Sterilization Of People With Mental Disabilities Issues Perspectives And Cases

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Rights of the Disabled - David M. Haugen 2008 Provides an overview, chronology of events, glossary and annotated bibliography for disability rights in the United States.

Trauma and the Struggle to Open Up: From Avoidance to Recovery and Growth - Robert T. Muller 2018-06-19

How to navigate the therapeutic relationship with trauma survivors, to help bring recovery and growth. In therapy, we see how relationships are central to many traumatic experiences, but relationships are also critical to trauma recovery. Grounded firmly in attachment and trauma theory, this book shows how to use the psychotherapy relationship, to help clients find self-understanding and healing from trauma. Offering candid, personal guidance, using rich case examples, Dr. Robert T. Muller provides the steps needed to build and maintain a strong therapist-client relationship –one that helps bring recovery and growth. With a host of practical tips and protocols, this book gives therapists a roadmap to effective trauma treatment.

Under the Influence - Preston Peet 2004-01-01

This myth-busting anthology cuts through the propaganda to tell the true story of drug use, abuse, and the costly war on friends, families and communities. Author and regular High Times contributor Preston Peet assembles an all-star cast of writers to shine a harsh light on the misinformation peddled by prohibitionists who profit from the War on Some Drugs and Users. Despite the anti-drug hysteria, drugs have been an integral aspect of human life for thousands of years. They cure diseases, ease pain, enhance intelligence, calm nerves, open the doors of perception and alter consciousness. Yet, even with the easing of marijuana restrictions, the War on Some Drugs and Users continues to persecute huge swaths of the population. The reasons why can be found in Under the Influence. The decades and trillions of dollars spent waging war on neighbors, friends and families have done nothing to eradicate drug use and abuse, but they have succeeded in overthrowing governments, tearing apart families and communities, and ensured the rise of international criminal cartels. Under the Influence explains how we came to this state of affairs and how we can bring about real reform. Contributors include Tom Robbins, Paul Krassner, Rick Doblin, Mike Gray, Lonny Shavelson, Daniel Forbes, Steve Wishnia, Cynthia Cotts, Russ Kick, Dr. Stanislav Grof, Daniel Pinchbeck, Paul Armentano, Jacob Sullum, Peter Dale Scott and Robert Anton Wilson.

Research Anthology on Physical and Intellectual Disabilities in an Inclusive Society - Management Association, Information Resources 2021-08-27

Discussions surrounding inclusivity have grown exponentially in recent years. In today's world where diversity, equity, and inclusion are the hot topics in all aspects of society, it is more important than ever to define what it means to be an inclusive society, as well as challenges and potential growth. Those with physical and intellectual disabilities, including vision and hearing impairment, Down syndrome, locomotor disability, and more continue to face challenges of accessibility in their daily lives, especially when facing an increasingly digitalized society. It is crucial that research is brought up to date on the latest assistive technologies, educational practices, work assistance, and online support that can be provided to those classified with a disability. The Research Anthology on Physical and Intellectual Disabilities in an Inclusive Society provides a comprehensive guide of a range of topics relating to myriad aspects, difficulties, and opportunities of becoming a more inclusive society toward those with physical or intellectual disabilities. Covering everything from disabilities in education, sports, marriages, and more, it is essential for psychologists, psychiatrists, pediatricians, psychiatric nurses, clinicians, special education teachers, social workers, hospital administrators, mental health specialists, managers, academicians,

rehabilitation centers, researchers, and students who wish to learn more about what it means to be an inclusive society and best practices in order to get there.

Disabled Persons and the Law - Bruce D. Sales 2013-06-29

<u>Mental Disability, Violence, and Future Dangerousness</u> - John Weston Parry 2013-09-26

Using a multi-disciplinary approach, this book documents and explains how, when and why adults and children with mental disabilities—including those with sexual disorders— who are perceived to be a future danger to others, the community, or themselves have become the most stigmatized, abused, and mistreated group in America, and what should be done to correct the resulting injustices. The author identifies and analyzes the key factors that should be understood when lawyers, judges, mental health professionals, policymakers, legislators, advocates, forensic experts, professors and their students consider the legal, treatment and policy decisions that affect this highly stigmatized group of people.

Eugenics and the Welfare State - Gunnar Broberg 2005 In 1997 Eugenics and the Welfare State caused an uproar with international repercussions. This edition contains a new introduction by Broberg and Roll-Hansen, addressing events that occurred following the original publication. The four essays in this book stand as a chilling indictment of mass sterilization practices, not only in Scandinavia but in other European countries and the United States--eugenics practices that remained largely hidden from the public view until recently. Eugenics and the Welfare State also provides an in-depth, critical examination of the history, politics, science, and economics that led to mass sterilization programs in Norway, Sweden, Denmark, and Finland; programs put in place for the "betterment of society" and based largely on the "junk science" of eugenics that was popular before the rise of Nazism in Germany. When the results of Broberg's and Roll-Hansen's book were widely publicized in August 1997, the London Observer reported, "Yesterday Margot Wallstrom, the Swedish Minister for Social Policy, issued a belated reaction to the revelations. She said: 'What went on is barbaric and a national disgrace.' She pledged to create a law ensuring that involuntary sterilisation would never again be used in Sweden, and promised compensation to victims." Ultimately, the Swedish government not only apologized to the many thousands who had been sterilized without their knowledge or against their will, but also put in place a program for the payment of reparations to these unfortunate victims.

Fit to Be Tied - Rebecca M. Kluchin 2011

The 1960s revolutionized American contraceptive practice. Diaphragms, jellies, and condoms with high failure rates gave way to newer choices of the Pill, IUD, and sterilization. Fit to Be Tied provides a history of sterilization and what would prove to become, at once, socially divisive and a popular form of birth control. During the first half of the twentieth century, sterilization (tubal ligation and vasectomy) was a tool of eugenics. Individuals who endorsed crude notions of biological determinism sought to control the reproductive decisions of women they considered "unfit" by nature of race or class, and used surgery to do so. Incorporating first-person narratives, court cases, and official records, Rebecca M. Kluchin examines the evolution of forced sterilization of poor women, especially women of color, in the second half of the century and contrasts it with demands for contraceptive sterilization made by white women and men. She chronicles public acceptance during an era of reproductive and sexual freedom, and the subsequent replacement of the eugenics movement with "neo-eugenic" standards that continued to influence American medical practice, family planning, public policy, and popular sentiment.

The Family in America -

Encyclopedia of Women in Today's World - Mary Zeiss Stange 2011-02-23

This work includes 1000 entries covering the spectrum of defining women in the contemporary world.

The Convention on the Rights of Persons with Disabilities - Ilias Bantekas 2018-09-20

This treatise is a detailed article-by-article examination of the United Nations Convention on the Rights of Persons with Disabilities (CRPD). Each article of the CRPD contains a methodical analysis of the preparatory works, followed by an exhaustive examination of the contents of each articlebased on case law and concluding observations from the CRPD Committee, judgments from national and international courts and tribunals, pertinent UN and other reports, the key literature on the article under review. The volume features commentary from a broad range of scholars across a variety of disciplines in order to provide a comprehensive study of the legal, psychological, education, sociological, and other aspects of the CPRD. This encyclopaedic commentary on the CRPD effectively covers all the issuesarising from international disability law and practice, and will be an ideal resource for all working in the field.

Sterilization of People with Mental Disabilities - Ellen A. Brantlinger 1995

An examination of the medical and legal trends in sterilization with an emphasis on people with disabilities.

Practical Ethics for General Practice - Wendy A Rogers 2009 "Practical Ethics for General Practice, second edition, is essential reading for GPs, trainees, community nurses, those interested in bioethics, and medical students." --Book Jacket.

Keeping America Sane - Ian Robert Dowbiggin 2018-10-18 What would bring a physician to conclude that sterilization is appropriate treatment for the mentally ill and mentally handicapped? Using archival sources, Ian Robert Dowbiggin documents the involvement of both American and Canadian psychiatrists in the eugenics movement of the early twentieth century. He explains why professional men and women committed to helping those less fortunate than themselves arrived at such morally and intellectually dubious conclusions. Psychiatrists at the end of the nineteenth century felt professionally vulnerable, Dowbiggin explains, because they were under intense pressure from state and provincial governments and from other physicians to reform their specialty. Eugenic ideas, which dominated public health policy making, seemed the best vehicle for catching up with the progress of science. Among the prominent psychiatrist-eugenicists Dowbiggin considers are G. Alder Blumer, Charles Kirk Clarke, Thomas Salmon, Clare Hincks, and William Partlow. Tracing psychiatric support for eugenics throughout the interwar years, Dowbiggin pays special attention to the role of psychiatrists in the fierce debates about immigration policy. His examination of psychiatry's unfortunate flirtation with eugenics elucidates how professional groups come to think and act along common lines within specific historical contexts.

The Human Rights of Persons with Intellectual Disabilities - Stanley S. Herr 2003

Inequality: Marcia H. Rioux

Sociopolitical Aspects of Disabilities - Willie V. Bryan 2010 The social and political history of disabilities reveals some of the historical roots that anchor some of our current beliefs, attitudes and perceptions of disabilities and persons who possess disabilities. An understanding of the social and political history of disabilities in the United States is important for rehabilitation professionals and other helping professionals who work with persons with disabilities not only to understand how history affects our current attitudes and behavior but also to provide a perspective on how current events and actions that have produced the present state of affairs for persons with disabilities. This new edition continues the discussion of the evolution of societal attitudes toward persons with disabilities and explains how social attitudes impact social interactions which in turn create a need for political action to correct injustices and/or increase opportunities for persons with disabilities. Divided into two parts, the first part begins with an introduction to the relationship between social issues and political actions and continues with discussions on the foundation of beliefs and treatment of persons with disabilities, the oppressed, social environment, the advocates, the disability rights movements, and family roles. The second part explores the political history of disabilities, the Developmental Disabilities Act, the Americans with Disabilities Act

Amendments Act of 2008, as well as the inclusion of acts concerning professional training and continuing education, vocational rehabilitation, medical rehabilitation, economic assistance, deinstitutionalization and independent living, civil rights and advocacy, and the new eugenics. Review/discussion questions are included at the end of each chapter along with a Suggested Readings section. It will be useful as a primary or secondary text in vocational rehabilitation training programs, social work programs, nursing programs, occupational and physical therapy assistant programs, and human services programs.

Pediatric Bioethics - Geoffrey Miller 2010

This volume offers a theoretical and practical overview of the ethics of pediatric medicine. It serves as a fundamental handbook and resource for pediatricians, nurses, residents in training, graduate students, and practitioners of ethics and healthcare policy. Written by a team of leading experts, Pediatric Bioethics addresses those difficult ethical questions concerning the clinical and academic practice of pediatrics, including an approach to recognizing boundaries when confronted with issues such as end of life care, life-sustaining treatment, extreme prematurity, pharmacotherapy, and research. Thorny topics such as what constitutes best interests, personhood, or distributive justice and public health concerns such as immunization and newborn genetic screening are also addressed.

Imbeciles - Adam Cohen 2017-03-07

Longlisted for the 2016 National Book Award for Nonfiction One of America's great miscarriages of justice, the Supreme Court's infamous 1927 Buck v. Bell ruling made government sterilization of "undesirable" citizens the law of the land In 1927, the Supreme Court handed down a ruling so disturbing, ignorant, and cruel that it stands as one of the great injustices in American history. In Imbeciles, bestselling author Adam Cohen exposes the court's decision to allow the sterilization of a young woman it wrongly thought to be "feebleminded" and to champion the mass eugenic sterilization of undesirable citizens for the greater good of the country. The 8-1 ruling was signed by some of the most revered figures in American law—including Chief Justice William Howard Taft, a former U.S. president; and Louis Brandeis, a progressive icon. Oliver Wendell Holmes, considered by many the greatest Supreme Court justice in history, wrote the majority opinion, including the court's famous declaration "Three generations of imbeciles are enough." Imbeciles is the shocking story of Buck v. Bell, a legal case that challenges our faith in American justice. A gripping courtroom drama, it pits a helpless young woman against powerful scientists, lawyers, and judges who believed that eugenic measures were necessary to save the nation from being "swamped with incompetence." At the center was Carrie Buck, who was born into a poor family in Charlottesville, Virginia, and taken in by a foster family, until she became pregnant out of wedlock. She was then declared "feebleminded" and shipped off to the Colony for Epileptics and Feeble-Minded. Buck v. Bell unfolded against the backdrop of a nation in the thrall of eugenics, which many Americans thought would uplift the human race. Congress embraced this fervor, enacting the first laws designed to prevent immigration by Italians, Jews, and other groups charged with being genetically inferior. Cohen shows how Buck arrived at the colony at just the wrong time, when influential scientists and politicians were looking for a "test case" to determine whether Virginia's new eugenic sterilization law could withstand a legal challenge. A cabal of powerful men lined up against her, and no one stood up for her—not even her lawyer, who, it is now clear, was in collusion with the men who wanted her sterilized. In the end, Buck's case was heard by the Supreme Court, the institution established by the founders to ensure that justice would prevail. The court could have seen through the false claim that Buck was a threat to the gene pool, or it could have found that forced sterilization was a violation of her rights. Instead, Holmes, a scion of several prominent Boston Brahmin families, who was raised to believe in the superiority of his own bloodlines, wrote a vicious, haunting decision upholding Buck's sterilization and imploring the nation to sterilize many more. Holmes got his wish, and before the madness ended some sixty to seventy thousand Americans were sterilized. Cohen overturns cherished myths and demolishes lauded figures in relentless pursuit of the truth. With the intellectual force of a legal brief and the passion of a front-page exposé, Imbeciles is an ardent indictment of our champions of justice and our optimistic faith in progress, as well as a triumph of American legal and social history.

Contraceptive Technology - Robert Anthony Hatcher 2007 The leading reference in reproductive health for over 30 years. Edited for ob/gyns, reproductive medicine MDs, and primary care physicians, this

guide covers the details of every method of contracetion, looking at the

benefits, risks, and reliability of each.

Challenging Choices - Erika Dyck 2020-11-18

Between the decriminalization of contraception in 1969 and the introduction of the Charter of Rights and Freedoms in 1982, a landmark decade in the struggle for women's rights, public discourse about birth control and family planning was transformed. At the same time, a transnational conversation about the "population bomb" that threatened global famine caused by overpopulation embraced birth control technologies for a different set of reasons, revisiting controversial ideas about eugenics, heredity, and degeneration. In Challenging Choices Erika Dyck and Maureen Lux argue that reproductive politics in 1970s Canada were shaped by competing ideologies on global population control, poverty, personal autonomy, race, and gender. For some Canadians the 1970s did not bring about an era of reproductive liberty but instead reinforced traditional power dynamics and paternalistic structures of authority. Dyck and Lux present case studies of four groups of Canadians who were routinely excluded from progressive, reformist discourse: Indigenous women and their communities, those with intellectual and physical disabilities, teenage girls, and men. In different ways, each faced new levels of government regulation, scrutiny, or state intervention as they negotiated their reproductive health, rights, and responsibilities in the so-called era of sexual liberation. While acknowledging the reproductive rights gains that were made in the 1970s, the authors argue that the legal changes affected Canadians differently depending on age, social position, gender, health status, and cultural background. Illustrating the many ways to plan a modern family, these case studies reveal how the relative merits of life and choice were pitted against each other to create a new moral landscape for evaluating classic questions about population control.

Ellen A. Brantlinger - 2019-10-01

Ellen A. Brantlinger: When Meaning Falter and Words Fail, Ideology Matters considers the impact of Ellen A. Brantlinger, a foundational leader of Disability Studies in Education upon the contributors efforts to advance DSE as a field of inquiry.

Sterilized by the State - Randall Hansen 2013-08-26

This book is the first comprehensive analysis of eugenics in North America focused on the second half of the twentieth century. Based on new research, Randall Hansen and Desmond King show why eugenic sterilization policies persisted after the 1940s in the United States and Canada. Through extensive archival research, King and Hansen show how both superintendents at homes for the 'feebleminded' and prosterilization advocates repositioned themselves after 1945 to avoid the taint of Nazi eugenics. Drawing on interviews with victims of sterilization and primary documents, this book traces the post-1940s development of eugenic policy and shows that both eugenic arguments and committed eugenicists informed population, welfare, and birth control policy in postwar America. In providing revisionist histories of the choice movement, the anti-population growth movement, and the Great Society programs, this book contributes to public policy and political and intellectual history.

Disability Studies Quarterly - 1997

Genomics and Health in the Developing World - Dhavendra Kumar 2012-06-14

Readership: Geneticists and clinicians worldwide in addition to graduate students and researchers interested in populations and genomics European Social Work - A Compendium - Fabian Kessl 2019-11-25 The publication takes account of the fundamental developments transforming social work in Europe at the beginning of the 21st century. A European standard of social work has already emerged, but models for future European social work are absent. Therefore the compendium gives an overview of the current transformation process for the first time, discusses the visible and invisible changes and maps out where social work is positioned in the emerging post-welfare states.

Eugenic Sterilization - Jonas B. Robitscher 1973

Beyond Guardianship - National Council on Disability 2018-03-27 In general, guardianship involves a state-court determination that an individual lacks the capacity to make decisions with respect to their health, safety, welfare, and/or property. This Beyond Guardianship report explains how guardianship law has evolved, explores the due process and other concerns with guardianships, offers an overview of alternatives to guardianship, and identifies areas for further study. This report covers people with mental illness or disabilities, to include children populations and aging adult populations Legal standards of incapacity are also

explored within this report. Discover more products related to this topic: Physically challenged collection and resources about persons that are disabled Aging resources collection Mental Health collection Childhood & Adolescence collection

 $\underline{\textbf{Psychosocial Aspects of Disability}} \text{ - Noreen M. Glover-Graf, RhD, CRC} \\ 2011\text{-}07\text{-}27$

"What a marvelous and amazing textbook. Drs. Marini, Glover-Graf and Millington have done a remarkable job in the design of this highly unique book, that comprehensively and very thoughtfully addresses the psychosocial aspects of the disability experience. These highly respected scholars have produced a major work that will be a central text in rehabilitation education for years to come." From the Foreword by Michael J. Leahy, Ph.D., LPC, CRC Office of Rehabilitation and Disability Studies Michigan State University "This is an excellent book, but the best parts are the stories of the disabled, which give readers insights into their struggles and triumphs." Score: 94, 4 Stars--Doody's Medical Reviews What are the differences between individuals with disabilities who flourish as opposed to those who never really adjust after a trauma? How are those born with a disability different from individuals who acquire one later in life? This is the first textbook about the psychosocial aspects of disability to provide students and practitioners of rehabilitation counseling with vivid insight into the experience of living with a disability. It features the first-person narratives of 16 people living with a variety of disabling conditions, which are integrated with sociological and societal perspectives toward disability, and strategies for counseling persons with disabilities. Using a minority model perspective to address disability, the book focuses on historical perspectives, cultural variants regarding disability, myths and misconceptions, the attitudes of special interest and occupational groups, the psychology of disability with a focus on positive psychology, and adjustments to disability by the individual and family. A wealth of counseling guidelines and useful strategies are geared specifically to individual disabilities. Key Features: Contains narratives of people living with blindness, hearing impairments, spinal cord injuries, muscular dystrophy, polio, mental illness, and other disabilities Provides counseling guidelines and strategies specifically geared toward specific disabilities, including "dos and don'ts" Includes psychological and sociological research relating to individual disabilities Discusses ongoing treatment issues and ethical dilemmas for rehabilitation counselors Presents thought-provoking discussion questions in each chapter Authored by prominent professor and researcher who became disabled as a young adult

Sexuality and Disabilities - Romel W. Mackelprang 1993 Individuals with disabilities are often "desexualized" in our society, yet they have the same need for intimacy, self-worth, and social belonging as people without disabilities. Sexuality and Disabilities addresses persons with physical, sensory, intellectual, and cognitive disabilities and their concerns in the areas of intimacy, family issues, sexuality, and sexual functioning. It offers suggestions for professionals who work with persons with these disabilities to help them work more competently with disabled persons in the sexuality arena. These concrete ideas are excellent for staff training and education and for enhancing professional development for those working with persons with physical disabilities. The contributing authors create an awareness that all people need individualized consideration and that the special needs of all individuals are important, especially for those who may have previously been left to discover things on their own--usually unsuccessfully. Sexuality and Disabilities focuses on a wide range of disabilities, including physical, developmental, and learning disabilities, mental retardation, and conditions that may have an impact on people later in life such as strokes, heart disease, or other chronic illness. Chapters discuss education and support issues for both practitioners and clients. Some of the topics examined include: components of a staff training program on sexuality and disability specific recommendations for sexuality education and counseling with people with spinal cord injuries and other acquired severe neurological disabilities a program model serving parents with mental retardation and their children specific ways educational programming, social work intervention, and policy efforts can address the special learning needs of people with cognitive impairments sources of support and stress for families caring for developmentally disabled children an analysis of special vulnerabilities and challenges relating to sexual victimization that confront people with disabilities An extremely helpful tool for human service practitioners, Sexuality and Disabilities is also a valuable resource for graduate and undergraduate students who have an interest in working with people with physical, cognitive, or

mental disabilities and helping them explore this basic facet of their lives

Beyond Testimony and Trauma - Steven High 2015-03-15 Survivors of terrible events are often portrayed as unsung heroes or tragic victims but rarely as complex human beings whose lives extend beyond the stories they have told. The contributors to Beyond Testimony and Trauma consider other ways to engage with survivors and their accounts based on valuable insights gained from their work on long-term oral history projects. While the contexts vary widely, they demonstrate that through deep listening, long-term relationship building, and collaborative research design, it is possible to move beyond the problematic aspects of "testimony" to shine a light on the more nuanced lives of survivors of mass violence.

Reconsidering Intellectual Disability - Jason Reimer Greig 2015-11-02

Drawing on the controversial case of "Ashley X," a girl with severe developmental disabilities who received interventionist medical treatment to limit her growth and keep her body forever small—a procedure now known as the "Ashley Treatment"—Reconsidering Intellectual Disability explores important questions at the intersection of disability theory, Christian moral theology, and bioethics. What are the biomedical boundaries of acceptable treatment for those not able to give informed consent? Who gets to decide when a patient cannot communicate their desires and needs? Should we accept the dominance of a form of medicine that identifies those with intellectual impairments as pathological objects in need of the normalizing bodily manipulations of technological medicine? In a critical exploration of contemporary disability theory, Jason Reimer Greig contends that L'Arche, a federation of faith communities made up of people with and without intellectual disabilities, provides an alternative response to the predominant bioethical worldview that sees disability as a problem to be solved. Reconsidering Intellectual Disability shows how a focus on Christian theological tradition's moral thinking and practice of friendship with God offers a way to free not only people with intellectual disabilities but all people from the objectifying gaze of modern medicine. L'Arche draws inspiration from Jesus's solidarity with the "least of these" and a commitment to Christian friendship that sees people with profound cognitive disabilities not as anomalous objects of pity but as fellow friends of God. This vital act of social recognition opens the way to understanding the disabled not as objects to be fixed but as teachers whose lives can transform others and open a new way of being human.

Handbook of Disability Studies - Gary L. Albrecht 2001

This path-breaking international handbook of disability studies signals the emergence of a vital new area of scholarship, social policy and activism. Drawing on the insights of disability scholars around the world and the creative advice of an international editorial board, the book engages the reader in the critical issues and debates framing disability studies and places them in an historical and cultural context. Five years in the making, this one volume summarizes the ongoing discourse ranging across continents and traditional academic disciplines. To provide insight and perspective, the volume is divided into three sections: The shaping of disability studies as a field; experiencing disability; and, disability in context. Each section, written by world class figures, consists of original chapters designed to map the field and explore the key conceptual, theoretical, methodological, practice and policy issues that constitute the field. Each chapter provides a critical review of an area, positions and literature and an agenda for future research and practice. The handbook answers the need expressed by the disability community for a thought provoking, interdisciplinary, international examination of the vibrant field of disability studies. The book will be of interest to disabled people, scholars, policy makers and activists alike. The book aims to define the existing field, stimulate future debate, encourage respectful discourse between different interest groups and move the field a step forward.

$\begin{tabular}{ll} \textbf{Women With Intellectual Disabilities} & -Rannveig\ Traustadottir\\ 2000-03 \end{tabular}$

I recommend this book to anyone engaged in working collaboratively with people with the label 'learning difficulty', particularly in women's; groups, self advocacy or rights bases/citizenship concerns. The plain English accounts are accessible, but I also found the main bulk of the text easily translatable and used it extensively in my recent research. For the women involved in this project it provided a framework of reference in which they recognized similar life events and experiences. Not only does this book fill this gap by providing a frame in which women can examine this exclusion, it also questions the marginalized position of

women classified as having 'learning difficulties' in feminist and disability literature.' - Disability and Society 'This is such a good read that it is difficult to be objective about the content, criticism was suspended! It is divided into parts and each part is helpfully introduced by the editors. There is also a short straightforward description of the content at the beginning of each chapter so that women with intellectual disabilities can be included in the readership. This book gives us a valuable insight into the lives of women with learning disabilities. It changes an often discriminated group into individuals of considerable interest and value. It is to be recommended to everyone who feels that difference is important in our community.' - Ann Craft Trust Bulletin This book provides the first comprehensive exploration of the issues affecting the lives of women with intellectual disabilities. Women from all over the world, with and without intellectual disabilities, have collaborated to write about their lives, their experiences and their hopes for the future. Different aspects of life - work, family, relationships and community involvement - are discussed. Some of the women have found, or are finding, fulfilling, happy, creative lifestyles. One message which emerging from many of their stories is that their intellectual disability is less of a problem than the social and economic discrimination these women experience. This book thus raises important questions about society's attitudes to women with intellectual disabilities. It is also a place where these women's stories - from the sad or disturbing to the happy, moving or inspirational - can be heard. The book's unique plain English versions of chapters will ensure that it is accessible to other women with intellectual disabilities. It is an important, interesting and readable addition to literature about intellectual disabilities and about women's lives across the world. **Developmental Challenges and Societal Issues for Individuals**

Developmental Challenges and Societal Issues for Individuals With Intellectual Disabilities - Gopalan, Rejani Thudalikunnil 2019-08-30

Intellectual disability is a generalized disorder appearing before adulthood characterized by significantly impaired cognitive functioning and deficits in two or more adaptive behaviors. With the current limitations in curative treatment for intellectual disabilities, the rehabilitation and management of affected individuals remains a major factor in the management and treatment of symptoms and for the improvement of daily life. Developmental Challenges and Societal Issues for Individuals With Intellectual Disabilities is a comprehensive academic resource that examines treatment and rehabilitation options for those who have intellectual disabilities and examines educational, vocational, and psychosocial needs that can improve quality of life for these individuals. Featuring a range of topics such as comorbidities, epidemiology, and stigma, this book is ideal for psychologists, psychiatrists, pediatricians, psychiatric nurses, clinicians, special ed teachers, social workers, hospital administrators, mental health specialists, managers, academicians, rehabilitation centers, researchers, and students.

The Development of Disability Rights Under International Law - Arlene S. Kanter 2014-11-27

The adoption of the Convention on the Rights of People with Disabilities (CPRD) by the United Nations in 2006 is the first comprehensive and binding treaty on the rights of people with disabilities. It establishes the right of people with disabilities to equality, dignity, autonomy, full participation, as well as the right to live in the community, and the right to supported decision-making and inclusive education. Prior to the CRPD, international law had provided only limited protections to people with disabilities. This book analyses the development of disability rights as an international human rights movement. Focusing on the United States and countries in Asia, Africa, the Middle East the book examines the status of people with disabilities under international law prior to the adoption of the CPRD, and follows the development of human rights protections through the convention's drafting process. Arlene Kanter argues that by including both new applications and entirely new approaches to human rights treaty enforcement, the CRPD is significant not only to people with disabilities but also to the general development of international human rights, by offering new human rights protections for all people. Taking a comparative perspective, the book explores how the success of the CRPD in achieving protections depends on the extent to which individual countries enforce domestic laws and policies, and the changing public attitudes towards people with disabilities. This book will be of excellent use and interest to researchers and students of human rights law, discrimination, and disability studies.

Challenges to the Human Rights of People with Intellectual **Disabilities** - Frances Owen 2008-12-15

'A book such as this both demonstrates the progress that has been made

over recent years, and will also serve to enhance respect for the human rights of persons with intellectual disabilities in the years to come.' -From the Foreword by Orville Endicott This wide-ranging volume provides a multidisciplinary examination of human rights and the lives of people with intellectual disabilities. The book combines historical, psychological, philosophical, social, educational, medical and legal perspectives to form a unique and insightful account of the subject. Initial chapters explain the historical context of rights for people with intellectual disabilities, including the right to life, and propose a conceptual framework to inform contemporary practice. Contributors then explore the many theoretical and practical challenges that people with intellectual disabilities face, in exercising their civil rights, educational rights or participatory rights, for instance. The implications arising from these issues are identified and practical guidelines for support and accommodation are provided. This book will be an essential resource for practitioners, advocates, lawyers, policy-makers and students on disability courses.

Ethical Issues in Neurology - James L. Bernat 2008

Written by an eminent authority from the American Academy of Neurology's Committee on Ethics, Law, and Humanities, this book is an excellent text for all clinicians interested in ethical decision-making. The book features outstanding presentations on dying and palliative care, physician-assisted suicide and voluntary active euthanasia, medical futility, and the relationship between ethics and the law. New chapters in this edition discuss how clinicians resolve ethical dilemmas in practice and explore ethical issues in neuroscience research. Other highlights include updated material on palliative sedation, advance directives, ICU withdrawal of life-sustaining therapy, gene therapy, the very-low-birth-weight premature infant, the developmentally disabled patient, informed consent, organizational ethics, brain death controversies, and fMRI and PET studies relating to persistent vegetative state.

Regulating Lives - John McLaren 2002

Nine essays investigate the history of law as an instrument of social control, moral regulation, and the government, focusing primarily on British Columbia, Canada, where most of the contributors work as scholars in law or criminology. Among the areas they tackle are the sex trade, the spread of venereal disease, the use and abuse of liquor, child welfare, mental disorder, intrafamily sexual abuse, Aboriginal culture and traditions, and Doukhobor beliefs and customs. The studies rely on forays into archival material at the national, provincial, and local levels. Annotation copyrighted by Book News, Inc., Portland, OR

Bioethics and the Law - Janet Dolgin 2018-12-14 Bioethics and the Law takes a multidisciplinary approach that combines legal discussion with jurisprudential, philosophical, and sociological materials. Strong expressions of different points of view highlight debates about bioethical issues. The text underscores the need to mediate between the law's focus on broad rules and the bioethicist's concern with context and detail. Students are required to consider the ethical implications of health care as a business, face the shifting parameters of the provider/patient relationship in healthcare, and understand the role of government in designing and implementing healthcare programs such as Medicaid and Medicare. Bioethics and the Law supplements the traditional focus of bioethics on the interest of the individual with a second focus on the socio-economic developments that shape healthcare. Connecting broad public healthcare issues to concerns of the individual patient/healthcare consumer, the text promotes understanding of unsettling and complex situations and shows the implications of bioethical developments for understandings of personhood. A helpful glossary defines basic terms and several short appendices summarize recent developments in science and technology. The Psychological and Social Impact of Illness and Disability, 6th Edition

- Dr. Irmo Marini 2012-02-24 Print+CourseSmart