

Compliance Checklist for IFU, Labelling and Packaging Requirements according to EU MDR 2017/745

IFU Requirements

EU MDR Section	What shall be included on the IFU	Applicability To Which Device	Applicable to the Organization (YES/NO)	Comments
23.4 (a) / [23.2 a]	The name or trade name of the device.	All the devices		
23.4 (a) / [23.2 c]	Name of manufacturer and registered place of business.	All the devices		
23.4 (a) / [23.2 e]	An indication if the device contains a) a medicinal substance, including human cells or blood b) tissues or cells, of human or animal origin.	Applicable to devices containing specific substances, as reported in the requirement description		
23.4 (a) / [23.2 f]	When devices contain substances which are carcinogenic, mutagenic or toxic to reproduction (Ref. category 1A or 1B, in accordance with Part 3 of Annex VI to Regulation (EC) No 1272/2008) or substances having endocrine-disrupting properties for which there is scientific evidence of probable serious effects to human health, these substances shall be listed in the IFU.	Applicable to devices containing specific substances, as reported in the requirement description		
23.4 (a) / [23.2 k]	Indication of any special storage and/or handling condition	All the devices		
23.4 (a) / [23.2 l]	Indication of device sterile state and the sterilisation method.	Sterile devices		
23.4 (a) / [23.2 n]	Indication of single use	Single Use Devices		
23.4 (a) / [23.2 r]	Overall qualitative composition of the device and quantitative information on the main constituent	For devices which are composed of substances or of combination of		

Essential Requirements Checklist Medical Device

Shasha Hu



Essential Requirements Checklist Medical Device:

Handbook of Medical Device Design Richard C. Fries, 2019-08-15 First published in 2001 This handbook has been written to give those professionals working in the development and use of medical devices practical knowledge about biomedical technology regulations and their relationship to quality health care

Reliable Design of Medical Devices Richard C. Fries, 2005-11-21 As medical devices increase in complexity concerns about efficacy safety quality and longevity increase in stride Introduced nearly a decade ago Reliable Design of Medical Devices illuminated the path to increased reliability in the hands on design of advanced medical devices With fully updated coverage in its Second Edition this practical guide continues to be the benchmark for incorporating reliability engineering as a fundamental design philosophy The book begins by rigorously defining reliability differentiating it from quality and exploring various aspects of failure in detail It examines domestic and international regulations and standards in similar depth including updated information on the regulatory and standards organizations as well as a new chapter on quality system regulation The author builds on this background to explain product specification liability and intellectual property safety and risk management design testing human factors and manufacturing New topics include design of experiments CAD CAM industrial design material selection and biocompatibility system engineering rapid prototyping quick response manufacturing and maintainability as well as a new chapter on Six Sigma for design Supplying valuable insight based on years of successful experience Reliable Design of Medical Devices Second Edition leads the way to implementing an effective reliability assurance program and navigating the regulatory minefield with confidence

Reliable Design of Medical Devices, Third Edition Richard C. Fries, 2012-09-06 As medical devices become even more intricate concerns about efficacy safety and reliability continue to be raised Users and patients both want the device to operate as specified perform in a safe manner and continue to perform over a long period of time without failure Following in the footsteps of the bestselling second edition Reliable Design of Medical Devices Third Edition shows you how to improve reliability in the design of advanced medical devices Reliability engineering is an integral part of the product development process and of problem solving activities related to manufacturing and field failures Mirroring the typical product development process the book is organized into seven parts After an introduction to the basics of reliability engineering and failures it takes you through the concept feasibility design verification and validation design transfer and manufacturing and field activity phases Topics covered include Six Sigma for design human factors safety and risk analysis and new techniques such as accelerated life testing ALT and highly accelerated life testing HALT What's New in This Edition Updates throughout reflecting changes in the field An updated software development process Updated hardware test procedures A new layout that follows the product development process A list of deliverables needed at the end of each development phase Incorporating reliability engineering as a fundamental design philosophy this book shares valuable insight from the author's more than 35 years of experience A practical guide it helps you develop a more effective reliability

engineering program contributing to increased profitability more satisfied customers and less risk of liability **Handbook of Medical Device Regulatory Affairs in Asia** Jack Wong, Raymond Tong Kaiyu, 2013-03-27 Medical device regulation in Asia has gained more importance than ever Governments and regulatory bodies across the region have put in place new regulatory systems or refined the existing ones A registered product requires a lot of technical documentation to prove its efficacy safety and quality A smooth and successful registration process demands soft skills for dealing with various key stakeholders in the government testing centers and hospitals and among doctors Handbook of Medical Device Regulatory Affairs in Asia covers medical device regulatory systems in different countries ISO standards for medical devices clinical trial and regulatory requirements and documentation for application Government bodies the medical device industry and academics and students will find this book immensely useful in understanding the global regulatory environment and in their research and development projects **Medical Device Quality Assurance and Regulatory Compliance** Richard C. Fries, 1998-08-11 Acquaints developers of medical devices with the basic concepts and major issues of medical quality assurance and regulatory documents describes the requirements listed in these documents and provides strategies for compliance with these requirements **Design of Biomedical Devices and Systems Second edition** Paul H. King, Richard C. Fries, 2008-08-22 The design and functional complexity of medical devices and systems has increased during the past half century evolving from the level of cardiac pacemakers to magnetic resonance imaging devices Such life saving advancements are monumentally advantageous but with so much at stake a step by step manual for biomedical engineers is essential This **Design of Biomedical Devices and Systems, 4th edition** Paul H. King, Richard C. Fries, Arthur T. Johnson, 2018-10-03 This fourth edition is a substantial revision of a highly regarded text intended for senior design capstone courses within departments of biomedical engineering bioengineering biological engineering and medical engineering worldwide Each chapter has been thoroughly updated and revised to reflect the latest developments New material has been added on entrepreneurship bioengineering design clinical trials and CRISPR Based upon feedback from prior users and reviews additional and new examples and applications such as 3D printing have been added to the text Additional clinical applications were added to enhance the overall relevance of the material presented Relevant FDA regulations and how they impact the designer's work have been updated Features Provides updated material as needed to each chapter Incorporates new examples and applications within each chapter Discusses new material related to entrepreneurship clinical trials and CRISPR Relates critical new information pertaining to FDA regulations Presents new material on discovery of projects worth pursuing and design for health care for low resource environments Presents multiple case examples of entrepreneurship in this field Addresses multiple safety and ethical concerns for the design of medical devices and processes **Risk Management for Medical Device Manufacturers** Joe W. Simon, 2022-01-20 As a quality professional in the medical device industry you know all too well the importance of a risk management process and how iterative it can be Industry regulations

and standards like ISO 14971 help medical device manufacturers define risk management processes but they don't make them bulletproof that is ensure the efficacy of their products while minimizing future liability This book can help you build a bulletproof risk process You will learn how Designing product and manufacturing processes controls risks Using consistent language in a holistic closed loop risk management system leads to greater efficiency Creating useable and audit ready risk documents can support verification validation V V sampling plans Developing labels and instructions can help end users and patients clearly understand the pertinent risks Creating post market surveillance PMS processes is essential to determine if additional clinical performance studies are necessary Joe Simon holds an MBA and has been a member of ASQ since 2008 Over his nearly 30 year career he worked with numerous companies as an employee and a consultant to build or improve complaint analysis trending post market surveillance PMS nonconformance NC corrective action preventive action CAPA stewardship and risk management processes

Medical Device Rommel Garcia, 2017-06-06 This book is meant to be a guide to all who want to learn about a highly regulated industry My approach is to give you the reader an example of a fictitious device and we will take it from a conceptual idea all the way to launch and beyond My intention is to incorporate the best experiences that I and other contributors have had into this book and convert them into laymans terms for those who are in need These experiences can and will be indispensable to beginners and professionals alike who are trying their hand in the medical device industry and to those who have not been out of their silo to help see how each of the systems relate to each as a whole However it should be noted that the contents of this book should be taken only as information and is not intended to demonstrate how companies can be in compliance In some instances there are multiple ways to go through the maze of regulations that are documented and made by agencies because the regulations are pretty much made and designed to be flexible and high level so that companies can adopt their systems which are solely designed for their purposes Therefore this book will try to avoid complicated words and complex technical details of engineering and statistics This book will strive to be an embodiment of the honest to goodness everyday experiences and issues that folks experience while working in the medical device industry

Design of Biomedical Devices and Systems Dragan Primorac, 2014-07-29 Apply a Wide Variety of Design Processes to a Wide Category of Design Problems Design of Biomedical Devices and Systems Third Edition continues to provide a real world approach to the design of biomedical engineering devices and or systems Bringing together information on the design and initiation of design projects from several sources this edition strongly emphasizes and further clarifies the standards of design procedure Following the best practices for conducting and completing a design project it outlines the various steps in the design process in a basic flexible and logical order What's New in the Third Edition This latest edition contains a new chapter on biological engineering design a new chapter on the FDA regulations for items other than devices such as drugs new end of chapter problems new case studies and a chapter on product development It adds mathematical modeling tools and provides new information on FDA regulations and standards

as well as clinical trials and sterilization methods Familiarizes the reader with medical devices and their design regulation and use Considers safety aspects of the devices Contains an enhanced pedagogy Provides an overview of basic design issues Design of Biomedical Devices and Systems Third Edition covers the design of biomedical engineering devices and or systems and is designed to support bioengineering and biomedical engineering students and novice engineers entering the medical device market

Design Controls for the Medical Device Industry, Third Edition Marie B. Teixeira, 2019-08-02 This third edition provides a substantial comprehensive review of the latest design control requirements as well as proven tools and techniques to ensure a company's design control program evolves in accordance with current industry practice It assists in the development of an effective design control program that not only satisfies the US FDA Quality Systems Regulation QSR and 13485 2016 standards but also meets today's Notified Body Auditors and FDA Investigators expectations The book includes a review of the design control elements such as design planning input output review verification validation change transfer and history as well as risk management inclusive of human factors and usability biocompatibility the FDA Quality System Inspection Technique QSIT for design controls and medical device regulations and classes in the US Canada and Europe Practical advice methods and appendixes are provided to assist with implementation of a compliant design control program and extensive references are provided for further study This third edition Examines new coverage of ISO 13485 2016 design control requirements Explores proven techniques and methods for compliance Contributes fresh templates for practical implementation Provides updated chapters with additional details for greater understanding and compliance Offers an easy to understand breakdown of design control requirements Reference to MDSAP design control requirements

Biotextiles as medical implants E. Duncan, 2013-10-31 The chapter describes the regulatory framework applicable to medical devices and specifically to biotextiles principally in the United States but also in the European Union With the emergence of new technologies and materials new forms of regulation have become necessary and some of the traditional classifications no longer match present day applications Problems encountered by manufacturers with regard to premarket approval clearance and material supply quality standards assurance are discussed as well as issues increasingly arising when one biotextile based medical device is evolved into another for a different use

Design Controls for the Medical Device Industry Marie B. Teixeira, Marie Teixeira, Richard Bradley, 2013-11-12 The second edition of a bestseller Design Controls for the Medical Device Industry provides a comprehensive review of the latest design control requirements as well as proven tools and techniques to ensure your company's design control program evolves in accordance with current industry practice The text assists in the development of an effective

Regulatory Affairs for Biomaterials and Medical Devices Stephen F. Amato, Robert M. Ezzell Jr, 2014-10-27 All biomaterials and medical devices are subject to a long list of regulatory practises and policies which must be adhered to in order to receive clearance This book provides readers with information on the systems in place in the USA and the rest of the world Chapters focus on a series of procedures and policies including topics

such as commercialization clinical development general good practise manufacturing and post market surveillance Addresses global regulations and regulatory issues surrounding biomaterials and medical devices Especially useful for smaller companies who may not employ a full time vigilance professional Focuses on procedures and policies including risk management intellectual protection marketing authorisation university patent licenses and general good practise manufacturing

Clinical Evaluation and Investigation of Medical Devices under the new EU-Regulation Wolfgang Ecker, Gerold Labek, Tarquin Mittermayr, Brigitte Raffener, Michael Ring, Bernhard Schwartz, 2020-06-04 The concept of clinical evaluation and the framework for clinical investigations have been significantly enforced within the new EU Medical Device Regulation MDR This book provides in depth and practice oriented guidance on the systematic identification and generation of clinical data through clinical investigations and other relevant sources It addresses the needs of all stakeholders be it manufacturers notified bodies or competent authorities when they have to plan perform or assess clinical evaluations and investigations for medical devices on the way to conformity assessment and CE marking It is a valuable tool of qualification for clinicians and related experts when preparing for a role of a clinical evaluator in the field either when serving any of the stakeholders or when trying to make their own involvement stand out in start ups spin offs or other development projects or in counselling services

Medical Innovation Rahul Kanegaonkar, James Tysome, 2023-03-28 An essential text for innovators this accessible book explains how medical and healthcare professionals and students who are new to innovation in healthcare can best progress their innovation projects and how those interested in healthcare innovation can develop an improved understanding of novel treatments and developments The book provides a clear framework for the innovation pathway describing step by step how projects are taken from concept to marketing and also includes a current assessment of emerging technologies that will influence medical innovation in the future Key Features Wide ranging and comprehensive coverage of the field from digital health and AI technologies through stem cell applications and robotic surgery to specialty specific innovations including those in cardiology public health and ophthalmology Illustrated with real life examples of success and failure and what can be learned from these projects Reflects a greater emphasis on clinical innovation within health systems and its inclusion in undergraduate and postgraduate medical curriculae and medically related courses Supports national and international initiatives to encourage innovation in healthcare and maximize the novel ideas generated by university staff and students as well as practicing clinicians Ideal for students at both undergraduate and postgraduate level and medical practitioners and allied medical health professionals it will also be of interest to clinical innovators and healthcare businesses seeking to increase uptake of their products both in the UK and internationally

Bioactive Glasses Heimo Ylänen, 2011-07-26 Due to their biocompatibility and bioactivity bioactive glasses are used as highly effective implant materials throughout the human body to replace or repair damaged tissue As a result they have been in continuous use since shortly after their invention in the late 1960s and are the subject of extensive research worldwide

Bioactive glasses provides readers with a detailed review of the current status of this unique material its properties technologies and applications Chapters in part one deal with the materials and mechanical properties of bioactive glass examining topics such as surface modification and cell interaction Part two is focussed on the applications of bioactive glasses covering their uses in wound healing maxillofacial surgery and bone tissue engineering among other topics With its distinguished editor and expert team of contributors Bioactive glasses is an invaluable reference for researchers and scientists in the field of biomaterials both in academia and in industry Provides a detailed review of bioactive glasses its properties technologies and applications An invaluable reference for researchers and scientists in the field of biomaterials both in academia and in industry Comprehensively covers materials and mechanical properties of bioactive glass and its applications including wound healing maxillofacial surgery and bone tissue engineering

The Regulatory Compliance Almanac Les Schnoll,2008 *Comprehensive Clinical Plasma Medicine* Hans-Robert Metelmann,Thomas von Woedtke,Klaus-Dieter Weltmann,2018-05-04 This book presents the state of the art in clinical plasma medicine and outlines translational research strategies Written by an international group of authors it is divided into four parts Part I is a detailed introduction and includes basic and recent research information on plasma sciences plasma devices and mechanisms of biological plasma effects Parts II and III provide valuable clinical insights f e into the treatment of superficial contaminations ulcerations wounds treatment of cells in cancer special indications like in heart surgery dentistry palliative treatment in head and neck cancer or the use of plasma in hygiene Part IV offers information on how and where to qualify in plasma medicine and which companies produce and supply medical devices and is thus of particular interest to medical practitioners This comprehensive book offers a sciences based practical to the clinical use of plasma and includes an extended selection of scientific medical data and translational literature

Polymers in Medical Applications B.J. Lambert,F.-W. Tang,W. J. Rogers,2001 The use of polymers in medical devices is growing at a steady rate These materials are generally relatively cheap and versatile qualities required in many bulk applications In more specialised medical devices polymeric components have been developed to meet challenging property and performance requirements This review describes the process of developing polymeric products for medical applications from design requirements through to specific examples of medical devices and packaging An additional indexed section containing several hundred abstracts from the Rapra Polymer Library database gives useful references for further reading

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