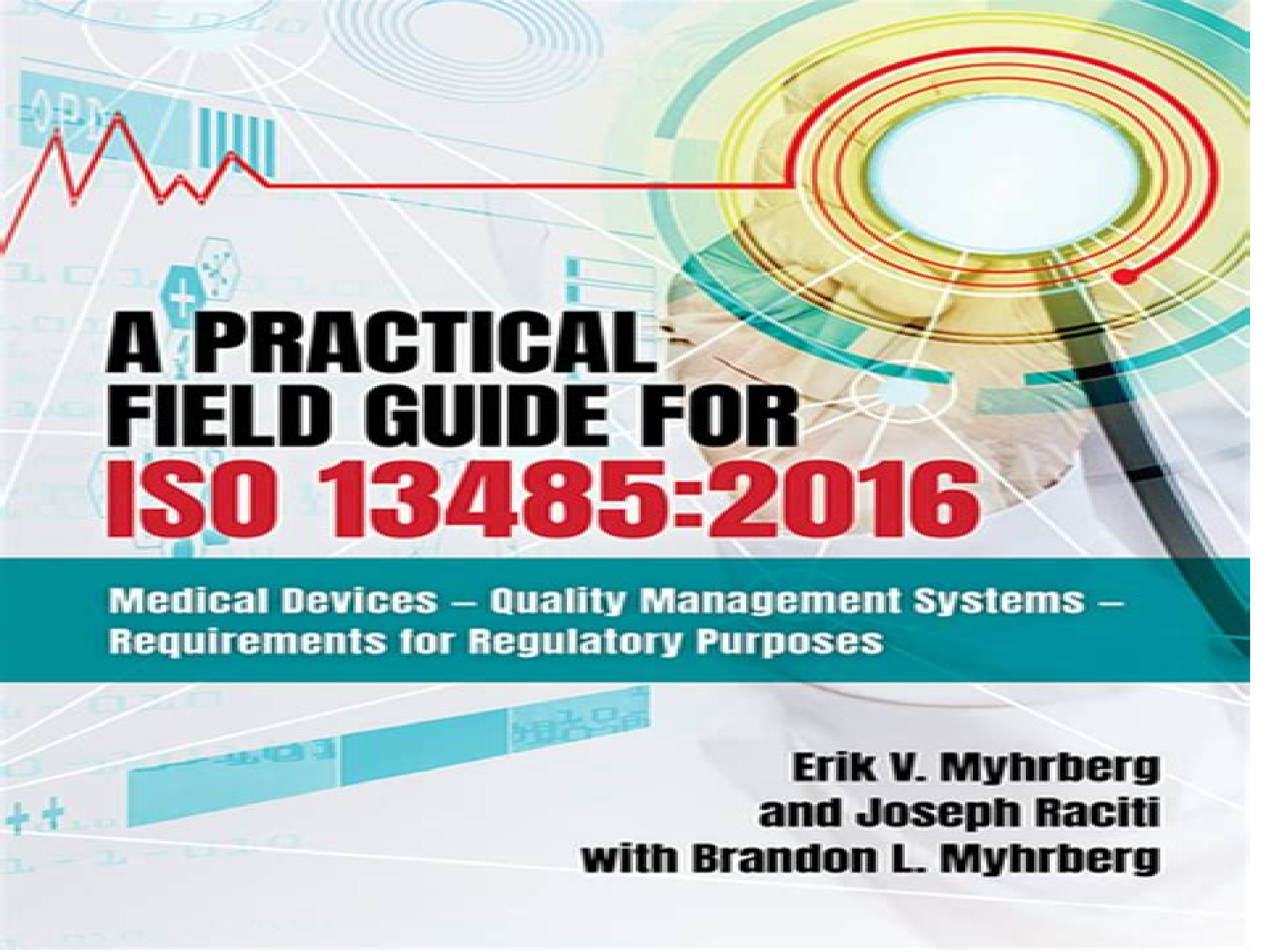


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A PRACTICAL FIELD GUIDE FOR ISO 13485:2016

**Medical Devices – Quality Management Systems –
Requirements for Regulatory Purposes**

**Erik V. Myhrberg
and Joseph Raciti
with Brandon L. Myhrberg**

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**Erik V. Myhrberg, Joseph
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The FDA and Worldwide Quality System Requirements Guidebook for Medical Devices Amiram Daniel, Ed Kimmelman, 2008-02-21 This new and expanded second edition maintains the organizational approach of the first and includes the requirements and guidance contained in the Quality System Regulation QsReg the ISO 13485 2003 standard the ISO TR 14969 2004 guidance document and as appropriate a number of the FDA and Global Harmonization

Task Force GHTF guidance documents This second edition also addresses a number of additional topics such as the incorporation of risk management into the medical device organization's QMS QMS issues related to combination products the key process interactions within a QMS effective presentation of and advocacy for a QMS during FDA inspections and third party assessments and future FDA compliance and standards activities The organization of the guidebook is based on the order of the requirements in the QSReg For each substantive requirement section there is A verbatim statement of the QSReg requirement A description of the comparable requirement in ISO 13485 2003 focusing on any additions to or differences from the requirements contained in the QSReg Excerpts of the FDA responses to relevant comment groups contained in the Preamble to the QSReg Excerpts from various FDA guidance documents related to quality management systems A description of the relevant guidance contained in ISO TR 14969 2004 focusing on any additions to or differences from the guidance in the Preamble and other FDA guidance documents and if useful excerpts from relevant GHTF guidances Authors notes giving guidance derived from the authors sixty years of regulatory compliance experience This guidance book is meant as a resource to manufacturers of medical devices providing up to date information concerning required and recommended quality system practices It should be used as a companion to the regulations standards themselves and texts on the specific processes and activities contained within the QMS

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Practical Process Validation Mark Allen Durivage, Bob (Bhavan) Mehta, 2016-07-11 For the past decade process validation issues ranked within the top six of Food and Drug Administration FDA form 483 observation findings issued each year This poses a substantial problem for the medical device industry and is the reason why the authors wanted to write this book The authors will share their collective knowledge to help organizations improve patient safety and increase profitability

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