SECOND EDITION

ACADEMIC PRESS SERIES IN BIOMEDICAL ENGINEERING



# MEDICAL DEVICE TECHNOLOGIES

A SYSTEMS BASED OVERVIEW USING ENGINEERING STANDARDS

GAIL BAURA



**Patrick Vollmar** 

Medical Device Technologies Gail Baura, 2011-09-28 Medical Device Technologies introduces undergraduate engineering students to commonly manufactured medical devices It is the first textbook that discusses both electrical and mechanical medical devices. The first 20 chapters are medical device technology chapters the remaining eight chapters focus on medical device laboratory experiments Each medical device chapter begins with an exposition of appropriate physiology mathematical modeling or biocompatibility issues and clinical need A device system description and system diagram provide details on technology function and administration of diagnosis and or therapy The systems approach lets students quickly identify the relationships between devices Device key features are based on five applicable consensus standard requirements from organizations such as ISO and the Association for the Advancement of Medical Instrumentation AAMI The medical devices discussed are Nobel Prize or Lasker Clinical Prize winners vital signs devices and devices in high industry growth areas Three significant Food and Drug Administration FDA recall case studies which have impacted FDA medical device regulation are included in appropriate device chapters Exercises at the end of each chapter include traditional homework problems analysis exercises and four questions from assigned primary literature Eight laboratory experiments are detailed that provide hands on reinforcement of device concepts Biomedical Engineering Design Joseph Tranquillo, Jay Goldberg, Robert Allen, 2022-02-19 Biomedical Engineering Design presents the design processes and practices used in academic and industry medical device design projects. The first two chapters are an overview of the design process project management and working on technical teams Further chapters follow the general order of a design sequence in biomedical engineering from problem identification to validation and verification testing The first seven chapters or parts of them can be used for first year and sophomore design classes The next six chapters are primarily for upper level students and include in depth discussions of detailed design testing standards regulatory requirements and ethics The last two chapters summarize the various activities that industry engineers might be involved in to commercialize a medical device Covers subject matter rarely addressed in other BME design texts such as packaging design testing in living systems and sterilization methods Provides instructive examples of how technical marketing regulatory legal and ethical requirements inform the design process Includes numerous examples from both industry and academic design projects that highlight different ways to navigate the stages of design as well as document and communicate design decisions Provides comprehensive coverage of the design process including methods for identifying unmet needs applying Design for X and incorporating standards and design controls Discusses topics that prepare students for careers in medical device design or other related medical fields

**Biomaterials Science** Buddy D. Ratner, Allan S. Hoffman, Frederick J. Schoen, Jack E. Lemons, 2012-12-31 The revised edition of this renowned and bestselling title is the most comprehensive single text on all aspects of biomaterials science It provides a balanced insightful approach to both the learning of the science and technology of biomaterials and acts as the

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