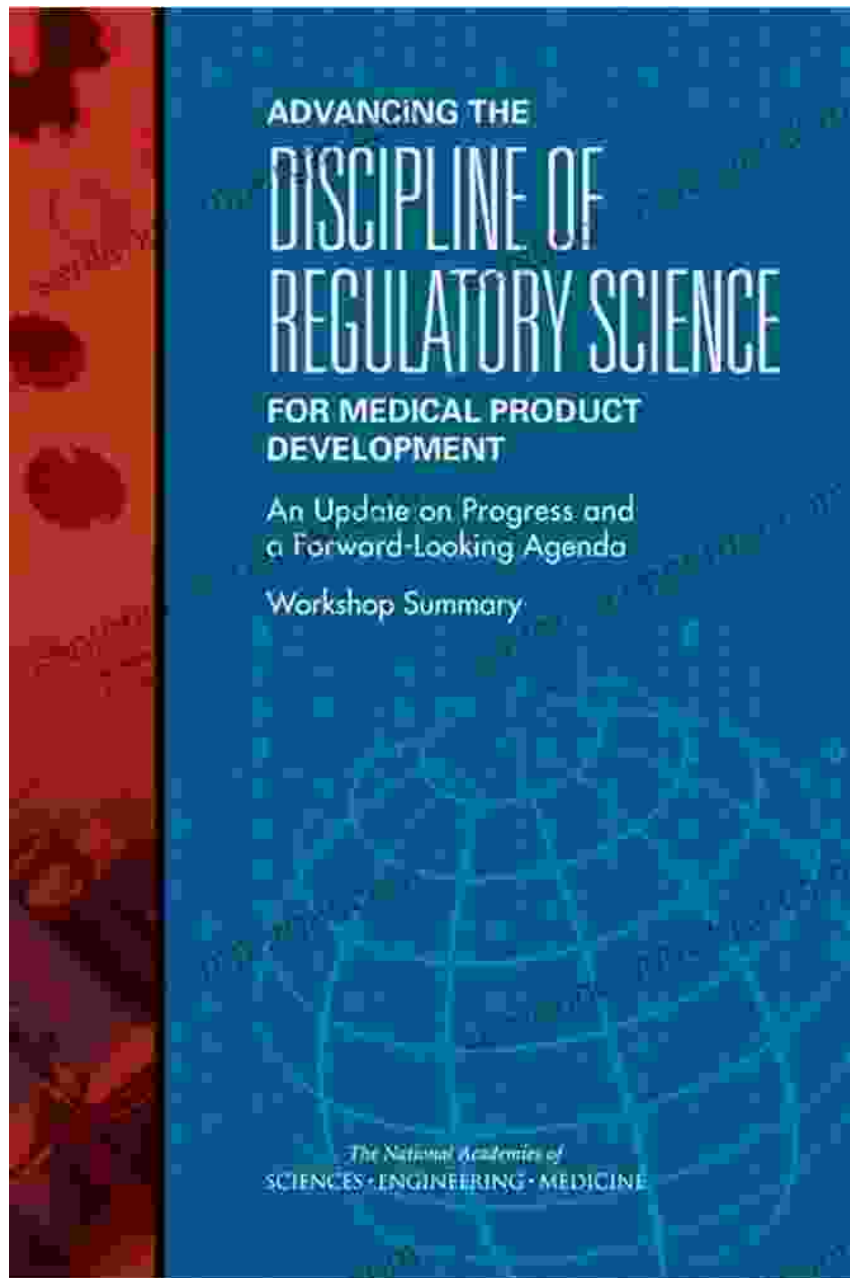


Advancing the Discipline of Regulatory Science for Medical Product Development



About the Book

This book provides a comprehensive overview of the field of regulatory science, with a focus on its application to medical product development. It

covers the history, principles, and current state of regulatory science, as well as the challenges and opportunities facing the field in the future.



Advancing the Discipline of Regulatory Science for Medical Product Development: An Update on Progress and a Forward-Looking Agenda: Workshop Summary

★★★★★ 5 out of 5

Language : English
File size : 4012 KB
Text-to-Speech : Enabled
Screen Reader : Supported
Enhanced typesetting : Enabled
Print length : 95 pages



The book is divided into three parts. The first part provides an overview of the field of regulatory science, including its history, principles, and current state. The second part covers the application of regulatory science to medical product development, including the different phases of clinical trials, the regulatory approval process, and post-market surveillance.

The third part of the book discusses the challenges and opportunities facing the field of regulatory science in the future. These challenges include the need for more efficient and effective clinical trials, the need for better communication between regulators and industry, and the need for more research on the safety and efficacy of medical products.

Benefits of Reading This Book

- Gain a comprehensive understanding of the field of regulatory science

- Learn how regulatory science is applied to medical product development
- Understand the challenges and opportunities facing the field of regulatory science in the future

Who Should Read This Book

This book is intended for a wide audience, including:

- Regulatory scientists
- Medical product developers
- Clinicians
- Researchers
- Students

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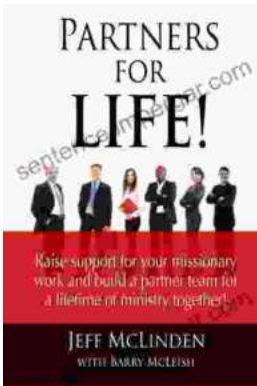
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