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Fundamentals in Regulatory Affairs for Medical Technologies

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.

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