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# HEALTH DATA IN THE INFORMATION AGE

## Use, Disclosure, and Privacy



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Social and Family History

INSTITUTE OF MEDICINE

# Health Data In The Information Age Use Disclosure And Privacy

**United States. National Bioethics  
Advisory Commission**



## **Health Data In The Information Age Use Disclosure And Privacy:**

**Health Data in the Information Age** Committee on Regional Health Data Networks, Institute of Medicine, 1994-01-15  
Regional health care databases are being established around the country with the goal of providing timely and useful information to policymakers physicians and patients But their emergence is raising important and sometimes controversial questions about the collection quality and appropriate use of health care data Based on experience with databases now in operation and in development Health Data in the Information Age provides a clear set of guidelines and principles for exploiting the potential benefits of aggregated health data without jeopardizing confidentiality A panel of experts identifies characteristics of emerging health database organizations HDOs The committee explores how HDOs can maintain the quality of their data what policies and practices they should adopt how they can prepare for linkages with computer based patient records and how diverse groups from researchers to health care administrators might use aggregated data Health Data in the Information Age offers frank analysis and guidelines that will be invaluable to anyone interested in the operation of health care databases

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administrators might use aggregated data Health Data in the Information Age offers frank analysis and guidelines that will be invaluable to anyone interested in the operation of health care databases **Legislating Privacy** Priscilla M.

Regan, 2000-11-09 While technological threats to personal privacy have proliferated rapidly legislation designed to protect privacy has been slow and incremental In this study of legislative attempts to reconcile privacy and technology Priscilla Regan examines congressional policy making in three key areas computerized databases wiretapping and polygraph testing In each case she argues legislation has represented an unbalanced compromise benefiting those with a vested interest in new technology over those advocating privacy protection *Legislating Privacy* explores the dynamics of congressional policy formulation and traces the limited response of legislators to the concept of privacy as a fundamental individual right According to Regan we will need an expanded understanding of the social value of privacy if we are to achieve greater protection from emerging technologies such as Caller ID and genetic testing Specifically she argues that a recognition of the social importance of privacy will shift both the terms of the policy debate and the patterns of interest group action in future congressional activity on privacy issues Originally published in 1995 A UNC Press Enduring Edition UNC Press Enduring Editions use the latest in digital technology to make available again books from our distinguished backlist that were previously out of print These editions are published unaltered from the original and are presented in affordable paperback formats bringing readers both historical and cultural value *Information Discovery on Electronic Health Records* Vagelis Hristidis, 2009-12-10 Exploiting the rich information found in electronic health records EHRs can facilitate better medical research and improve the quality of medical practice Until now a trivial amount of research has been published on the challenges of leveraging this information Addressing these challenges *Information Discovery on Electronic Health Records* exp *For the Record* National Research Council, Division on Engineering and Physical Sciences, Computer Science and Telecommunications Board, Commission on Physical Sciences, Mathematics, and Applications, Committee on Maintaining Privacy and Security in Health Care Applications of the National Information Infrastructure, 1997-07-09 When you visit the doctor information about you may be recorded in an office computer Your tests may be sent to a laboratory or consulting physician Relevant information may be transmitted to your health insurer or pharmacy Your data may be collected by the state government or by an organization that accredits health care or studies medical costs By making information more readily available to those who need it greater use of computerized health information can help improve the quality of health care and reduce its costs Yet health care organizations must find ways to ensure that electronic health information is not improperly divulged Patient privacy has been an issue since the oath of Hippocrates first called on physicians to keep silence on patient matters and with highly sensitive data genetic information HIV test results psychiatric records entering patient records concerns over privacy and security are growing *For the Record* responds to the health care industry's need for greater guidance in protecting health information that increasingly flows through the national information infrastructure

from patient to provider payer analyst employer government agency medical product manufacturer and beyond This book makes practical detailed recommendations for technical and organizational solutions and national level initiatives For the Record describes two major types of privacy and security concerns that stem from the availability of health information in electronic form the increased potential for inappropriate release of information held by individual organizations whether by those with access to computerized records or those who break into them and systemic concerns derived from open and widespread sharing of data among various parties The committee reports on the technological and organizational aspects of security management including basic principles of security the effectiveness of technologies for user authentication access control and encryption obstacles and incentives in the adoption of new technologies and mechanisms for training monitoring and enforcement For the Record reviews the growing interest in electronic medical records the increasing value of health information to providers payers researchers and administrators and the current legal and regulatory environment for protecting health data This information is of immediate interest to policymakers health policy researchers patient advocates professionals in health data management and other stakeholders

*Research Involving Human Biological Materials: Commissioned papers* United States. National Bioethics Advisory Commission,1999

*Research Involving Human Biological Materials: Commissioned papers. Privacy and the analysis of stored tissues* United States. National Bioethics Advisory Commission,1999

*H.R. 52, the Fair Health Information Practices Act of 1997* United States. Congress. House. Committee on Government Reform and Oversight. Subcommittee on Government Management, Information, and Technology,1998

**Beyond the HIPAA Privacy Rule** Institute of Medicine,Board on Health Care Services,Board on Health Sciences Policy,Committee on Health Research and the Privacy of Health Information: The HIPAA Privacy Rule,2009-03-24 In the realm of health care privacy protections are needed to preserve patients dignity and prevent possible harms Ten years ago to address these concerns as well as set guidelines for ethical health research Congress called for a set of federal standards now known as the HIPAA Privacy Rule In its 2009 report *Beyond the HIPAA Privacy Rule Enhancing Privacy Improving Health Through Research* the Institute of Medicine s Committee on Health Research and the Privacy of Health Information concludes that the HIPAA Privacy Rule does not protect privacy as well as it should and that it impedes important health research

**Biomedical Engineering Handbook 2** Joseph D. Bronzino,2000-02-15

**Research Involving Human Biological Materials** United States. National Bioethics Advisory Commission,1999

**Digital Health Care** Phillip Olla,Joseph K. H. Tan,2022-05-18 Binding NVA

*Medical Records Confidentiality Act of 1995* United States. Congress. Senate. Committee on Labor and Human Resources,1996

*Ethics and Information Technology* James G. Anderson,Kenneth Goodman,2007-05-28 This series is directed to health care professionals who are leading the transformation of health care by using information and knowledge Launched in 1988 as *Computers in Health Care* the series offers a broad range of titles some addressed to specific professions such as nursing medicine and health administration others to special areas of practice

such as trauma and radiology. Still other books in the series focus on interdisciplinary issues such as the computer based patient record, electronic health records and networked health care systems. Renamed Health Informatics in 1998 to reflect the rapid evolution in the discipline now known as health informatics, the series will continue to add titles that contribute to the evolution of the field. In the series, eminent experts serving as editors or authors offer their accounts of innovations in health informatics. Increasingly, these accounts go beyond hardware and software to address the role of information in influencing the transformation of healthcare delivery systems around the world. The series also increasingly focuses on peopleware and the organizational, behavioral and societal changes that accompany the diffusion of information technology in health services environments.

**The History and Heritage of Scientific and Technological Information Systems** W. Boyd Rayward, Mary Ellen Bowden, 2004. Emphasis for the second conference on the history of information science systems was on scientific and technical information systems in the period from the Second World War up through the early 1990s. These proceedings present the papers of historians of science and technology, information scientists and scientists in other fields on a wide range of topics: informatics in chemistry, biology and medicine; information developments in multinational industrial and military settings; biographical studies of pioneering individuals and the transformation of information systems and formats in the twentieth century.

*The Fair Health Information Practices Act of 1994* United States. Congress. House. Committee on Government Operations. Information, Justice, Transportation, and Agriculture Subcommittee, 1994.

**Protecting Health Information** United States. Congress. House. Committee on Government Reform and Oversight. Subcommittee on Government Management, Information, and Technology, 1999.

**H.R. 3137, Data Needs and Related Issues for Implementing Health Care Reform** United States. Congress. House. Committee on Post Office and Civil Service. Subcommittee on Census, Statistics, and Postal Personnel, 1994.

**The role of FCRA in employee background checks and the collection of medical information** United States. Congress. House. Committee on Financial Services. Subcommittee on Financial Institutions and Consumer Credit, 2003.

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