

GENETIC INFORMATION

Acquisition, Access, and Control

Edited by
Alison K. Thompson
and
Ruth F. Chadwick

Genetic Information Acquisition Access And Control

Randall E. Auxier, Lewis Edwin Hahn



Genetic Information Acquisition Access And Control:

Genetic Information Alison K. Thompson, Ruth F. Chadwick, 2007-07-23 It is difficult to think of an example of an advancement in the biological sciences that has had an impact on society similar to that of the new genetics. Recent developments in biotechnology have occasioned much discussion among academics, professionals and lay people alike. In particular, many questions and concerns have arisen over the acquisition, access and control of genetic information. There are several reasons why the new genetics has commanded such widespread attention and why it is now the subject of considerable debate. Special reference is given in this volume to the implications of genetic information for five different subject areas: eugenics, the insurance industry, the commercialisation of genetic testing, strategies for raising public awareness and the value of theoretical, ethical and sociological frameworks in the debate. This diverse collection of papers attempts to address and critically discuss issues surrounding the control of and access to genetic information from ethical, medical, legal and theoretical points of view. The first and shortest section of the book attempts to address concerns over the eugenic potential of new biotechnologies. It also provides a historical context for the debate for controversy over the subject of eugenics predates the current debate over genetic information by a considerable length of time. Indeed, by the time the first patent was issued for Chakrabarty's strain of oil-eating bacteria in the early 1970s, the term eugenics had already acquired strong pejorative connotations.

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Governing Biobanks Jane Kaye, Susan Gibbons, Catherine Heeney, Andrew Smart, 2012-02-03 Biobanks are proliferating rapidly worldwide because they are powerful tools and organisational structures for undertaking medical research. By linking samples to data on the health of individuals, it is anticipated that

biobanks will be used to explore the relationship between genes environment and lifestyle for many diseases as well as the potential of individually tailored drug treatments based on genetic predisposition However they also raise considerable challenges for existing legal frameworks and research governance structures This book critically examines the current governance structures in place for biobanks in England and Wales It shows that the technologies techniques and practices involved in biobanking do not always conform neatly to existing legal principles and frameworks that apply to other areas of medical research Using a socio legal approach including interview data gathered from the scientific community this book provides unique insights and makes recommendations about appropriate governance mechanisms for biobanking in the future It also explores the issues around the secondary use of information such as consent and how to protect privacy when biobanks are accessed by a number of different third parties These issues have relevance both within England and Wales and to a wide international audience as well as for other areas where large datasets are used

Genetic Data and the Law

Mark Taylor,2012-03-08 Research using genetic data raises various concerns relating to privacy protection Many of these concerns can also apply to research that uses other personal data but not with the same implications for failure The norms of exclusivity associated with a private life go beyond the current legal concept of personal data to include genetic data that relates to multiple identifiable individuals simultaneously and anonymous data that could be associated with any number of individuals in different but reasonably foreseeable contexts It is the possibilities and implications of association that are significant and these possibilities can only be assessed if one considers the interpretive potential of data They are missed if one fixates upon its interpretive pedigree or misunderstands the meaning and significance of identification This book demonstrates how the public interest in research using genetic data might be reconciled with the public interest in proper privacy protection

Genetic Information Values and Rights Niklas Juth,2005

Genes and Insurance Marcus

Radetzki,Marian Radetzki,Niklas Juth,2003-11-06 The result of two key social developments in recent years are examined here the partial dismantling of the welfare state and the progress of genetics Genetic insights are increasingly valuable for risk assessment and insurers would like to use these insights to help determine premiums Combined with the fact that social welfare is being curtailed this could potentially create an uninsured high risk population Along with considerations of autonomy and privacy this is the basis for an ethical critique of insurer s access to information The result has often been regulation of such information but the authors argues that due to adverse selection regulation will not solve these problems and this may jeopardize the survival of private personal insurance Instead we should look towards the resurrection of social insurance a key component of the welfare state This will interest academic researchers as well as professionals involved with genetics and insurance

Encyclopedia of Applied Ethics ,2012-01-10

The Encyclopedia of Applied Ethics Second Edition Four Volume Set addresses both the physiological and the psychological aspects of human behavior Carefully crafted well written and thoroughly indexed the encyclopedia helps users whether they are students just beginning formal study of the

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Handbook of the Philosophy of Medicine Thomas Schramme, Mary Jean Walker, 2025-09-30 This is the second edition of Springer's first wide ranging multi authored handbook on philosophy of medicine It covers the underlying conceptual issues of many important social political and ethical issues in health care and as such provides a single source of information for this far ranging and still developing field This new edition of this authoritative handbook includes topics that have gained momentum in the last couple of years This second edition introduces and develops nearly one hundred topics concepts and issues in the field It is written by distinguished specialists from multiple disciplines including philosophy health sciences nursing sociology political theory and medicine Many complicated social and ethical issues in health care are based on conceptual problems most prominently on the definitions of health and disease or on epistemological issues regarding causality or diagnosis Philosophy is the discipline that deals with such conceptual metaphysical epistemological methodological and axiological matters This handbook covers all the central concepts in medicine such as ageing death disease mental disorder and well being It is an invaluable resource for health care specialists who want to be informed and stay up to date with the relevant discussions as well as philosophers and ethicists with an interest in medicine and members of the general public with an interest in health care and related issues The text also advances these debates and sets the agenda for years to come

Autonomy and Trust in Bioethics Onora O'Neill, 2002-04-18 Why has autonomy been a leading idea in philosophical writing on bioethics and why has trust been

marginal In this important book Onora O'Neill suggests that the conceptions of individual autonomy so widely relied on in bioethics are philosophically and ethically inadequate and that they undermine rather than support relations of trust She shows how Kant's non-individualistic view of autonomy provides a stronger basis for an approach to medicine science and biotechnology and does not marginalize untrustworthiness while also explaining why trustworthy individuals and institutions are often undeservingly mistrusted Her arguments are illustrated with issues raised by practices such as the use of genetic information by the police or insurers research using human tissues uses of new reproductive technologies and media practices for reporting on medicine science and technology Autonomy and Trust in Bioethics will appeal to a wide range of readers in ethics bioethics and related disciplines

The Handbook of Information and Computer Ethics Kenneth E. Himma, Herman T. Tavani, 2008-05-27 Discover how developments in information technology are raising new ethical debates Information and computer ethics has emerged as an important area of philosophical and social theorizing combining conceptual meta-ethical normative and applied elements As a result academic interest in this area has increased dramatically particularly in computer science philosophy and communications departments business schools information and library schools and law schools The Handbook of Information and Computer Ethics responds to this growing interest with twenty-seven chapters that address both traditional and current issues in information and computer ethics research It is organized into six parts Foundational Issues and Methodological Frameworks Theoretical Issues Affecting Property Privacy Anonymity and Security Professional Issues and the Information Related Professions Responsibility Issues and Risk Assessment Regulatory Issues and Challenges Access and Equity Issues Each chapter written by one or more of the most influential ethicists in their fields of expertise explains and evaluates the central positions and arguments on the respective issues Chapters end with a bibliography that identifies the most important supplementary books and papers available on the topic This handbook provides an accessible yet sophisticated overview of the most important issues we face in information and computer ethics today It is an ideal supplemental text for advanced undergraduate and graduate level courses in information and computer ethics and is also of interest to readers who are involved in library science computer science or philosophy

The North Carolina journal of international law and commercial regulation, 2003 Open Consent Ants Nõmper, 2005

Cloning, Selection, and Values Matti Häyry, 2007 **Rutgers Computer & Technology Law Journal**, 2008

Social Theory and Practice, 2003 Regulating Healthcare Quality John Tingle, Charles Foster, Kay Wheat, 2004 The key focus of this edited text is on the legal and professional conflicts and issues that can arise from regulating health care quality Doctors and nurses all increasingly face a number of dilemmas with regulating health quality issues such as increasing levels of complaints and litigation scarcity of resources under staffing professional discipline clinical governance clinical risk management etc This book spells out and discusses these issues taking an academic approach though this will be tempered with a practical focus on issues Discusses ethical approaches to regulating health care quality Examines health

care rights in the UK Looks at complaints procedures in the new NHS Presents alternatives to the present clinical negligence system Compares health care regulation in the US and UK Benefit-sharing Kadri Simm,2005 Information Communication Technologies Craig Van Slyke,2008 Information Communication Technologies Concepts Methodologies Tools and Applications is the essential compilation of breaking research in the pivotal areas of social adaptation to information technology This all inclusive reference source examines through case studies empirical analysis and conceptual models the successes and consequences associated with the growth of information communication technologies in the world today

The Philosophy of Marjorie Grene Randall E. Auxier,Lewis Edwin Hahn,2002 Professor Grene is the 29th subject in a series that includes Albert Einstein John Dewey Jean Paul Sartre and Bertrand Russell This book includes photographs 25 critical essays by Grene s distinguished peers and her biting replies She is known for her work and her controversial views in several areas including the philosophy of law anthropology and biology *Nature Encyclopedia of the Human Genome: Genome databases - Mitochondrial genome: Evolution* David Neil Cooper,2003

Reviewing **Genetic Information Acquisition Access And Control**: Unlocking the Spellbinding Force of Linguistics

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web during this monthlong project students learned how mrs parks sparked the montgomery bus boycott by not giving up her bus seat to a white passenger in 1955 and how as a result of that brave act in 1956 the supreme court ruled segregation on buses was illegal rosa parks died on october 25 2005 at age 92

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web shortly after the bus boycott rosa and her family moved from alabama to detroit to escape the harassment and threatening phone calls sylvester had found an apartment for rosa parks and their mother to live in rosa continued to travel and do speaking engagements

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web rosa louise mccauley was born on february 4th 1913 in tuskegee alabama as a child she went to an industrial school for girls and later enrolled at alabama state teachers college for negroes present day alabama state university unfortunately parks was forced to withdraw after her grandmother became ill

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parks she was arrested for not giving up her seat for a white passenger

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