

## Fundamentals Of Eu Regulatory Affairs Sixth Edition 2012

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The pharmaceutical and biotech industries are the most regulated in the world due to the nature of the product. Patient safety is central to all decisions, processes, procedures, guidance and regulations. Regulatory agencies such as the MHRA and EMA are in place to ensure new and existing products meet stringent safety requirements.

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## Content - Agenda | Fundamentals Of EU Regulatory Affairs

Regulatory Affairs Professionals Society v Table of Contents Section I: General Information Chapter 1 EMA and Other EU Regulatory Bodies ..... 1

## Table of Contents - Regulatory Affairs Professionals Society

Ruthanne Vendy, RAC, a Senior Principal Specialist at R&Q, was selected to be a contributing author to the Regulatory Affairs Professionals Society's (RAPS) Fundamentals of EU Regulatory Affairs, Ninth Edition, which was recently updated to cover the upcoming implementation of the European Union's new Medical Devices Regulation (EU MDR) and In Vitro Diagnostics Regulation (EU IVDR), and the subsequent impact for manufacturers, Notified Bodies, competent authorities, and other stakeholders.

## R&Q's Ruthanne Vendy Contributes to RAPS Fundamentals of ...

Regulatory Affairs Professionals Society vii Table of Contents Section I: General Information Chapter 1 FDA and Related Regulatory Agencies.....1 By Mitchell S. Berger, MPH, JD, RAC and Barry A. Berger, JD, MBA

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