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Such as the MHRA and EMA are in place to ensure new and existing products meet stringent safety requirements.

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Ruthanne Vendy, RAC, a Senior
Principal Specialist at R&Q, was

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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

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Devices Regulation (EU MDR) and
In Vitro Diagnostics Regulation
(EU IVDR), and the subsequent
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R&Q's Ruthanne Vendy

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Mitchell S. Berger, MPH, JD, RAC

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