonomy in the relational structure of the medical profession. Here, the focus is on the reception of patient autonomy in the European context and on European alternatives for the radical understanding of patient autonomy. In the second part of this book, the contributions deal with the ethical problems of clinical research with patients and with the ethical values involved in human experimentation. In the third part of the book, the just organisation of the health care system and the consequent resource allocation are critically examined. With their common European background, the authors want to enrich and encourage the ongoing discussion on the physician-patient relationship, human experimentation and justice in health care from various perspectives and different points of departure. Therefore, this volume is suitable for the interested reader as well as for educational purposes.

The Right to Health Care: Ethical Considerations Eike-Henner W. Kluge.2022-01-13 This book provides an in-depth ethical analysis of the right to health care by contrasting privatized with socialized approaches. It pays special attention to how a socialized approach can be implemented in

the context of limited resources and offers a way of integrating allocation decisions at the policy level with institutional and hands-on decision-making. It also discusses how the right to health care translates into duties on part of the members of society. In an Appendix, it suggests how, in time of need, the TRIPS Agreement allows countries to side-step patent regulations that would otherwise raise the cost of patented healthcare products beyond what a particular society is able to afford. The book is of interest not only to scholars but also to healthcare policy makers, administrators and healthcare professionals, as well as to patients themselves.

Theories of the Self and Autonomy in Medical Ethics Michael Kühler, Veselin L.

Mitrović. 2020-09-29 This book engages in a critical discussion on how to respect and promote patients' autonomy in difficult cases such as palliative care and end-of-life decisions. These cases pose specific epistemic, normative, and practical problems, and the book elucidates the connection between the practical implications of the theoretical debate on respecting autonomy, on the one hand, and specific questions and challenges that arise in medical practice, on the other hand. Given that the idea of personal autonomy includes the notion of authenticity as one of its core components, the book explicitly includes discussions on underlying theories of the self. In doing so, it brings together original contributions and novel insights for "applied" scenarios based on interdisciplinary collaboration between German and Serbian scholars from philosophy, sociology, and law. It is of benefit to anyone cherishing autonomy in medical ethics and medical practice.

The Different Faces of Autonomy Maartje Schermer. 2002-11-30 Patient autonomy is a much discussed and debated subject in medical ethics, as well as in healthcare practice, medical law, and healthcare policy. This book provides a detailed and nuanced analysis of both the concept of autonomy and the principle of respect for autonomy, in an accessible style. The unique feature of

this book is that it combines empirical research into hospital practice with thorough philosophical analyses. As such, it is an example of a new movement in applied ethics, that of 'empirical ethics'. The key themes are informed consent and medical decision making, personal well-being, competence, paternalism and decision making for incompetent patients. Much attention is also devoted to autonomy in non-decision making situations - patient control over small everyday aspects of care, authenticity and existential aspects of illness, autonomy and the 'ethics of care', and the relationship between autonomy and trust in the physician-patient relationship. This book will be of interest to those working or studying in the field of medical ethics and applied ethics but also to healthcare professionals and health policy makers.

International Biolaw and Shared Ethical Principles Cinzia Caporale, Ilja Richard Pavone. 2018-01-12
The Universal Declaration on Bioethics and Human Rights, 2005, marked a significant step towards the recognition of universal standards in the field of science and medicine. This book provides an overview of the ethical and legal developments which have occurred in the field of bioethics and human rights since then. The work critically analyzes the Declaration from an ethical and legal perspective, commenting on its implementation, and discussing the role of non-binding norms in international bioethics. The authors examine whether the Declaration has contributed to the understanding of universal or global bioethics, and to what degree states have implemented the principles in their domestic legislation. The volume explores the currency of the Declaration vis-à-vis the more recent developments in technology and medicine and looks ahead to envisage the major bioethical challenges of the next twenty years. In this context, the book offers a comprehensive ethical and legal study of the Declaration with an in-depth analysis of the meaning of the provisions, in order to clarify the extension of human rights in the field of medicine and the obligations

incumbent upon UNESCO member States, with reference to their implementation practice.

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.

African Traditional Medicine: Autonomy and Informed Consent Peter Ikechukwu Osuji.2014-07-18 This book focuses on informed consent in African Traditional Medicine (ATM). ATM forms a large portion of the healthcare systems in Africa. WHO statistics show that as much as 80% of the population in Africa uses traditional medicine for primary health care. With such a large constituency, it follows that ATM and its practices should receive more attention in bioethics. By

comparing the ethics of care approach with the ATM approach to Relational Autonomy In Consent (RAIC), the authors argue that the ATM focus on consent based on consensus constitutes a legitimate informed consent. This book is distinctive insofar as it employs the ethics of care as a hermeneutic to interpret ATM. The analysis examines the ethics of care movement in Western bioethics to explore its relational approach to informed consent. Additionally, this is the first known study that discusses healthcare ethics committees in ATM.

Self-Determination, Dignity and End-of-Life Care Stefania Negri.2012-02-03 By providing an interdisciplinary reading of advance directives regulation in international, European and domestic law, this book offers new insights into the most controversial legal issues surrounding the debate over dignity and autonomy at the end of life.

Handbook of Racism, Xenophobia, and Populism Adebowale Akande.2022-12-08 This handbook presents the roots of symbolic racism as partly in both anti-black antagonism and non-racial conservative attitudes and values, representing a new form of racism independent of older racial and political attitudes. By doing so, it homes in on certain historical incidents and episodes and presents a cogent analysis of anti-black, Jim Crowism, anti-people of color (Black, Latino, Native Americans), and prejudice that exists in the United States and around the world as a central tenet of racism. The book exposes the reader to the nature and practice of stereotyping, negative bias, social categorization, modern forms of racism, immigration law empowerment, racialized incarceration, and police brutality in the American heartland. It states that several centuries of white Americans' negative socializing culture marked by widespread negative attitudes toward African Americans, are not eradicated and are still rife. Further, the book provides a panoramic view of trends of racial discrimination and other negative and desperate challenges that Black, Indigenous, and People of

Color face across the world. Finally, the volume examines xenophobia, racism, prejudice, and stereotyping in different contexts, including topics such as Covid-19, religion and racism, information manipulation, and populism. The book, therefore, is a must-read for students, researchers, and scholars of political science, psychology, history, sociology, communications/media studies, diplomatic studies, and law in general, as well as ethnic and racial studies, American politics, global affairs, populism, and discrimination in particular.

Global Bioethics and Human Rights Wanda Teays, Alison Dundes Renteln. 2020 The ethical issues we face in healthcare, justice, and human rights extend beyond national boundaries--they are global and cross-cultural in scope. The second edition of this interdisciplinary and international collection features new essays on gender identity, vaccines, stem cells, bioterror, and other pressing contemporary concerns.

Public Health Policy and Ethics Michael Boylan. 2006-05-17 Public Health Policy and Ethics brings together philosophers and practitioners to address the foundations and principles upon which public health policy may be advanced. What is the basis that justifies public health in the first place? Why should individuals be disadvantaged for the sake of the group? How do policy concerns and clinical practice work together and work against each other? Can the boundaries of public health be extended to include social ills that are amenable to group-dynamic solutions? These are some of the crucial questions that form the core of this volume of original essays sure to cause practitioners to engage in a critical re-evaluation of the role of ethics in public health policy. This volume is unique because of its philosophical approach. It develops a theoretical basis for public health and then examines cutting-edge issues of practice that include social and political issues of public health. In this way the book extends the usual purview of public health. Public Health Policy and Ethics is of

interest to those working in public health policy, ethics and social philosophy. It may be used as a textbook for courses on public health policy and ethics, medical ethics, social philosophy and applied or public philosophy.

Doctors and Torture Wanda Teays.2019-07-22 This book brings into sharp relief the extent to which the medical profession has enabled or participated in actions that are at moral crossroads. Physical and psychological abuse and violations of medical codes have already been brought to light by concerned bioethicists responding to ethical lapses of the “war on terror.” This book goes to the next level by looking at three areas that also merit our attention and call us to speak out against abuses. These are (1) dehumanization (such as forced nudity, hooding, sensory deprivation, exploitation of phobias, waterboarding, and environmental manipulation), (2) non-consensual forced-feeding, and (3) solitary confinement. Each area raises important questions for the medical profession. Author Wanda Teays calls upon doctors and nurses to reflect on the role they play in the unethical treatment of prisoners and detainees by crossing moral boundaries around each of these areas. In the process, we are reminded that bioethics is global, not local — and the concerns of the discipline encompass issues with a wider scope.

Global Health and Human Rights John Harrington, Maria Stuttaford.2010-06-10 The right to health, having been previously neglected is now being deployed more and more often in litigation, activism and policy-making across the world. International bodies such as the WHO, UNAIDS, World Bank and WTO are increasingly using or being evaluated with reference to health rights, and international NGOs frequently use the language of rights in campaigning and in more concrete litigation. This book brings together an impressive array of internationally renowned scholars in the areas of law, philosophy and health policy to critically interrogate the development of rights based

approaches to health. The volume integrates discussion of the right to health at a theoretical level in law and ethics, with the difficult substantive issues where the right is relevant, and with emerging systems of global health governance. The contributions to this volume will add to our theoretical and practical understanding of rights based approaches to health.

International Public Health Policy and Ethics Michael Boylan. 2008-07-20 Public Health Policy and Ethics brings together philosophers and practitioners to address the foundations and principles upon which public health policy may be advanced – especially in the international arena. What is the basis that justifies public health in the first place? Why should individuals be disadvantaged for the sake of the group? How do policy concerns and clinical practice work together and work against each other? Can the boundaries of public health be extended to include social ills that are amenable to group-dynamic solutions? What about political issues? How can international finance make an impact? These are some of the crucial questions that form the core of this volume of original essays sure to cause practitioners to engage in a critical re-evaluation of the role of ethics in public health policy.

International Health Law and Ethics André den Exter. 2009 This book contains a collection of treaty documents and soft law on health care rights and health ethics used in health law training programs. Regional documents and explanatory reports on health care rights, which are derived from international human rights law, provide a way of unwrapping government obligations in health care, making rights more specific, accessible, and (judicially) accountable. In addition, soft law declarations and medical ethics contribute to understanding the moral meaning of human rights in health care. As such, the principles and standards provide practical guidance for States when dealing with equal access to health care services, the rights of patients, biomedical research, organ

donation and transplantation, genetics, and public health. The book's general comments and explanatory reports amplify the principles embodied in human rights treaties. The authoritative interpretations clarify a 'European approach' on a State's obligations concerning health care rights and ethics. This volume is an initiative of the Erasmus Observatory on Health Law. It will be a helpful guide for all trainers, health care professionals, and students interested in human rights issues in health care.

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