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Prenatal Testing And Disability Rights

Kenneth Goodman

Prenatal Testing And Disability Rights:

Prenatal Testing and Disability Rights Erik Parens, Adrienne Asch, 2000-09-28 As prenatal tests proliferate the medical and broader communities perceive that such testing is a logical extension of good prenatal care it helps parents have healthy babies But prenatal tests have been criticized by the disability rights community which contends that advances in science should be directed at improving their lives not preventing them Used primarily to decide to abort a fetus that would have been born with mental or physical impairments prenatal tests arguably reinforce discrimination against and misconceptions about people with disabilities In these essays people on both sides of the issue engage in an honest and occasionally painful debate about prenatal testing and selective abortion. The contributors include both people who live with and people who theorize about disabilities scholars from the social sciences and humanities medical geneticists genetic counselors physicians and lawyers Although the essayists don't arrive at a consensus over the disability community s objections to prenatal testing and its consequences they do offer recommendations for ameliorating some of the problems Prenatal Testing and Disability Rights Erik Parens, Adrienne Asch, 2000 In these essays associated with the practice health care professionals scholars and members of the disability community debate the implications of prenatal testing for people with disabilitties and for parent child relationships generally Cover Prenatal Testing and Disability Rights Erik Parens, Adrienne Asch, 2009 Prenatal Genetic Testing, Abortion, and Disability Justice Amber Knight, Joshua Miller, 2023 Prenatal Genetic Testing Abortion and Disability Justice explores the intent and effects of prenatal screening in connection to women's bodily autonomy and disability rights addressing themes at the intersection of genetic medicine policymaking critical disabilities studies and political theory The Eugenic Mind Project Robert A. Wilson, 2021-02-02 An examination of eugenic thinking past and present from forced sterilization to prenatal screening drawing on experience with those who survived eugenics Part science and part social movement eugenics emerged in the late nineteenth century as a tool for human improvement In response to perceived threats of criminality moral degeneration feeble mindedness and the rising tide of color eugenic laws and social policies aimed to better the human race by regulating reproductive choice through science and technology In this book Rob Wilson examines eugenic thought and practice from forced sterilization to prenatal screening drawing on his experience working with eugenics survivors Using the social sciences standpoint theory as a framework to understand the intersection of eugenics disability social inclusiveness and human variation Wilson focuses on those who have lived through a eugenic past and those confronted by the legacy of eugenic thinking today By doing so he brings eugenics from the distant past to the ongoing present Wilson discusses such topics as the conceptualization of eugenic traits the formulation of laws regulating immigration and marriage and requiring sexual sterilization the depiction of the targets of eugenics as subhuman the systematic construction of a concept of normality the eugenic logic in prenatal screening and contemporary bioethics and the incorporation of eugenics and disability into standpoint theory Law and

the Contradictions of the Disability Rights Movement Samuel R. Bagenstos, 2009-06-23 The passage of the Americans with Disabilities Act in 1990 was hailed as revolutionary legislation but in the ensuing years restrictive Supreme Court decisions have prompted accusations that the Court has betrayed the disability rights movement The ADA can lay claim to notable successes yet people with disabilities continue to be unemployed at extremely high rates In this timely book Samuel R Bagenstos examines the history of the movement and discusses the various often conflicting projects of diverse participants He argues that while the courts deserve some criticism some may also be fairly aimed at the choices made by prominent disability rights activists as they crafted and argued for the ADA The author concludes with an assessment of the limits of antidiscrimination law in integrating and empowering people with disabilities and he suggests new policy directions to make these goals a reality **Disability Rights and Wrongs** Tom Shakespeare, 2006-12-05 Over the last thirty years the field of disability studies has emerged from the political activism of disabled people In this challenging review of the field leading disability academic and activist Tom Shakespeare argues that the social model theory has reached a dead end Drawing on a critical realist perspective Shakespeare promotes a pluralist engaged and nuanced approach to disability Key topics discussed include dichotomies the dangerous polarizations of medical model versus social model impairment versus disability and disabled people versus non disabled people identity the drawbacks of the disability movement s emphasis on identity politics bioethics in disability choices at the beginning and end of life and in the field of genetic and stem cell therapies care and social relationships questions of intimacy and friendship This stimulating and accessible book challenges orthodoxies in British disability studies promoting a new conceptualization of disability and fresh research agenda It is an invaluable resource for researchers and students in disability studies and sociology as well as professionals policy makers and activists

Disability, Health, Law, and Bioethics I. Glenn Cohen, Carmel Shachar, Anita Silvers, Michael Ashley Stein, 2020-04-23 Historically and across societies people with disabilities have been stigmatized and excluded from social opportunities on a variety of culturally specific grounds In this collection the authors explore the impact that the philosophical framing of disability can have on public policy questions in the clinic in the courtroom and elsewhere They examine the implications of this understanding for legal and policy approaches to disability strategies for allocating and accessing health care the implementation of the Americans with Disabilities Act health care rights and other legal tools designed to address discrimination This volume should be read by anyone seeking a balanced view of disability and an understanding of the connection between the framing of disability and policies that have a real world impact on individuals Studies in Law, Politics, and Society Austin Sarat, 2018-06-29 Studies in Law Politics and Society provides a vehicle for the publication of scholarly articles in interdisciplinary legal scholarship This volume features a special section with papers dedicated to law and disability The chapters examine issues of HIV obesity disability rights assisted suicide and prenatal testing Life Histories of Genetic Disease Andrew J. Hogan, 2016-11 A history of genetic testing warns that such tests may tell us more

than we want to know Medical geneticists began mapping the chromosomal infrastructure piece by piece in the 1970s by focusing on what was known about individual genetic disorders Five decades later their infrastructure had become an edifice for prevention allowing today s expecting parents to choose to test prenatally for hundreds of disease specific mutations using powerful genetic testing platforms In Life Histories of Genetic Disease Andrew J Hogan explores how various diseases were made genetic after 1960 with the long term aim of treating and curing them using gene therapy In the process he explains these disorders were located in the human genome and became targets for prenatal prevention while the ongoing promise of gene therapy remained on the distant horizon In narrating the history of research that contributed to diagnostic genetic medicine Hogan describes the expanding scope of prenatal diagnosis and prevention He draws on case studies of Prader Willi fragile X DiGeorge and velo cardio facial syndromes to illustrate that almost all testing in medical genetics is inseparable from the larger and increasingly big data oriented aims of biomedical research Hogan also reveals how contemporary genetic testing infrastructure reflects an intense collaboration among cytogeneticists molecular biologists and doctors specializing in human malformation Hogan critiques the modern ideology of genetic prevention which suggests that all pregnancies are at risk for genetic disease and should be subject to extensive genomic screening He examines the dilemmas and ethics of the use of prenatal diagnostic information in an era when medical geneticists and biotechnology companies have begun offering whole genome prenatal screening essentially searching for any disease causing mutation Hogan's focus and analysis is animated by ongoing scientific and scholarly debates about the extent to which the preventive focus in contemporary medical genetics resembles the aims of earlier eugenicists Written for historians sociologists and anthropologists of science and medicine as well as bioethics scholars physicians geneticists and families affected by genetic conditions Life Histories of Genetic Disease is a profound exploration of the scientific culture surrounding malformation and Disability Dialogues Andrew J. Hogan, 2022-11-29 A historical look at how activists influenced the adoption of mutation more positive inclusive and sociopolitical views of disability Disability activism has fundamentally changed American society for the better and along with it the views and practices of many clinical professionals After 1945 disability self advocates and family advocates pushed for the inclusion of more positive inclusive and sociopolitical perspectives on disability in clinical research training and practice In Disability Dialogues Andrew J Hogan highlights the contributions of disabled people along with their family members and other allies in changing clinical understandings and approaches to disability Hogan examines the evolving medical social and political engagement of three postwar professions clinical psychology pediatrics and genetic counseling with disability and disability related advocacy Professionals in these fields historically resisted adopting a more inclusive and accepting perspective on people with disabilities primarily due to concerns about professional role identity and prestige In response to the work of disability activists however these attitudes gradually began to change Disability Dialogues provides an important contribution to historical sociological and bioethical accounts of disability and clinical

professionalization Moving beyond advocacy alone Hogan makes the case for why present day clinical professional fields need to better recruit and support disabled practitioners Disabled clinicians are uniquely positioned to combine biomedical expertise with their lived experiences of disability and encourage greater tolerance for disabilities among their colleagues The Dream of the Perfect Child Joan Rothschild, 2005 A feminist critique of bioethics and students and institutions attitudes toward reproductive technologies Genetic Governance Robin Bunton, Alan R. Petersen, 2005 Contributors look at how the new genetics healthcare and arising health policy can or should be governed and regulated rather than focusing on individualistic ethical issues this text takes a broader international public health approach The Case of Terri Schiavo Kenneth Goodman, 2009-10-29 The case of Terri Schiavo a young woman who spent 15 years in a persistent vegetative state has emerged as a watershed in debates over end of life care While many observers had thought the right to refuse medical treatment was well established this case split a family divided a nation and counfounded physicians legislators and many of the people they treated or represented In renewing debates over the importance of advance directives the appropriate role of artificial hydration and nutrition and the responsibilities of family members the case also became one of history s most extensively litigated health care disputes The Case of Terri Schiavo assembles a team of first hand participants and content experts to provide thoughtful and nuanced analyses In addition to a comprehensive overview the book includes contributions by Ms Schiavo s guardian ad litem a neurologist and lawyer who participated in the case and scholars who examine issues related to litigation faith gender and disability The volume also includes a powerful dissent from the views of many scholars in the bioethics community The book is intended for students health care professionals policy makers and other in search of carefully reasoned analyses of the case that will shape our view of death and end of life medical care for decades

Encyclopedia of Disability Gary L Albrecht,2006 Presents current knowledge of and experience with disability across a wide variety of places conditions and cultures to both the general reader and the specialist In Search of Common Ground on Abortion Robin West, Justin Murray, Meredith Esser, 2016-05-23 This book brings together academics legal practitioners and activists with a wide range of pro choice pro life and other views to explore the possibilities for cultural philosophical moral and political common ground on the subjects of abortion and reproductive justice more generally It aims to rethink polarized positions on sexuality morality religion and law in relation to abortion as a way of laying the groundwork for productive and collaborative dialogue Edited by a leading figure on gender issues and emerging voices in the quest for reproductive justice a broad concept that encompasses the interests of men women and children alike the contributions both search for common ground between opposing positions in our struggles around abortion and seek to bring balance to these contentious debates The book will be valuable to anyone interested in law and society gender and religious studies and philosophy and theory of law Routledge Handbook of Disability Law and Human Rights Peter Blanck, Eilionóir Flynn, 2016-07-07 This handbook provides a comprehensive and authoritative state of the art review of the current and

emerging research and policy on disability law Bringing together a team of respected and experienced experts the handbook offers a range of jurisdictional and multidisciplinary perspectives The authors consider historical and contemporary as well as comparative perspectives of disability law Divided into three parts the contributors provide a comprehensive reference to the theoretical underpinnings ongoing debates and emerging fields within the subject The study provides a strong basis for consideration of contemporary disability law its research foundations and progressive developments in the area The book incorporates interdisciplinary and comparative country perspectives to capture the breadth of current discourse on disability law This handbook provides a valuable resource for a wide range of scholars public and private researchers NGOs and practitioners working in the area of disability law and across national and transnational disability schemes The work will be of important interest to those in the fields of sociology history psychology economics political science rehabilitation sciences medicine technology and law among others The Routledge Handbook of Dehumanization Maria Kronfeldner, 2021-02-25 A striking feature of atrocities as seen in genocides civil wars or violence against certain racial and ethnic groups is the attempt to dehumanize to deny and strip human beings of their humanity Yet the very nature of dehumanization remains relatively poorly understood The Routledge Handbook of Dehumanization is the first comprehensive and multidisciplinary reference source on the subject and an outstanding survey of the key concepts issues and debates within dehumanization studies Organized into four parts the Handbook covers the following topics The history of dehumanization from Greek Antiquity to the 20th century contextualizing the oscillating boundaries dimensions and hierarchies of humanity in the history of the West How dehumanization is contemporarily studied with respect to special contexts as part of social psychology as part of legal studies or literary studies and how it connects to the idea of human rights disability and eugenics the question of animals and the issue of moral standing How to tackle its complex facets with respect to the perpetrator s and the target s perspective metadehumanization and selfdehumanization rehumanization social death status and interdependence as well as the fear we show toward robots that become too human for us Conceptual and epistemological questions on how to distinguish different forms of dehumanization and neighboring phenomena on why dehumanization appears so paradoxical and on its connection to hatred essentialism and perception Essential reading for students and researchers in philosophy history psychology and anthropology this Handbook will also be of interest to those in related disciplines such as politics international relations criminology legal studies literary studies gender studies disability studies or race and ethnic studies as well as readers from social work political activism and public policy Disability Rights and Wrongs Revisited Tom Shakespeare, 2013-10-30 Over the last forty years the field of disability studies has emerged from the political activism of disabled people In this challenging review of the field leading disability academic and activist Tom Shakespeare argues that disability research needs a firmer conceptual and empirical footing This new edition is updated throughout reflecting Shakespeare s most recent thinking drawing on current research and responding to controversies

surrounding the first edition and the World Report on Disability as well as incorporating new chapters on cultural disability studies personal assistance sexuality and violence Using a critical realist approach Disability Rights and Wrongs Revisited promotes a pluralist engaged and nuanced approach to disability Key topics discussed include dichotomies going beyond dangerous polarizations such as medical model versus social model to achieve a complex multi factorial account of disability identity the drawbacks of the disability movement s emphasis on identity politics bioethics choices at the beginning and end of life and in the field of genetic and stem cell therapies relationships feminist and virtue ethics approaches to questions of intimacy assistance and friendship This stimulating and accessible book challenges disability studies orthodoxy promoting a new conceptualization of disability and fresh research agenda It is an invaluable resource for researchers and students in disability studies and sociology as well as professionals policy makers and activists **Disability and Equality Law** ElizabethF. Emens, 2017-07-05 This interdisciplinary collection of essays addresses the theoretical practical and legal dimensions of equality for persons with disabilities The issues covered include the central problem of defining disability and impairment the dilemma of same versus different treatment the balance between autonomy and external influence and support linkages to other anti discrimination categories such as race and sex the place of disability theory within identity politics and issues of life death and our most intimate relationships The articles reflect a wealth of international viewpoints and interdisciplinary areas which include philosophy economics memoirs cultural studies empirical studies and legal scholarship The selection also includes classic texts which set out foundational ideas such as the social model of disability or the goal of integration alongside essays that critique these conceptual mainstays This volume brings into sharp focus a wide range of contentious and complex issues in the field of disability studies and is of interest to researchers and students from a wide range of fields

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