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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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