

Medical Devices Used In Home Health Care

Sabine Zange

Medical Devices Used In Home Health Care:

The Role of Human Factors in Home Health Care National Research Council, Division of Behavioral and Social Sciences and Education, Committee on Human-Systems Integration, Committee on the Role of Human Factors in Home Health Care, 2010-11-14 The rapid growth of home health care has raised many unsolved issues and will have consequences that are far too broad for any one group to analyze in their entirety Yet a major influence on the safety quality and effectiveness of home health care will be the set of issues encompassed by the field of human factors research the discipline of applying what is known about human capabilities and limitations to the design of products processes systems and work environments To address these challenges the National Research Council began a multidisciplinary study to examine a diverse range of behavioral and human factors issues resulting from the increasing migration of medical devices technologies and care practices into the home Its goal is to lay the groundwork for a thorough integration of human factors research with the design and implementation of home health care devices technologies and practices On October 1 and 2 2009 a group of human factors and other experts met to consider a diverse range of behavioral and human factors issues associated with the increasing migration of medical devices technologies and care practices into the home This book is a summary of that workshop representing the culmination of the first phase of the study **Federal Policies and the Medical Devices** Human Factors Interventions for the Health Care of Older Adults Wendy A. Rogers, Arthur D. Industry ,1984 Fisk, 2001-08-01 Written by experts in the field this book explores the relevance and contributions of the field of human factors to health care It begins with overviews of the field of human factors and the primary research methodologies of that field and goes on to review the cognitive issues that must be considered in the context of the health care environment and the potential for exercises to improve such cognitive functions. The remaining chapters cover a range of cutting edge topics including care giving telecommunication issues design of medical devices computer monitoring of patients automated communication systems computer interface issues in general and the use of the Web as a source for health information

Naturalistic Decision Making Caroline E. Zsambok, Gary Klein, 2014-01-02 If you aren t using the term naturalistic decision making or NDM you soon will be Even as a very young field NDM has already had far reaching applications in areas as diverse as management aviation health care nuclear power military command and control corporate teamwork and manufacturing Put simply NDM is the way people use their experience to make decisions in the context of a job or task Of particular interest to NDM researchers are the effects of high stake consequences shifting goals incomplete information time pressure uncertainty and other conditions that are present in most of today s work places and that add to the complexity of decision making Applications of NDM research findings target decision aids and training that help people in their decision making processes This book reports the findings of top NDM researchers as well as many of their current applications In addition the book offers a historical perspective on the emergence of this new paradigm describes recent theoretical and

methodological advancements and points to future developments It was written for people interested in decision making research and applications relative to a diverse array of work settings and products such as human computer interfaces decision support systems individual and team training product designs and organizational development and planning

Medical Devices World Health Organization, 2010 Background papers 1 to 9 published as technical documents Available in separate records from WHO HSS EHT DIM 10 1 to WHO HSS EHT DIM 10 9 Plastics in Medical Devices Vinny R. Sastri, 2021-11-24 Plastics in Medical Devices Properties Requirements and Applications Third Edition provides a comprehensive overview on the main types of plastics used in medical device applications. The book focuses on the applications and properties that are most important in medical device design such as chemical resistance sterilization capability and biocompatibility The roles of additives stabilizers and fillers as well as the synthesis and production of polymers are covered and backed up with a wealth of data tables The book also covers other key aspects in detail including regulations compliance purchasing controls and supplier controls and process validation This updated edition has been thoroughly revised with regard to new plastic materials applications and requirements This is a valuable resource for engineers scientists and managers involved in the design and manufacture of medical devices Presents detailed coverage of commercially available plastics used in medical device applications organized by polymer type and supported by data Includes up to date regulatory requirements and practical information on purchasing and supplier controls process validation and risk management Supports the development marketing and commercialization of medical devices and materials for use The Future of Medical Device Regulation I. Glenn Cohen, Timo Minssen, W. Nicholson Price in medical devices II, Christopher Robertson, Carmel Shachar, 2022-04-07 Regulators have been more permissive for medical devices compared to their drug and biologic counterparts While innovative products can thereby reach consumers more guickly this approach raises serious public health and safety concerns Additionally the nature of medical devices is rapidly changing as software has become as important as hardware Regulation must keep pace with the current developments and controversies of this technology This volume provides a multidisciplinary evaluation of the ethical legal and regulatory concerns surrounding medical devices in the US and EU For medical providers policymakers and other stakeholders the book offers a framework for the opportunities and challenges on the horizon for medical device regulation Readers will gain a nuanced overview of the latest developments in patient privacy and safety innovation and new regulatory laws This book is also available as Open Access on Cambridge Core Handbook of Human Factors in Medical Device Design Matthew Bret Weinger, Michael E. Wiklund, Daryle Jean Gardner-Bonneau, 2010-12-13 Developed to promote the design of safe effective and usable medical devices Handbook of Human Factors in Medical Device Design provides a single convenient source of authoritative information to support evidence based design and evaluation of medical device user interfaces using rigorous human factors engineering principles It offers guidance Healthcare and Medical Devices Jay Kalra and Nancy J. Lightner, 2022-07-24

Healthcare and Medical Devices Proceedings of the 13th International Conference on Applied Human Factors and Ergonomics AHFE 2022 July 24 28 2022 New York USA Encyclopedia of Medical Devices and Instrumentation John G. Webster, 2006-04-07 The articles in The Encyclopedia of Medical Devices and Instrumentation focus on what is currently useful or is likely to be useful in future medicine. They answer the question What are the branches of medicine and how does technology assist each of them Articles focus on the practice of medicine that is assisted by devices rather than including for example the use of drugs to treat disease The title is the only resource on the market dealing with the subject in encyclopedic detail Accessible to practitioners with a broad range of backgrounds from students to researchers and physicians Articles cover the latest developments such as nanotechnology fiber optics and signal processing Healthcare Technology Management - A Systematic Approach Francis Hegarty, John Amoore, Paul Blackett, Justin McCarthy, Richard Scott, 2017-01-06 Healthcare Technology Management A Systematic Approach offers a comprehensive description of a method for providing safe and cost effective healthcare technology management HTM The approach is directed to enhancing the value benefit in relation to cost of the medical equipment assets of healthcare organizations to best support patients clinicians and other care providers as well as financial stakeholders. The authors propose a management model based on interlinked strategic and operational quality cycles which when fully realized delivers a comprehensive and transparent methodology for implementing a HTM programme throughout a healthcare organization The approach proposes that HTM extends beyond managing the technology in isolation to include advancing patient care through supporting the application of the technology The book shows how to cost effectively manage medical equipment through its full life cycle from acquisition through operational use to disposal and to advance care adding value to the medical equipment assets for the benefit of patients and stakeholders This book will be of interest to practicing clinical engineers and to students and lecturers and includes self directed learning questions and case studies Clinicians Chief Executive Officers Directors of Finance and other hospital managers with responsibility for the governance of medical equipment will also find this book of interest and value For more information about the book please visit the website Monthly Catalog of United States Government Publications ,1994

Monthly Catalogue, United States Public Documents ,1994 Medical Devices Bulletin , Safe Medical Devices for Children Institute of Medicine, Board on Health Sciences Policy, Committee on Postmarket Surveillance of Pediatric Medical Devices,2005-12-20 Innovative medical devices have helped reduce the burden of illness and injury and improve the quality of life for countless children Mechanical ventilators and other respiratory support devices rescue thousands of fragile newborns every year Children who once would have died of congenital heart conditions survive with the aid of implanted pacemakers mechanical heart valves and devices that close holes in the heart Responding to a Congressional request the Institute of Medicine assesses the system for postmarket surveillance of medical devices used with children The book specifically examines The Food and Drug Administration s monitoring and use of adverse event reports

The agency's monitoring of manufacturers fulfillment of commitments for postmarket studies ordered at the time of a device s approval for marketing The adequacy of postmarket studies of implanted devices to evaluate the effects of children's active lifestyles and their growth and development on device performance Postmarket surveillance of medical devices used with children is a little investigated topic in part because the market for most medical products is concentrated among older adults Yet children differ from adults and their special characteristics have implications for evaluation and monitoring of the short and long term safety and effectiveness of medical devices used with young patients Pharmacoepidemiology Brian L. Strom, 2006-01-04 The fourth edition of Pharmacoepidemiology is an outstanding and fully comprehensive textbook which will be an essential resource for all interested in the field in academia in regulatory agencies in industry and in the law Brian Strom s classic textbook continues both to reflect their reased maturation of pharmacoepedemiology and to help shapeit s direction Reviews of previous editions of his celebrated textbookinclude The book is essential reading for anyone interested inpharmacoepidemiology INTERNATIONAL JOURNAL OF EPIDEMIOLOGY an excellent textbook and a comprehensivereference which belongs in the library of everypharmaceutical manufacturer and regulator EUROPEAN Developing Drug Products in an Aging Society Sven Stegemann, 2016-10-20 This book IOURNAL OF PUBLIC HEALTH aims to address the major aspects of future drug product development and therapy for older adults giving practical guidance for the rational product and clinical development and prescribing of drug products to this ever growing segment of the population With authors coming from key aging markets such as Europe the USA China and Japan the book will provide valuable information for students scientists regulators practitioners and other healthcare professionals from academia industry and regulatory bodies Improving Patient Care by Reporting Problems with Medical Devices, 1997

Corporate Compliance in Home Health Fay Adrienne Rozovsky,1998 Litigation is rampant in the home health field and it s not just about billing and coding It s also about other practices that could lead to compliance problems such as substandard employment practices or failing to run background checks A lawsuit against your agency could come completely out of left field But you can avoid getting blind sided and protect your agency Corporate Compliance in Home Health Establishing a Plan Managing the Risks gives home health care providers practical down to earth standards for controlling and preventing losses growing out of corporate compliance With this new resource from Aspen get an up to date and easy to understand review of the law of corporate compliance find out how the new OIG Office of the Inspector General of HCFA model compliance guidance for hospitals impacts home health and uncover where your agency is at risk You ll find out what to do if and when your agency comes under investigation and get important timely federal reference material from the GAO to HHS and key parts of the Federal Sentencing Guidelines

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