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This User's Guide is intended to support the design, implementation, analysis, interpretation, and quality evaluation of registries created to increase understanding of patient outcomes. For the purposes of this guide, a patient registry is an organized system that uses observational study methods to collect uniform data (clinical

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and other) to evaluate specified outcomes for a population defined by a particular disease, condition, or exposure, and that serves one or more predetermined scientific, clinical, or policy purposes. A registry database is a file (or files) derived from the registry. Although registries can serve many purposes, this guide focuses on registries

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created for one or more of the following purposes: to describe the natural history of disease, to determine clinical effectiveness or cost-effectiveness of health care products and services, to measure or monitor safety and harm, and/or to measure quality of care. Registries are classified according to how their populations are defined. For example,

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product registries include patients who have been exposed to biopharmaceutical products or medical devices. Health services registries consist of patients who have had a common procedure, clinical encounter, or hospitalization. Disease or condition registries are defined by patients having the same diagnosis, such as cystic fibrosis or



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heart failure. The User ' s Guide was created by researchers affiliated with AHRQ ' s Effective Health Care Program, particularly those who participated in AHRQ ' s DEcIDE (Developing Evidence to Inform Decisions About Effectiveness) program. Chapters were subject to multiple internal and external independent reviews.

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This manual is aimed at helping medical record workers in the development and management of medical records services of health care facilities in developing countries in an effective and efficient manner. It has not been designed as an introductory text to medical record management, but rather as

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an aid to medical record officers (MROs) and medical record clerks by describing appropriate systems for Medical Records Departments in developing countries. It covers manual procedures and may be used as an adjunct to computerized systems. It does not provide all of the options for medical record management, but it does

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provide one option in each area for the management of medical records in developing countries. A list the textbooks that provide detailed information on medical record management is also provided.

This manual has been designed as a basic

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reference for use when exploring the development and implementation of electronic health record (EHR) systems. It provides a general overview, some basic definitions and examples of EHR practices. Also covered are points for consideration when moving towards the introduction of an EHR, some issues and challenges which

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may need to be addressed and some possible strategies, along with steps and activities to implementation. There is a particular focus on setting goals, revising policies, developing an action plan and outlining implementation procedures.

When hospitals began implementing their

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electronic medical records/electronic health records systems (EMR/HER) the pharmaceutical companies that were conducting clinical trials at those hospitals wanted to sue the date from those systems instead of having the hospitals enter the data in their EHR systems and also in the study data entry system. However, the FDA

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regulations would require that the hospital systems be "validated". The hospitals and the companies developing the systems argued that was "over-regulation." HIMSS published their Developer Code of Conduct where they said instead they would use Quality Management techniques. This book covers how to use Quality Management



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(ISO 9001) to develop computer systems, specifically EMR systems. It gives a basic introduction to how to implement computer systems. It also covers the topic of compliance because the hospitals are required to comply with regulations other than FDS regulations. The book also discusses the topics of risk management and

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conducting audits, both of which are part of ISO 9001 quality management of computer systems. The book is designed to give the reader an introduction to the things you have to do when implementing a computer system that has to satisfy some standards and where the accuracy of the information could impact the accuracy of a person ' s medical

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Providing policies and guidance for all aspects of EHR, including: Understanding meaningful use incentives Improving patient care Ensuring privacy and security protection Addresses real-life daily challenges from the field Puts policies into context for the user Policies in the second edition have been updated to reflect

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An EHR transformation touches virtually every aspect of a medical practice and brings



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about an entirely new way of thinking and managing a practice. Regardless of where you are at in your EHR implementation journey--adopting a new EHR or trying to optimize an existing EHR, this book explores the process in a practical, easy-to-follow way, offering proven strategies for success. Readers will learn methods for

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developing an implementation plan and project budget, selecting the right vendor and preparing your medical practice for transitioning from paper records. This book also addresses federal standards and policies to ensure readers fully understand compliance requirements and the opportunities to take advantage of financial

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Health Informatics (HI) focuses on the application of Information Technology (IT) to the field of medicine to improve individual and population healthcare delivery, education and research. This extensively updated fifth edition reflects the

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